

**Caracterização molecular de linhagens clínicas de *Providencia rettgeri*
transportando os genes *bla*_{NDM} e *bla*_{TEM}**

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São Luís-MA

2017

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Defesa apresentada ao programa de Pós-Graduação Meio Ambiente como parte dos requisitos para obtenção do Título de Mestre em Meio Ambiente

Orientador Prof^ª Dra. Andrea de Souza Monteiro.

São Luís-MA

2017

Este trabalho de dissertação foi dividido em dois capítulos, artigo 1: *Providencia rettgeri* e a sua emergência como patógeno transportador de genes de resistências a carbapenêmicos e artigo 2: Caracterização molecular de linhagens clínicas de *Providencia rettgeri* transportando os genes *bla*_{NDM} e *bla*_{TEM}

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“O que não tem remédio, remediado está! ”

Andrea Monteiro

Caracterização molecular de linhagens clínicas de *Providencia rettgeri* transportando os genes *bla_{NDM}* e *bla_{TEM}*

Resumo

Providencia rettgeri é um bacilo Gram-negativo amplamente distribuído no meio ambiente e atualmente é considerado um patógeno emergente, associado a infecções nosocomiais. O objetivo deste estudo foi analisar o perfil de susceptibilidade antimicrobiana de linhagens de *P. rettgeri* isoladas de amostras clínicas e realizar uma caracterização genômica das linhagens *P. rettgeri* PR01 e *P. rettgeri* PR02. Neste estudo foram isoladas oito (08) linhagens de *P. rettgeri* a partir de amostras clínicas de pacientes atendidos em unidades de terapia intensiva. A presença do gene *bla_{NDM}* foi determinada pela reação de PCR. Além disso, as concentrações mínimas inibitórias (CIM) dos antimicrobianos foram determinadas usando o sistema automatizado VITEK-02. Os genomas das linhagens PR01 e PR02, caracterizadas como transportadora dos genes *bla_{NDM-1}* e *bla_{TEM}* foram sequenciados utilizando a plataforma Illumina - MiSeq. Após o sequenciamento, as sequências de DNA genômico pré-montadas foram anotadas usando o software Prokka. Uma análise complementar do genoma foi realizada utilizando a tecnologia de subsistema de anotação rápida (RAST). Todas as oito (08) linhagens de *P. rettgeri*, as 08 estavam albergando o gene *bla_{NDM}* e apresentaram resistência ao meropenem. Os valores de CIM, realizados pelo teste de crescimento em placa variaram de 8-128 µg/mL para o antimicrobiano meropenem. Uma caracterização genômica parcial para as linhagens *P. rettgeri* PR01 e *P. rettgeri* PR02 utilizando o RAST indicou uma ampla gama de genes relacionados a bombas de efluxo de drogas antimicrobianas. Os sistemas genéticos foram caracterizados, como a extrusão de múltiplas drogas (MATE), a divisão de família (RND), e a grande superfamília do facilitador (MFS). Nestas análises foram detectados em média 91 genes que estavam relacionados ao subsistema de virulência, doença e defesa. A análise genômica parcial de *P. rettgeri* PR01 e *P. rettgeri* PR02 confirmou a presença do gene *bla_{NDM-1}* e outros genes para a resistência à β-lactâmicos, como *bla_{TEM}*. Além disso, foram identificados vários sistemas genéticos para a produção e liberação de sideróforos, que estão associados à absorção de ferro, e outros genes, como para receptores e transportadores do anel heme e de hemina. Cerca de 43 genes foram identificados e expressaram proteínas para absorção e metabolismo de ferro. A análise do genoma das linhagens PR01 e PR02 mostrou uma alta concentração de sequências de inserção, elementos móveis e pro-fagos, sugerindo que a espécie apresenta uma plasticidade genética considerável. Ademais, ainda foi verificado a presença de diversas ilhas genômicas, associadas a marcadores de persistência frente a ao estresse químico e nutricional, como o sistema-toxina e antitoxina, contendo os genes *PasI* e *PasT*. A diversidade de elementos genéticos em *P. rettgeri* relacionados aos mecanismos de resistência a antimicrobianos, pode levar a curto prazo uma ineficiência das terapias antimicrobianas no combate a infecções causadas por este micro-organismo, e sua rápida acessão como patógeno é resultado de sua plasticidade genética.

Palavras-chave: *Providencia rettgeri*, gene *bla_{NDM}*, gene *bla_{TEM}* resistência antimicrobiana.

Abstract

Providencia rettgeri is a Gram-negative bacillus widely distributed in the environment and is currently considered an emerging pathogen, mainly associated with nosocomial infections. The objective of this study was to analyze the antimicrobial susceptibility profile of *P. rettgeri* strains isolated from clinical samples and to perform a genomic characterization of the *P. rettgeri* PR01 and *P. rettgeri* PR02 strains. In this study eight (08) strains of *P. rettgeri* were isolated from clinical samples of patients seen in intensive care units. The presence of the *bla*_{NDM} gene was determined by the PCR reaction. In addition, the minimum inhibitory concentrations (MIC) of antimicrobials were determined using the VITEK-02 automated system. The genomes of the PR01 and PR02 strains, characterized as carrier of the *bla*_{NDM-1} and *bla*_{TEM} genes were sequenced using the Illumina-MiSeq platform. After sequencing, the pre-assembled genomic DNA sequences were annotated using the Prokka software. A complementary genome analysis was performed using the Fast Annotation Subsystem technology (RAST). All eight (08) of *P. rettgeri* isolates, 08 (eight) were harboring the *bla*_{NDM} gene and showed resistance to meropenem. The MIC values, performed by the plaque growth test ranged from 8-128 µg/mL for the antimicrobial meropenem. A partial genomic characterization for the strains *P. rettgeri* PR01 and *P. rettgeri* PR02 using RAST serve indicated a wide range of genes related to efflux pumps of antimicrobial drugs, as genetic systems were characterized, such as multiple drug extrusion (MATE), the and the large family of the facilitator (MFS), in which a total of 91 genes were identified that were related to the virulence, disease and defense subsystem. The partial genomic analysis of *P. rettgeri* PR01 and *P. rettgeri* PR02 confirmed the presence of the *bla*_{NDM-1} gene and other genes for β-lactam resistance, such as *bla*_{TEM}. In addition, several genetic systems were identified for the production and release of siderophores, which are associated with iron absorption, and other genes, as well as for heme ring and hemine receptor and transporters. About 43 genes were identified and expressed proteins for uptake and metabolism of iron. Genome analysis of the PR01 and PR02 strains showed a high concentration of insertion sequences, motile elements and prophages, suggesting that the species presents a considerable genetic plasticity. It was also verified the presence of several genomic islands, associated to persistence markers in relation to chemical and nutritional stress, such as the toxin and antitoxin system, containing the *PasI* and *PasT* genes. The diversity of genetic elements in *P. rettgeri* related to mechanisms of antimicrobial resistance can lead in the short term to an inefficiency of antimicrobial therapies in the fight against infections caused by this microorganism and its rise rapid as pathogen is a result of genetic plasticity.

Key words: *Providencia rettgeri*, *bla*_{NDM} gene, *bla*_{TEM} gene, antimicrobial resistance

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LISTA DE ABREVIATURAS

- 1
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- 3 AcrB- Proteína de resistência à Acriflavina.
- 4 CIM- Concentração inibitória mínima.
- 5 CmeA- Sistema de efluxo RND, proteína de fusão de membrana CmeA.
- 6 CmeB- Sistema de efluxo RND, transportador de membrana interna.
- 7 FhuA - Receptor de membrana externa hidroxamato férrico fhuA.
- 8 FhuB - Transportador de hidroxamato férrico ABC FhuB componente permease.
- 9 FhuC - transportador de hidroxamato férrico ABC, Proteína de ligação de ATP.
- 10 FhuD - Transportador de hidroxamato férrico ABC, substrato periplasmático de
- 11 alimentos de proteína de ligação.
- 12 KPC- *Klebsiella pneumoniae* carbapenemase.
- 13 MacA- proteína de efluxo específicos de macrólídeos.
- 14 MacB- Exportação de Macrólídeos pelo ATP/ Proteína Permease MacB.
- 15 MATE_family_MDR_Pum - Proteína de extrusão antimicrobiana (Na(+)
- 16 /antiporter drogas), Bombas de Efluxo Resistentes a Múltiplas drogas da família
- 17 MATE.
- 18 MFS- Importante transportador de múltiplas drogas, superfamília (MFS).
- 19 PCR- Reação em cadeia da polimerase.
- 20 RAST- Anotações rápidas usando tecnologia de subsistemas.
- 21 RND - Proteína de fusão da membrana de bombas de efluxo a multidrogas
- 22 ShIA/ShIB- Genes de hemolisina de *Serratia marcescens*.
- 23 TolC - O tipo I de secreção de proteínas da membrana externa
- 24

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Palavras-chave: *Providencia rettgeri*, *bla_{NDM}*, *bla_{TEM}* resistência antimicrobiana.

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Artigo 1

2017

1. INTRODUÇÃO

As bactérias são micro-organismos que se caracterizam por apresentar constantes mudanças bioquímicas e fisiológicas devido a uma grande plasticidade do seu material genético (PILLAI et al., 2011). Esta plasticidade genética associada a evolução dinâmica do genoma proporciona uma resposta adaptativa celular rápida frente a pressão seletiva do meio circundante, ampliando de uma maneira significativa o repertório de espécies bacterianas que possuem amplos mecanismos para a resistência a antimicrobianos, além de um amplo espectro de fatores de virulência (PILLAI et al., 2011). Por sua vez, os efeitos das drogas antimicrobianas utilizadas para combater estes patógenos se tornam ineficazes, gerando a uma classe bacteriana conhecida como os *superbugs* (PILLAI et al., 2011).

A partir do surgimento dos “*superbugs*”, foram criados dois novos conceitos com relação a características da resistência a drogas antimicrobianas. Estes conceitos se denominam como *Extensive Drug Resistance* (XDR), que se refere a isolados de bactérias que permanecem como suscetíveis a uma ou duas categorias de drogas antimicrobianas; e o termo *Pandrug Resistance* (PDR), que se refere a isolados bacterianos não sensíveis a todas as categorias de drogas antimicrobianas conhecidas (ARMBRUSTER et al., 2014).

A circulação de bactérias em ambientes nosocomiais com perfis de resistência a drogas antimicrobianas e que se enquadram tanto no padrão XDR quanto no padrão PDR tem causado uma apreensão na comunidade médica, uma vez que drogas de última escolha estão sendo ineficazes para o tratamento de pacientes com infecções graves (ARMBRUSTER. 2014).

Dentre as drogas antimicrobianas de última escolha, os carbapenêmicos são uma classe bem distinta de fármacos, principalmente introduzidos na antibioticoterapia de uso restrito em hospitais (OLAITAN. 2015).

Os carbapenêmicos são utilizados principalmente para o tratamento de infecções causadas por linhagens de bactérias da família Enterobacteriaceae multirresistentes a drogas e produtoras de β -lactamases de espectro estendido (ESBL, do inglês *Extended spectrum beta-lactamase*) (OLAITAN. 2015). Entretanto, o surgimento de resistência bacteriana aos carbapenêmicos tem sido cada vez mais

1 relatado entre as espécies de bactérias da família Enterobacteriaceae, sendo uma
2 questão de grande preocupação na clínica médica (OLAITAN., 2015).

3 A resistência bacteriana aos carbapenêmicos e cefalosporinas de uso clínico
4 observada em espécies de bactérias da família Enterobacteriaceae está relacionada
5 principalmente com a expressão das enzimas serino- β -lactamases, das quais a uma
6 das mais importantes, é a enzima *Klebsiella pneumoniae carbapenemase* (KPC)
7 (CORDOVA., 2015) A enzima KPC foi detectada pela primeira vez no ano 2000 uma
8 estirpe de *Klebsiella* na Carolina do Norte, nos Estados Unidos (CORDOVA 2015).
9 Até o presente momento o gene blaKPC-2, e suas variantes foram detectado em
10 todos os países, se tornando endêmico em praticamente todas as regiões (POIREL.,
11 2010).

12 As enzimas do tipo KPC são classificadas no *grupo funcional 2f de Bush* e
13 conferem resistência a todos os tipos de penicilinas, diversas cefalosporinas,
14 monobactâmicos e carbapenêmicos (NORDMAM 2009). As enzimas
15 carbapenemases do tipo KPC constituem um dos mecanismos de resistência a
16 carbapenêmicos mais relevantes em espécies de bactérias da família
17 Enterobacteriaceae. Sendo, que atualmente, os genes codificantes para enzimas
18 KPC, são classificados em 21 variantes, todas estas variantes foram detectados em
19 linhagens *K. pneumoniae* e é um dos mais distribuídos em todo o mundo (NAAS.,
20 2008)..

21 Na América do Sul, a ocorrência do gene blaKPC em linhagens bacterianas
22 obtidas de amostras clínicas é atualmente considerada endêmica, e sua distribuição
23 ocorre em quase todas as regiões (ROSSI et al., 2012). Ademais, na América do Sul
24 existe uma alta prevalência para o gene blaKPC-2, em países como Colômbia,
25 Argentina e Brasil, quando se analisa espécies de bactérias com perfil de resistência
26 a multidrogas (MDR) obtidas de surtos hospitalares, sendo que as espécies mais
27 frequentemente detectadas são *Klebsiella pneumoniae*, *Acinetobacter baumannii* e
28 *Pseudomonas aureuginosa* (RIBEIRO., 2016). Entre as espécies transportadoras do
29 gene blaKPC menos frequentes estão *Serratia marcescens* e *Enterobacter cloacae*
30 (RIBEIRO., 2016).

31 Além da enzima KPC, várias espécies de bactérias Gram-negativas

expressam metallo- β -lactamases (M β LS). As M β LS são enzimas que apresentam um largo espectro de atividade de hidrólise de antibióticos β -lactâmicos, como penicilinas, cefalosporinas (SAVARD. 2014), tendo a necessidade de cations divalentes (Zn⁺⁺) como co-fatores enzimáticos, sendo inibidas pela ação de agentes quelantes ou por componentes derivados de tiois como o ácido tiolático ou ácido 2-mercaptopropiônico (2-MPA) (MENDES et al., 2006).

As M β LS estão classificadas no grupo funcional 3 de Busch e apresentam-se subdivididas em 3 subgrupos (A, B e C). Dentre os tipos de M β LS adquiridas, as de maior importância para a disseminação epidemiológica e relevância clínica são as enzimas IMP (imipenemase metallo- β -lactamase), VIM (Verona integron-borne metallo- β -lactamase), SPM (São Paulo metallo- β -lactamase), e NDM (Nova Delhi metallo- β -lactamase) (MOOSAVIAN; RAHIMZADEH, 2015). Dentre as M β LS, a enzima NDM-1 constitui atualmente uma preocupação nos ambientes hospitalares devido à sua tendência à disseminação intercontinental (CUZON et al., 2013).

As M β LS conferem resistência a diferentes carbapenêmicos, como imipenem, meropenem e ertapenem, os quais se constituem em um suporte fundamental no tratamento de infecções bacterianas resistentes aos demais antibióticos de primeira escolha. As M β LS são enzimas produzidas intrinsecamente por algumas bactérias como *Bacillus cereus*, *Chryseobacterium meningosepticum* e *Stenotrophomonas maltophilia* (WOODFORD et al., 2005; CHEN et al., 20015). Entretanto, desde o início da década de 1990, genes que codificam M β LS têm sido descritos com grande frequência em bactérias isoladas de amostras clínicas e implicadas em infecções hospitalares, como *Acinetobacter* spp., *Pseudomonas* spp (CAMPOS. 2015).

A enzima NDM-1 e suas variantes são carbapenemases do tipo M β L capaz de inativar todos os antimicrobianos da classe dos β -lactamâmicos, sendo amplamente distribuída entre as bactérias Gram-negativas (FOMDA. 2014). O alelo do gene NDM possui atualmente 17 variantes distribuídas entre várias famílias de bactérias Gram-negativas, e são responsáveis por desencadear um maior nível de resistência bacteriana aos β -lactamâmicos (DORTET, 2014) Este padrão se deve a capacidade destas enzimas em inativar praticamente todos os antibióticos β -lactâmicos, pois a existência do gene confere um padrão de resistência a multidrogas (MDR) (DORTET., 2014). A enzima NDM codificada pelo gene *bla*_{NDM-1}, foi primeiramente detectada em

1 uma linhagem de *Klebsiella pneumoniae* isolada a partir de amostras clínicas de um
2 paciente sueco que tinha sido anteriormente hospitalizado na Índia em 2009 (YONG.,
3 2009).

4 Até o momento no Brasil, foram identificadas três linhagens de bactérias de
5 espécies distintas que transportam o gene *bla*_{NDM-1}. Todos esses isolados são
6 pertencentes às espécies *Enterobacter hormaechei*, *Providencia rettgeri* e
7 *Acinetobacter baumannii* (CARVALHO-ASSEF., 2013, CARVALHO., 2014). Todas as
8 linhagens bacterianas foram isoladas na cidade de Londrina, no sul do Brasil. Em
9 qualquer caso, o modo de aquisição de isolados portadores do gene *bla*_{NDM-1} por
10 pacientes nestes estudos não está claro.

11 *P. rettgeri* é uma espécie de bactéria patogênica considerada emergente, no
12 mundo e algumas estirpes isoladas de algumas amostras clínicas tem apresentado o
13 gene *bla*_{NDM-1}. Os primeiros relatos de isolados de *P. rettgeri* de origem clínica
14 portadores do gene *bla*_{NDM-1} ocorreram em 2008 em hospitais em Israel (ROLAIN.,
15 2010). A partir de 2012, observa-se um aumento da detecção do gene *bla*_{NDM-1} em *P.*
16 *rettgeri* principalmente em linhagens isoladas de pacientes de hospitais do Nepal
17 (TADA. 2013), e em 2013 na Colômbia foi notificada a presença do gene *bla*_{NDM-1} em
18 uma estirpe de *P. rettgeri* nomeada de RB151, isolada da amostra de urina de uma
19 paciente de 53 anos (MARQUEZ-ORTIZ. 2017). Com relação aos isolados de *P.*
20 *rettgeri* de Israel, foi observado uma divergência filogenética entre as linhagens,
21 demonstrando que existe uma ampliação de clones não relacionados aos portadores
22 do gene *bla*_{NDM-1} (GEFEN 2013).

23 No Brasil recentemente foram detectadas linhagens de *P. rettgeri* portadoras
24 do gene *bla*_{NDM-1} em dois indivíduos no estado do Rio Grande do Sul, sendo um
25 paciente colonizado e outro infectado. Sendo que a análise molecular foi conduzida
26 pela Fundação Oswaldo Cruz (Fiocruz/RJ) (BRASIL, 2013). Considera-se então, que
27 a detecção de estirpes bacterianas portadoras do gene NDM no Brasil são esporádicas
28 até o momento. Diferentemente da presença do gene *bla*_{KPC}, que se tornou endêmica
29 (CARVALHO-ASSEF et al., 2013, RIBEIRO et al., 2016).

2. Gênero *Providencia*

O gênero *Providencia* é constituído de bactérias Gram-negativas pertencentes à família Enterobacteriaceae. Sendo bacilos residentes no solo, águas poluídas presentes em estações de tratamento de esgotos e em águas residuais (CLIFORD et al, 2012). As espécies que compõem o gênero *Providencia* podem ser isolados a partir de uma ampla gama de organismos vivos, sendo considerado para seres humanos microrganismos patógenos oportunistas (SHIMA et al., 2016). Dentre as espécies do gênero, *Providencia alcalifaciens* é responsável por causar infecções agudas no trato gastrointestinal e urinário, mais frequentemente em crianças e em pacientes com imunodeficiências ou que estejam sobre tratamento com imunossupressores após cirurgias (CHOI et al., 2017).

As primeiras espécies do gênero *Providencia* foram primeiramente descritas em 1920 por Ornstein, que as classificou como *Bacillus inconstans*. Em 1943-1946, Stuart isolou culturas bacterianas puras a partir de amostras de fezes de pacientes com infecções do trato gastrointestinal. Estes micro-organismos foram nomeados de "paracolon 29911". Em 1944, Gomes descreveu a espécie *Eberthella alcalifaciens*. Em 1952, Kauffmann e Edwards reclassificou este grupo como gênero *Providencia* a partir da observação das características bioquímicas específicas e sorológicas. Durante a década seguinte, a posição de bactérias do gênero *Providencia* na classificação taxonômica e suas relações com outros gêneros estreitamente relacionados como *Proteus* e *Morganella* foram revistas várias vezes culminando com a transferência de espécies de bactérias destes gêneros para gênero *Providencia* (O'HARA et al., 2000).

No ano de 1962, o gênero *Providencia* foi aceito como um gênero independente, e incluiu em conjunto com o gênero *Proteus*, à tribo Proteae; depois, o gênero *Morganella* foi anexado à tribo. Dois bio-grupos de *Providencia* foram descritos por Ewing e reconhecidas como as espécies *Providencia alcalifaciens* e *Providencia stuartii* (O'HARA et al., 1999).

1 As mais recentes mudanças na taxonomia em *Providencia* ocorreu após a
2 introdução de hibridação DNA-DNA para a classificação de espécies bacterianas.
3 Com base nas semelhanças no genoma, *Proteus rettgeri* foi reclassificado como
4 *Providencia rettgeri* e *Providencia alcalifaciens* (FARMER et al., 1977), o biogrupo 3
5 tornou-se uma espécie separada nomeada de *Providencia rustigianii* (HICKMAN-
6 BRENNER et al., 1983). Em 1986, Muller, enquanto estudava bactérias isoladas de
7 fezes dos pinguins, descreveram uma nova espécie, *Providencia heimbachae*; mais
8 tarde, uma estirpe desta espécie foi também encontrada em um paciente com diarreia
9 idiopática (O'HARA et al., 1999). Em 2006, uma nova espécie, *Providencia vermicola*,
10 foi proposto para as estirpes que infectam formas juvenis de um nematóide
11 entomopatogênico (SOMVANSI et al., 2006). Já, em 2009, representantes de duas
12 novas espécies, *Providencia sneebia* e *Providencia burhodogranariae*, foram isoladas
13 de hemolinfa de moscas de fruta (JUNEJA et al., 2009). Em 2013, uma nova espécie
14 denominada de *Providencia thailandensis*, foi descoberta após a obtenção de
15 amostras de água em um sistema de tratamento de efluentes de uma fábrica de frutos
16 do mar na província de Songkhla, na Tailândia (KHUNTHONGPAN et al., 2013).
17 Deste modo, o número de espécies do gênero *Providencia* aumentaram para nove
18 espécies reconhecidas pela taxonomia clássica.

19 O gênero *Providencia* é formado por bacilos Gram-negativos produtores de
20 urease e pertencentes a família *Enterobacteriaceae*. Algumas espécies do gênero
21 contribuem para uma série de infecções humanas, destacando, entre elas as
22 espécies *Providencia stuartii* e *Providencia rettgeri* (PILLAI. 2011). A espécie *P.*
23 *rettgeri* pode ser encontrada em diversos ambientes como água, solo e ainda
24 habitando tecidos de plantas (PILLAI. 2011). *P. rettgeri* também faz parte da
25 microbiota normal do intestino de humanos. Frequentemente tem sido relatado a
26 associação de *P. rettgeri* com infecções das vias urinárias, e em alguns casos estas
27 infecções estão associadas a quadros de gastroenterite e bacteremia (OLAITAN.
28 2015).

29 *P. rettgeri* é considerada uma bactéria patogênica emergente que pode
30 apresentar uma elevada taxa de resistência a agentes antimicrobianos comumente
31 empregados na clínica médica (OLAITAN 2015). *P. rettgeri* apresenta resistência
32 intrínseca aos antibióticos polimixinas B e E, que são drogas antimicrobianas de

1 última escolha para o tratamento de infecções causadas por bactérias resistentes a
2 antibióticos β -lactâmicos (OLAITAN. 2015), essa resistência no ambiente hospitalar
3 vem se tornando um evento de grande preocupação, principalmente pois tais genes
4 relacionados a resistência se encontram em plasmídeos moveis.

5 Dentre as espécies clínicas do gênero *Providencia*, a espécie *P. rettgeri* tem
6 emergido nas duas últimas décadas como bactéria patogênica nosocomial, que
7 frequentemente causa infecções do trato urinário em pacientes hospitalizados e
8 recentemente foi relacionada a disseminação de genes de resistência por
9 transferência plasmidial (LAHLAOUI.2014).

10 *P. rettgeri* é uma bactéria com motilidade, capaz de crescer em agar
11 MacConkey, e catalisar a dissociação da ureia em amônia e dióxido de carbono, além
12 de desaminar a fenilalanina e de produzir gás a partir da fermentação de glicose. No
13 entanto, a maioria das linhagens são não fermentadoras de lactose, *uma*
14 *característica importante utilizada para classificar micro-organismos no gênero*
15 *Providencia* (FARMER et al., 1977).

16 Os primeiros membros da espécie *P. rettgeri* foram isolados por Leo F. Rettger
17 do *Sheffield Laboratory* da Universidade de Yale. Esses isolamentos bacterianos
18 foram realizados como parte de uma investigação epidemiológica sobre uma
19 epidemia de cólera de aves em 1904 (O'HARA et al., 2000). Entretanto, os micro-
20 organismos *não foram caracterizados até o ano de 1918 quando Phillip Hadley*
21 *realizou uma avaliação superficial do gênero e propôs o nome de Bacterium rettgeri*
22 para se referir a uma nova estirpe produtoras de uréase (O'HARA et al., 2000). Em
23 1943, Rustigian e Stuart recomendaram a inclusão de *Bacterium rettgeri* no gênero
24 *Proteus*, sendo denominada de *Proteus rettgeri*, com base em características
25 bioquímicas comuns (STUART. 1945).

26 A primeira descrição de uma infecção humana provocada por *P. rettgeri* foi
27 publicado em 1951 (Goldfarb E Bakey, 1951). Este relatório, de Goldfarb e De Bakey,
28 descreve um caso de empiema (pleurite purulenta) associada a *P. rettgeri*.
29 Entretanto, as primeiras descrições de linhagens de *P. rettgeri* resistentes a drogas
30 antimicrobianas só foram relatadas já em 1971 (TRAUB. et al., 1971), e a notificação
31 de surtos hospitalares relacionados à *P. rettgeri* resistente a antibióticos em 1974,

1 com a identificação da espécie entre os pacientes de uma enfermaria cirúrgica
2 (TRAUB.1974).

3 O segundo grande surto de *P. rettgeri* foi relacionada a infecções do trato
4 urinário relatada por Edwards e colaboradores, em 1974 (EDWARDS et al., 1974).
5 Mesmo com a *P. rettgeri* implicada na etiologia em infecções do trato gastrointestinal
6 em 1986, diarreia em um viajante em 2004, e em casos de infecção ocular em 2006
7 (YOH et al., 2005; KOREISHI et al., 2006).

8 Com relação à susceptibilidade antimicrobiana, *P. rettgeri* é uma bactéria
9 tipicamente resistente à gentamicina, a colistina e a tobramicina, mas susceptível à
10 amicacina. Estirpes de *P. rettgeri*, produtoras de enzimas beta-lactamase de espectro
11 estendido (ESBL) foram relatadas com muita frequência na Europa Oriental
12 (MARCHANDIN et al., 1999). A detecção de estirpes de *P. rettgeri* portadoras da
13 enzima NDM-1 foram relatados na América do Sul a partir de 2013 (CARVALHO-
14 ASSEF et al., 2013)

15 O primeiro relato de uma estirpe de *P. rettgeri* transportando o gene *bla_{NDM}* foi
16 notificado a partir de 2013 em Porto Alegre/RS, região Sul do Brasil. Esta estirpe foi
17 obtida de uma amostra de ferida do dedo de paciente diabético com doença vascular
18 periférica (CARVALHO-ASSEF et al., 2013). O segundo relato de uma linhagem *P.*
19 *rettgeri* NDM-positiva também foi obtida de um paciente hipertenso de 55 anos
20 internado em um hospital público com cuidado terciário (Hospital Heliópolis) na cidade
21 de São Paulo, para sofrer amputação do 4º dedo devido à complicação da
22 osteomielite (CARMO. 2015).

23 3. Carreamento do gene *bla_{NDM}*

24 Alguns plasmídeos carreadores do gene *bla_{NDM}* e de suas variantes genéticas
25 têm sido extensivamente caracterizados em espécies da família *Enterobacteriaceae*
26 (YONG 2009). O histórico de descrições para o carreamento do gene *bla_{NDM}* mostra
27 uma ampla variação no contexto genético para o gene *bla_{NDM}* que estão alocados em
28 plasmídeos (YONG 2009). Os genes das variantes da enzima NDM são translocados
29 por vários tipos de plasmídeos em espécies de *Enterobacteriaceae*, como *Inc*
30 (WAILAN 2014), *IncF* (HISHINUMA 2013), *IncL/M* (HO 2011), *IncH* (VILLA 2012),
31 *IncN* (SCHULTZ 2017), e *IncX* (WANG 2014). Entretanto, observa-se que existe uma

1 variação sobre os mecanismos de replicação nestes plasmídeos e também uma
2 elevada diversidade de sequências de inserção nestes segmentos genéticos. Para os
3 plasmídeos encontrados nas espécies de *Enterobacteriaceae* que carregam o gene
4 *bla_{NDM}*, existe duas características predominantes. Em primeiro lugar, o gene *bla_{NDM}*
5 é frequentemente associado ao transposon Tn125 de 10.099 pb, este possui dois
6 elementos flanqueadores denominados de *ISAba125*, que são sequências de
7 inserção (PARTRIDGE. 2012).

8 A estrutura do transposon Tn125 contendo o gene *bla_{NDM}* em
9 *Enterobacteriaceae* é frequentemente truncada em vários comprimentos (SCHULTZ.
10 2017). Em segundo lugar, a sequência flanqueando a estrutura Tn125 envolve vários
11 mecanismos de aquisição de genes, incluindo diferentes elementos de transposição,
12 como integrons de classe 1, sequências de inserção flanqueadoras (IS), elementos
13 de transposição de repetição inversa em miniatura (MITes) (WAILAN 2016). Estas
14 características do ambiente genético do gene *bla_{NDM}* têm contribuído para os
15 diferentes contextos genéticos do gene *bla_{NDM}* relatados até o presente momento,
16 mesmo considerando o mesmo tipo de plasmídeo transportador (WAILAN. 2015).

17 O transporte do gene *bla_{NDM}* por plasmídeos em *P. rettgeri* foi recentemente
18 caracterizado em algumas linhagens bacterianas de origem clínica. Contudo, ainda
19 existem poucos estudos que enfocam o contexto genético de genes de metalo- β -
20 lactamases em *P. rettgeri*. Em um estudo utilizando como modelo a linhagem H1736
21 foi possível verificar a existência de 5 plasmídeos simultaneamente na bactéria, sendo
22 que o plasmídeo transportador do gene *bla_{NDM}* apresentava um tamanho de 48.5-kb
23 e foi relacionado ao plasmídeo pPrY2001 (*GenBank* número de acesso: KF295828.1)
24 (OLAITAN. 2015). Por sua vez, outra linhagem de origem clínica denominada de *P.*
25 *rettgeri* RB151 isolada de amostra clínica de um paciente da Colômbia em 2013, teve
26 o seu genoma sequenciado e caracterizado recentemente, nesta linhagem o
27 transporte do gene *bla_{NDM}* foi relacionado com o plasmídeo pRB151 com tamanho de
28 1.08 Kb (MARQUEZ ORTIS 2017). Além disso, estes estudos evidenciaram a
29 presença de outros elementos genéticos móveis, como genes de bacteriófagos
30 associados ao genoma de linhagens de *P. rettgeri*. Estes genes associados às
31 sequências de inserção estão ligados a uma extensa transferência de genes por
32 eventos de transferências horizontais. Estas inserções no genoma de *P. rettgeri* tem
33 contribuído para as alterações genéticas e a evolução da patogenicidade desta
34 espécie bacteriana (OLAITAN 2015).

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Artigo 2

2017

1 Introdução

2 O gênero *Providencia* faz parte da família Enterobacteriaceae e alberga bacilos
3 Gram-negativos móveis produtores de urease. Atualmente o gênero *Providencia* é
4 composto por nove espécies nomeadas: *P. alcalifaciens*, *P. stuartii*, *P. rettgeri*, *P.*
5 *rustigianii*, *P. heimbachae*, *P. vermicola*, *P. sneebia*, *P. burhodogranariea*, e *P.*
6 *thailandensis* (Shima et al., 2016). Algumas espécies do gênero contribuem para uma
7 série de infecções humanas, destacando-se, entre as espécies *Providencia stuartii* e
8 *Providencia rettgeri* (Washington; Barnhill, 2015).

9 A espécie *P. rettgeri* pode ser encontrada em diversos ambientes como água,
10 solo e ainda habitando tecidos de plantas, fazendo também parte da microbiota
11 normal do trato gastrointestinal humano (Clifford et al.; 2012). *P. rettgeri* tem sido
12 implicada em infecções das vias urinárias, e em alguns casos estas infecções estão
13 associadas a quadros de gastroenterite e bacteremia (Yoh et al., 2005; Tada et al.,
14 2014). *P. rettgeri* é considerada uma bactéria patogênica emergente e que pode
15 apresentar uma elevada taxa de resistência a agentes antimicrobianos comumente
16 empregados na clínica médica (Tshisevhe et al., 2017). *P. rettgeri* apresenta
17 resistência intrínseca aos antibióticos polimixina B e E (colistina), que são drogas
18 antimicrobianas de última escolha para o tratamento de infecções causadas por
19 bactérias resistentes a antibióticos β -lactâmicos, como carbapenêmicos (Olaitan et
20 al., 2016).

21 *P. rettgeri* é uma espécie bacteriana que tem sido descrita nos últimos anos como
22 um micro-organismo importante para a disseminação do gene *New Delhi Metallo- β -*
23 *lactamase* (comumente chamada de *bla_{NDM}*) na América do Sul (Carvalho-Assef et al,
24 2013; Marquez-Ortiz, 2017). As enzimas classificadas como NDM, são caracterizadas
25 como carbapenemases, que atualmente possuem 16 variantes detectáveis em
26 bactérias Gram-negativas. Tais enzimas são capazes de inativar alguns dos
27 antibióticos da classe dos β -lactâmicos (Khan; Maryam; Zarrilli, 2017), sendo que as
28 NDM's são amplamente distribuídas entre as bactérias Gram-negativas, e são
29 responsáveis por desencadear um maior nível de resistência bacteriana aos
30 antimicrobianos (Khan; Maryam; Zarrilli, 2017).

1 A enzima NDM-1, que é codificada pelo gene *bla*_{NDM-1}, foi primeiramente
2 detectada em linhagens de *Escherichia coli* e *Klebsiella pneumoniae* isoladas a partir
3 de amostras clínicas de um paciente sueco, cujo o histórico médico demonstrava sua
4 hospitalização na Índia no ano de 2009 (Pillai; Mcgeer; Low, 2011). A disseminação
5 do gene *bla*_{NDM-1} e suas variantes apresenta um desafio para profissionais de saúde
6 sobre como lidar com pacientes infectados (Olaitan et al., 2016).

7 A alta mobilidade do gene *bla*_{NDM} está relacionada com os mecanismos de
8 transferência horizontal de genes (THG), que ocorre principalmente por via plasmidial
9 (eventos de conjugação), atrelado ao encontro das bactérias no ambiente (Rolain;
10 Parola; Cornaglia, 2010). Estes genes de resistência são facilmente mobilizados
11 devido a sua localização no material genético móvel (MGEs), como o transposon
12 Tn125 contendo 10092pb, também encontrado na espécie *Acinetobacter baumannii*
13 (Poirel et al., 2012). Germinalmente estes encontros contribuem bastante para a
14 sobrevivência bacteriana (Jackson et al., 2011). O gene *bla*_{NDM-1} já foi detectado nos
15 plasmídeos pMR0211 em *P. stuartii* (Gann et al., 2012), e nos plasmídeos M15628 e
16 pPrY2001 portados por *P. rettgeri* (Mataseje, 2014). Entretanto, o gene *bla*_{NDM-1} em
17 algumas linhagens bacterianas pode estar integrado ao cromossomo bacteriano,
18 como evidenciado em linhagens de *P. rettgeri* (Gefen-Halevi et al., 2013).

19 Após sua descrição em 2009, o gene *bla*_{NDM-1} foi identificado em bactérias
20 isoladas de amostras clínicas de pacientes de hospitais do Reino Unido, Índia e outras
21 regiões do globo, principalmente em isolados de *E. coli* e *K. pneumoniae* (Pillai;
22 McGeer, Low, 2011). No Brasil, em 2013 foram detectadas linhagens bacterianas
23 portadoras do gene *bla*_{NDM-1} em amostras clínicas de dois pacientes de um hospital
24 do estado do Rio Grande do Sul, estirpe: CCBH11880 com número de acesso no
25 *GenBank*: GCA000805715.1 (Carvalho-Assef et al., 2013). A segunda detecção do
26 gene *bla*_{NDM-1} em espécies de bactérias circulantes em pacientes no Brasil foi
27 reportada no estado de São Paulo. Nesta descrição um paciente estava colonizado e
28 outro infectado por *P. rettgeri* (Carmo-Junior et al., 2015).

Este trabalho teve como objetivo a caracterização dos genes e mecanismos moleculares envolvidos na disseminação de resistência à carbapenêmicos e outras drogas antimicrobianas em isolados clínicos de *Providencia rettgeri*. Bem como caracterizar os genes relacionados a virulência em duas linhagens de *P. rettgeri* denominadas de PR01 e PR02 a partir de sequenciamento parcial do genoma.

Material e Métodos

Obtenção e identificação dos isolados bacterianos

Neste estudo foram avaliadas 8 (oito) linhagens de *Providencia rettgeri*, obtidas de hemoculturas e amostras de urina de pacientes atendidos em hospitais do estado do Maranhão. Estes micro-organismos foram fornecidos pelo serviço de rotina do Laboratório de Análises Clínica Cedro, localizado na cidade de São Luis-MA.

A identificação da espécie bacteriana foi obtida pelo sistema MALDI-TOF MS (*Matrix-Assisted Laser Desorption Ionization–Time of Flight Mass Spectrometry*) utilizando o sistema Biotyper (Bruker, Billerica, MA). A célula das bactérias foram cultivadas em placas de meio ágar TSB (Himedia, India), durante 24 h, a 37 °C. Para o teste, uma amostra da colônia foi transferida com auxílio de uma alça bacteriológica calibrada de 1 µL para uma lamina metálica de modo a apresentar uma fina camada sobre a placa de detecção, e seguida, foram adicionados a amostra 1 µL de uma solução de matriz (ácido α -ciano-4-hidroxicinâmico). As amostras foram submetidas à incidência de um feixe de laser para extração dos peptídeos ribossomais energizados. Finalmente, os espectros de massa adquiridos para cada linhagem bacteriana foram comparados com os espectros de massa conhecida contidas no software para classificação (Versão 3.1, Library 1.0).

O perfil de suscetibilidade a drogas antimicrobianas das linhagens bacterianas foi determinado com os cartões AST N° 105 para o sistema automatizado VITEK 2 (BioMerieux SA, Marcy-l'Etoile, França). Utilizando os antibióticos, amicacina, ampicilina, ampicilina/sulbactam, cefepima, ceftazidima, ceftriaxona, cefuroxima, cefuroxima axetil, ciprofloxacina, ertapenem, gentamicina, imipenem, meropenem, piperacilina/tazobactam. As análises de suscetibilidade foram feitas de acordo com as recomendações do fabricante e de acordo com as normas do *Clinical Laboratory Standards Institute* (CLSI ,2016). As bactérias foram estocadas em caldo BHI

acrescido de 20% de glicerol e armazenadas em criotubos à temperatura de -80 °C.

Identificação molecular dos genes *bla*_{KPC}, *bla*_{NDM-1}, *bla*_{TEM}, *bla*_{SHV} e *bla*_{AMPC}

Extração do DNA e PCR

Para a amplificação dos fragmentos dos genes alvos relacionados às enzimas β -lactamases, foram utilizadas culturas bacterianas obtidas a partir de um crescimento bacteriano a 37°C por até 18 horas em meio líquido BHI (Difco, Detroit, MI, USA). Após a incubação, 500 μ L da cultura foi centrifugada a 10.000 rpm por 20 minutos para obtenção do pellet para extração de DNA utilizando kit “Wizard® Genomic DNA Purification” (Promega Corporation, Madison, WI USA), seguindo-se o protocolo original do fabricante. O DNA genômico foi quantificado em Nanodrop™ 1000 a 260 e 280 nm e utilizado para as reações de amplificação de fragmentos de genes relacionados a resistência aos β -lactâmicos. As reações de PCR foram realizadas utilizando-se um par de específico de iniciadores (*primers*) em um volume final de reação de 25 μ L, contendo 12,5 μ L de uma solução de GoTaq® Green Master Mix (Promega, Madison, WI, USA). As sequências dos iniciadores específicos que foram utilizados estão neste estudo estão listadas na Tabela 1.

Sequenciamento genômico

As reações de amplificação foram conduzidas de acordo com métodos padronizados pelos seguintes autores: Cunningham et al. (2013), Liu et al. (2012), Khalilzadegan et al., (2015) e Dallenne et al. (2010). Os produtos de PCR foram separados em gel de agarose na concentração de 1,5% e visualizados sob exposição a luz ultravioleta. Para o sequenciamento do gene *bla*_{NDM}, os fragmentos obtidos foram submetidos a reações de sequenciamento usando o kit DYEnamic *ET Terminator Cycle Sequencing* (GE Healthcare Life Sciences, Buckinghamshire, UK), de acordo com as instruções do fabricante. Os fragmentos foram analisados no sistema de sequenciamento automático Genetic Analyze ABI PRISM® 3100 (Applied Biosystems, USA).

A qualidade dos eletroferogramas das sequências obtidas durante o processo de sequenciamento foi analisada com o software ChromasPro (<http://www.technelysium.com.au/chromas.html>). Para o alinhamento, pelo menos três sequências consenso de cada fragmento sequenciado do tipo NDM foram

1 escolhidas a partir do banco de dados para alinhamentos usando MEGA 6.0 (Tamura
2 et al., 2013). As semelhanças entre as sequências de nucleotídeos obtidas foram
3 verificadas usando BLASTn disponível em <https://blast.ncbi.nlm.nih.gov/Blast.cgi>.

4 Para identificar as variantes do gene *bla_{NDM}*, todas as sequências foram
5 traduzidas em aminoácidos usando o software ExPASy [ferramenta de tradução
6 (<http://web.expasy.org/translate/>)]. A tradução correta foi escolhida com base nos
7 dados disponíveis no *GenBank*. As sequências de aminoácidos foram comparadas
8 com as sequências da proteína NDM obtidas pelo *GenBank* usando BLASTx. Os
9 valores de similaridade para as sequências de aminoácidos variaram de 99 a 100%,
10 indicando uma região altamente conservada.

11 **Caracterização dos Grupos de Incompatibilidade dos Plasmídeos**

12 A determinação dos grupos de incompatibilidade dos plasmídeos foi realizada
13 de acordo com o método descrito por Carattoli et al. (2005). Utilizando 8 reações de
14 amplificações por cadeia da polimerase de ácidos nucleicos (5 multiplex-PCR e 3
15 simplex-PCR). Para esta reações são testados os tipos de plasmídeos mais
16 representativos da família Enterobacteriaceae, tais como: FIA, FIB, FIC, HI1, HI2, I1-
17 Iy, L/M, N, P, W, T, A/C, K, B/O, X, Y, F e FIIA. As reações de PCR foram preparadas
18 em um volume final de reação de 25 µL, contendo 12,5 µL de uma solução de GoTaq®
19 Green Master Mix (Promega, Madison, WI, USA), e 100 ng de de DNA molde por
20 reação. O programa do termociclador, à exceção do que se utilizou para a reação
21 simplex-F (cuja diferença é a temperatura de anelamento de 52 °C), consistiu num
22 passo inicial de 5 minutos a 94 °C seguido de 30 ciclos de 1 minuto à 94 °C, 30
23 segundos à 60 °C e 1 min. a 72 °C e um período de elongação final de 10 min.

Reação em Cadeia da Polimerase Multiplex-para detecção de integrons de Classe 1, 2 e 3

A caracterização dos integrons foi realizada tendo como alvos as sequencias intI1, intI2 intI3. Estas foram amplificadas pelo método de reação de PCR-Multiplex utilizando os iniciadores descritos por Goldstein et al (2001). As reações de PCR foram realizadas utilizando-se um par específico de iniciadores (*primers*) em um volume final de reação de 25 µL, contendo 12,5 µL de uma solução de GoTaq® Green Master Mix (Promega, Madison, WI, USA contendo 12,5 µL de uma solução de DNA Polimerase Martex-mix (Promega), 25 pmol de cada iniciador e 1 µL de molde de DNA (100 ng de DNA/reação). As condições de PCR foram utilizadas como descrito por Khoramrooz et al. (2016). Os produtos amplificados, 280 pb para a classe íntegron 1, 233 pb para a classe integron 2 e 600 pb para a classe *integron* 3 foram separados por eletroforese em gel de agarose a 1% contendo 0,5 µg/mL de brometo de etídio. O gel foi então fotografado sob exposição de iluminação UV.

Sequenciamento do genoma e anotações

O genoma das linhagens PR01 e PR02 portadoras dos genes *bla_{NDM}* foram submetidos ao processo de montagem de dados obtidos pelo MiSeq, como auxílio do pipeline A5 (<http://www.ncbi.nlm.nih.gov/pubmed/23028432>). Este utiliza uma abordagem de sequenciamento *de novo* para genomas procariotos A5 (*Andrew And Aaron's Awesome Assembly pipeline*). O *pipeline* e os programas associados são de códigos abertos distribuídos sob licença GPLv3, executado em ambiente Linux, e está instalado no servidor local para análise bioinformática de Neopropecta (Configuração: 03 servidores Linux, Core i7, 64 GB RAM, 2 TB HD).

Os genes sequenciados foram previstos utilizando como inferência de proteínas constante no BLAST (<http://www.ncbi.nlm.nih.gov/>). A partir dessa análise, foi identificado sequências relacionadas em outras espécies de bactérias de interesse constantes no banco de dados RAST (*Rapid Annotation using Subsystem Technology*), que é uma importante ferramenta de bioinformática para a previsão de genes putativos (acessível em <http://rast.nmpdr.org>).

1 **Construção da árvore filogenética**

2 As sequências de *Providencia rettgeri* PR01 e PR02 foram submetidas a
3 análise ao software RNAmmer presente em:
4 <http://www.cbs.dtu.dk/services/RNAmmer> (Lagesen et al., 2007) para identificação do
5 gene ribossomal 16S. As análises foram realizadas com o *algoritmo BLAST*,
6 utilizando-se pelo menos uma sequência do gene 16S rDNA. Algumas sequências do
7 gene 16S rDNA de espécies do gênero *Providencia* foram utilizadas para a
8 elaboração das relações filogenéticas. Para a determinação da raiz da Árvore
9 (*outgroup*) foi utilizada a sequência parcial do gene 16S rDNA (RNA ribossomal) de
10 *Serratia plymuthica* K-7 (NR_037111). As sequências obtidas das espécies
11 bacterianas foram então alinhadas no software MEGA 7.0 (Kumar; Stecher; Tamura,
12 2016). Para a construção da árvore filogenética foi utilizado o método de máxima
13 verossimilhança (*Maximum likelihood*) aplicando o algoritmo *MUSCLE (Multiple*
14 *Sequence Comparison by Log-Expectation)* (Edgar; Drive; ValleY, 2004). Os valores
15 de *bootstrap* menores que 70% foram ocultos na formação da figura da árvore
16 filogenética.

17 **Análise de ilhas genômicas e ilhas de patogenicidade**

18 Os genomas das linhagens *P. rettgeri* PR01 e *P. rettgeri* PR02 foram
19 analisados com relação a presença de ilhas genômicas, ilhas de patogenicidade
20 (PAIs) e de resistência a antibióticos (RIs). Para estas análises, os genomas *draft*
21 foram trabalhados utilizando *in house scripts* para a concatenação dos
22 *contigs/scaffolds*. Para estas análises primeiramente foi uma busca por genomas de
23 bactérias do mesmo gênero ou mais próximo possível aos genomas das linhagens *P.*
24 *rettgeri* PR01 e *P. rettgeri* PR02. Nesta busca chegou-se a espécie *Providencia*
25 *alcalifaciens* DSM 30120. As análises foram realizadas tendo *P. rettgeri* PR01 (query)
26 e um genoma de referência, neste caso *P. alcalifaciens* (subject) para predição das
27 ilhas genômicas. Depois de preditas as ilhas, uma figura no BRIG foi gerada,
28 utilizando, *P. rettgeri* PR01 como referência. Nesta mesma figura, foram adicionadas
29 as coordenadas das ilhas preditas anteriormente. Adicionalmente, foram plotados
30 mais dois anéis correspondentes aos *contigs* do genoma de *P. rettgeri* PR01, onde,
31 os *contigs* pares foram marcados em azul e os ímpares, em verde. Uma curadoria
32 manual foi feita em cima dos dados gerados pelo software GIPSy. Assim, foram

identificadas PAIs (ilhas de patogenicidade), GEIs (ilhas genômicas) e ilhas mistas (MSIs) utilizando o software GIPSy. Paralelamente as ilhas genômicas de *P. rettgeri* PR01 e *P. rettgeri* PR02 foram preditas utilizando o software IslandViewer 4 (Bertelli et al., 2017) associado aos programas IslandPick, SIGI-HMM, e IslandPath-DIMOB.

Resultados

Determinação do perfil de suscetibilidade a agentes antimicrobianos

Neste estudo foi observado que as 08 (oito) linhagens de *P. rettgeri* designadas como PR01, PR02, PR12, PR13, PR18, PR20, PR21 e PR27 apresentaram resistência a múltiplas drogas antimicrobianas de diferentes classes (Tabela 2), amicacina e gentamicina (aminoglicosídeos), resistência com CIM ≥ 16 $\mu\text{g/mL}$, cefalosporinas com CIM ≥ 8 $\mu\text{g/mL}$ e carbapenêmicos com CIM ≥ 4 $\mu\text{g/mL}$. Onde a PR27 foi o único isolado que apresentou um padrão de sensibilidade para todos os antibióticos testados, incluindo os β -lactâmicos.

Detecção de genes relacionados a enzimas β - lactamases

Os genes *bla*_{TEM}, *bla*_{AmpC} e *bla*_{NDM} foram identificados nas linhagens de *P. rettgeri*. Entretanto para os genes *bla*_{Shv} e *bla*_{KPC} os resultados foram negativos em todas as linhagens de *P. rettgeri*, como observado na tabela 3.

Identificação e caracterização dos grupos de incompatibilidade e integrons

Todas as linhagens de *Providencia rettgeri* apresentaram integrons de classe 1 e 2, assim como o grupo de incompatibilidade plasmidial FIIA como visto na tabela 3. Na caracterização dos grupos de incompatibilidade plasmidial, foi observado que todos os isolados de *P. rettgeri* eram portadores do *replicon* FIIA juntamente com integrons de classe 1 e 2 (Tabela 3).

Análises filogenéticas de *P. rettgeri* PR01 e *P. rettgeri* PR02

As relações evolutivas entre os isolados *P. rettgeri* PR01 e *P. rettgeri* PR02 foram determinadas pelo método da máxima verossimilhança, os resultados indicaram que as linhagens mais próximas aos isolados foram *P. rettgeri* DSM4542 *P. rettgeri* NCTC11801 (89% de *bootstrap*), estirpes já identificadas e protocoladas no genbank, apresentando uma similaridade genética com a estirpe RB151 como visto na figura 2.

1 **Caracterização genômicas das linhagens PR01 e PR02**

2 Os dados gerais da anotação dos genomas parciais das linhagens PR01 e
3 PR02 estão representados na Tabela 04. A identificação das sequências genicas
4 codificantes ou CDS (do inglês, *coding sequence*) presentes nas linhagens foi de
5 4425 e 4477 para as linhagens PR01 e PR02, respectivamente. Nas análises
6 realizadas pelo RAST foram observados vários genes associados a diferentes
7 categorias de sub-sistemas (Figuras 3,4). Contudo, neste estudo destacamos as
8 proteínas associados a bombas de efluxo para antibióticos e biocidas, proteínas para
9 captura de ferro, proteínas associadas a motilidade e elementos associados aos
10 mecanismos de transferência gênica. Estes elementos foram comparados pelo
11 BLASTn e por fim suas identidades foram então indicadas no suplemento I. Em
12 adição, verificou-se a presença de genes associados a mobilidade do genoma tais
13 como aqueles relacionados a proteína de mobilidades e bacteriófagos. Por fim, as
14 análises dos genes e proteínas relacionadas também foram direcionadas ao sub-
15 sistema relacionado ao metabolismo e captura de ferro.

16 **Caracterização do perfil de resistência e virulência das linhagens *P. rettgeri* PR01** 17 **e *P. rettgeri* PR02**

18 **Bombas de efluxo de drogas e biocidas**

19 Para uma análise mais específica sobre as características genéticas das
20 linhagens de *P. rettgeri* PR01 e PR02, estas foram caracterizadas no nível genômico
21 após o sequenciamento de nova geração, utilizando o sistema Illumina- Miseq. Nestas
22 análises foram identificadas 16 (dezesesseis) tipos de bombas de efluxo para drogas
23 antimicrobianas e biocidas para a PR01 e PR02 (Figuras 5 e 6).

24 Entre as 16 (dezesesseis) bombas de efluxo presentes nas estirpes PR01 e
25 PR02 foram identificadas a MacA (*Macrolide-specific efflux protein MacA*), MacB
26 (*Macrolide export ATP-binding/permease protein Mac*, TolC (*RND efflux system, outer*
27 *membrane lipoprotein CmeC*), AcrB (*RND multidrug efflux transporter, Acriflavin*
28 *resistance protein*), MATE_Family_MDR_Pump (Multi antimicrobial extrusion protein
29 (Na(+)) / drug antiporter, MATE family of MDR efflux pumps).

Caracterização do sistema de captação de ferro

As linhagens PR01 e PR02, apresentaram 6 categorias de genes relacionados a captação e metabolismo de ferro, tendo dois grupos majoritários representados nas categorias de captação de Heme e Hemina com 22 regiões genicas relacionadas seguido de transporte de Hemina com 14 regiões. Como visto na figura 7, possuindo genes associados as proteínas IutA, e FhuC com a localização no BLASTn de gi|490378242|WP_004257840.1 com 99% de identidade para o IutA, e dois genes para o FhuC com as respectivas localizações gi|490382157|WP_004261670.1 e gi|757595221|WP_042843418.1 ambos com 100% de identidade. Apresentando uma vasta gama de sistemas de sideróforos do tipo aerobactina e para transporte de ferro (TonB) apresentados no suplemento I.

As linhagens PR01 e PR02, possuem genes associados as proteínas IutA, e FhuC com a localização no BLASTn de gi|490378242|WP_004257840.1 com 99% de identidade para o IutA, e dois genes para o FhuC com as respectivas localizações gi|490382157|WP_004261670.1 e gi|757595221|WP_042843418.1 ambos com 100% de identidade.

Caracterização Ilhas genômicas

Nas análises de ilhas genômicas realizadas pelo programa IslandViewer 4 foi observado 22 ilhas genômicas nos genomas das linhagens PR01 e PR02 como visto nas figuras 11 e 12 com uma grande diversidade de genes relacionados a virulência bacteriana, resistência a drogas antimicrobianas e a mercúrio, sistema toxina-antitoxina, regulação genica, entre outros no suplemento II. A maior ilha genômica identificada na linhagem PR01 apresentou 114.821 pb (>10 Kb) e albergando os genes para resistência a aminoglicosídeos e β -lactâmicos. Nesta ilha genômica se encontram o gene *bla*_{TEM} e *bla*_{NDM}, além de um gene para resolvase associada ao transposon Tn3.

Paralelamente, a análise pelo programa BRIG comparou 6 ilhas de patogenicidade com 6 estirpes, sendo 2 *P. rettgeri* e 4 *P. stuartii*, com uma identidade de 100% como mostrado na figura 10.

Análise do mobiloma bacteriano

Sequências de inserção

Nas análises realizadas no genoma da linhagem *P. rettgeri* PR01 utilizando o *Software* ISSaga (*Insertion Sequence semi-automatic genome annotation*) foram detectadas 12 (doze) tipos de sequências putativas de inserção, como visto na figuras 13 e 14, com um total de 54 elementos, sendo que a maior porcentagem para os elementos de inserção foram para IS3 com 19,64% para PR01 e 19,23% para PR02 seguido de IS5 com 12,5% para PR01 e 13,46 para PR02. Cerca de 8,93% das sequências de inserção em PR01 e 11,54 % em PR02 não foram identificadas pela base de dados, sendo denominadas de ISNCY (do inglês, IS *not classified yet*, IS não classificadas).

Também foram identificadas que cerca de 10,71% e 13,46% dos elementos de inserção em PR01 e PR02, respectivamente, pertencem ao transposon Tn3. O *transposon* Tn3 foi caracterizado com uma variação de média de 2.306 a 3.282 pb apresentando uma identidade de 99% (2892/2896) com a sequência albergada no plasmídeo pNDM15-1091 transportado por *P. rettgeri* linhagem N15-01091 (*GenBank*, código: CP012903.1).

Sequências de DNA associadas à bacteriófagos

Nas análises de sequências de CDs de bacteriófagos realizadas pelo programa PHAST, foram observadas 6 regiões relacionados a estes elementos. Das sequências encontradas cinco estavam incompletas apresentando o tamanho de 2.4 Kb até 30 Kb, sendo relacionados a sequências encontradas nas bactérias da família Enterobacteriaceae (como por exemplo *Salmonella* spp), além de gram-positivos como *Staphylococcus* spp. A única sequência de bacteriófago intacta encontrada, foi referente ao fago (PHAGE_Salmon_RE_2010_NC_019488), uma sequência de 33.7 Kb (posição no genoma: 4233321-4267110). Esta sequência apresenta 49 CDs e a maioria destas sequências se encontram derivadas de um bacteriófago de *Salmonella* spp. (código no *GenBank* NC019488).

Elementos transponíveis

Nas análises realizadas pelo RAST, foram observados vários genes associados a elementos transponíveis ou transposons. Estes elementos tiveram suas

identidades comparadas pelo algoritmo BLASTn. As linhagens PR01 e PR02 albergam dois tipos de transposons, estes elementos são responsáveis pela transposição de proteínas, sendo eles Tn7 e Tn21, como pode ser visto no suplemento I.

Nas análises de elementos transponíveis foi possível verificar que sequências associadas ao Tn7 estão presentes em ambas as linhagens PR01 e PR02 (Suplemento I). Estas sequências foram relacionadas com os fragmentos de 882 e 1026 pb, e estão localizados no transposon Tn7. Contudo a linhagem PR02 contém apenas uma sequência associada ao Tn21, indicado pela sequência 372 pb, já o PR01, possui três sequências do Tn21 relacionados aos fragmentos de 372, 252 e 240 pb com suas respectivas localizações e % de identidade no BLASTn de [gi|519733601|EPP24804.1](#) e 98% de ID, [gi|1119200185|WP_072209121.1](#) com 100% de ID [gi|1119200185|WP_072209121.1](#) com 99% de ID.

Discussão

Todos os isolados bacterianos foram resistentes à carbapenêmicos, como ertapenem, meropenem e imipenem. Uma característica particular destes isolados bacterianos é que eles são originários de dois hospitais localizados a uma distância 250 km. Uma vez que os carbapenêmicos estão entre as melhores opções para tratamento de infecções por de micro-organismos Gram-negativos resistentes a múltiplas drogas antimicrobianas (Johnson, Woodford, 2017; Zafer et al, 2014), evidencia-se assim uma situação preocupante para a saúde pública, onde esses micro-organismos são uma fonte comum de infecções adquiridas em hospitais (Rolain; Parola; Cornaglia, 2010). O perfil MDR é advindo de genes relacionados a bombas de efluxo, ou genes mais específicos, como aqueles codificantes para enzimas β -lactamases, como por exemplo o gene *bla_{NDM}* e o gene *bla_{KPC}*. Estes genes são comumente encontrados em espécies de bactérias da família Enterobacteriaceae (Wu; Feng; Carattoli, 2015).

Os genes *bla_{TEM}* e *bla_{AmpC}* são responsáveis pela expressão de uma cefalosporinase do tipo ESBL, de uma AmpC β -lactamase, e uma metallo- β -lactamase, respectivamente. A resistência aos antibióticos amicacina, gentamicina, ampicilina, sulbactam, cefepima, ceftazidima, ceftriaxona, cefuroxma, ciprofloxacina, piperacilina e tazobactam se dá pela correlação de vários mecanismos de defesa,

que os MDR`s possuem, como bombas de efluxos, e enzimas específicas (Li; PlésiaT, 2015).

O gene *bla*_{TEM} expressa uma cefalosporinase do tipo ESBL, frequentemente detectado em *Escherichia coli* e *Klebsiella spp.* sendo todos micro-organismos Gram negativos (Oduro-Mensah et al., 2016). A presença do gene *bla*_{TEM} em *P. rettgeri*, indica que há troca de material genético via transferência de elementos transponíveis moveis de maneira horizontal. Uma característica dos genes de resistência é sua associação com uma região *Cassette* dos *integrans*, região a qual está localizada entre dois sítios de recombinação (*attI* e *attC*), sítios de recombinação inicial e de finalização respectivamente, desta maneira os genes moveis podem fazer parte integral da molécula de DNA (Deng et al., 2015).

O gene *bla*_{NDM} tem apresentado uma relevância clínica nos últimos anos, uma vez que este gene é responsável pela produção da enzima carbapenemase, uma metallo- β -lactamase dependente de zinco, que atua na hidrólise dos antibióticos carbapenêmicos (Galdiero et al., 2012). As carbapenemases são usualmente capazes de hidrolisar não só carbapenêmicos, mas também todos os antibióticos β -lactâmicos, como cefalosporinas, penicilinas e monobactâmicos (Lorenzoni et al., 2016).

Neste estudo foram identificados integrans de classe I e II nas linhagens de *P. rettgeri* de todas as amostras clínicas obtidas, pelo aspecto da resistência apresentada possivelmente os marcadores de resistência a β -lactâmicos estão presentes entre os sítios de recombinações dos integrans, que são comumente genes de resistência a fármacos ou genes de patogenicidades (Rajpara et al., 2015). Tais genes podem estar alocados no cromossomo uma vez que a presença dos integrans de classe I em conjunto com transposons da família Tn3 possibilitam tal inserção genica (Deng et al., 2015; Jackson et al., 2017; Rajpara et al., 2015).

O grupo de incompatibilidade FIIA detectado nas linhagens *P. rettgeri* confere um padrão único com relação a barreiras de transconjugação entre estas estirpes. O replicon FIIA (*replicon typing*) é denominado IncF ou plasmídeo conjugativo IncF é o de referência para o desenho dos iniciadores (*primers*) (Carattoli et al., 2005). Este grupo de incompatibilidade foi identificado primeiramente em *Salmonella enterica*

(Typhimurium). O *replicon* FIIA também já foi identificado em um plasmídeo recuperado de uma linhagem de *Salmonella enterica* serovar Kentucky, isolada a partir de amostras de fezes de galinha (Fricke et al., 2009). O grupo de incompatibilidade FIIA foi relacionado com fatores de virulência, e resistência a estreptomicina e tetraciclina nestas linhagens de *Salmonella enterica* (Fricke et al., 2009). Contudo, existem poucos relatos sobre a presença do grupo de incompatibilidade FIIA em outras espécies de bactérias da família Enterobacteriaceae.

Ademais, acredita-se que a presença do *replicon* FIIA seja rara em outras espécies de bactérias Gram-negativas, contudo sua aquisição por outras espécies, como *P. rettgeri* possa ser possível, principalmente se estas espécies se relacionarem em ambientes como o trato gastrointestinal de animais como as aves ou hospedeiros humanos. Recentemente, o *replicon* FIIA foi identificado em linhagens de *Enterobacter* spp. de origem clínica (Logan et al., 2016). Estas linhagens bacterianas eram portadoras dos genes *bla_{ACT/MIR}*, que conferem resistência a oximiino- β -lactâmicos e outros antibióticos β -lactâmicos (Logan et al., 2016). Verifica-se então, que existe a possibilidade do *replicon* FIIA não ser restrito a *Salmonella* spp. como anteriormente havia se proposto, podendo circular em bactérias de outros gêneros, entretanto com menor frequência de transmissão.

As linhagens PR01 e PR02 apresentaram uma grande semelhança genética com a estirpe de *P. rettgeri* RB151 que é um patógeno transportador do gene *bla_{NDM-1}*, sendo relacionada com (89% de *bootstrap*), que transporta o gene *metallo*- β -lactamase (Castro-Cardozo et al., 2017). A linhagem RB151 foi isolada no ano 2013, a partir uma amostra de urina de uma paciente do sexo feminino de 58 anos idade. Esta paciente foi diagnosticada com infecção do trato urinário e estava sendo atendida no serviço de emergência de um hospital na cidade de Bucaramanga, Colômbia (Saavedra-Rojas et al., 2015). O interessante é que o gene *bla_{NDM-1}* presente na linhagem RB151, está albergado em um plasmídeo denominado de pRB151-NDM (Marquez-Ortiz et al., 2017), o que talvez não ocorra com as linhagens PR01 e PR02. Foi observado em uma primeira análise após um sequenciamento parcial, que o plasmídeo de PR01 (pPR01 9,8 Kb) não apresentou o gene *bla_{NDM}*, somente albergava gene *bla_{TEM}* localizado no suplemento II. Este plasmídeo

apresentou 99% de Identidade com o plasmídeo pCR14_2 transportado por *Klebsiella pneumoniae* linhagem CR14 (ID no gene Bank: CP015394.1). O gene *bla*_{NDM-1} estar integrado ao cromossomo da linhagem *P. rettgeri* PR01, como tem sido relatado anteriormente para outras espécies, já que ensaios de conjugação falharam em demonstrar a transferência horizontal entre linhagens clínicas *Escherichia coli* (NDM positivas) e a linhagem receptora permissiva *E. coli* J53 (Shaheen et al., 2013).

Analises dos subsistemas de *P. rettgeri* PR01 e *P. rettgeri* PR02

Em *P. rettgeri* PR01 e *P. rettgeri* PR02, possivelmente os sistemas MATE, TolC, MacAB, AcrB RND estão relacionadas de maneira direta a resistência a drogas antimicrobianas, tais como fluoroquinolonas (ciprofloxacina, norfloxacina, norfloxacina), rifampicina, e antibióticos macrolídeos (ritromicina, azitromicina e claritromicina), além da resistência bacteriana à cobalto e zinco, esta evidencia foi correlacionada com o padrão de resistência no teste de CIM. Uma vez que as linhagens foram resistentes a maioria dos antimicrobianos testados.

Nos genomas dos isolados PR01 e PR02 também foi possível detectar a presença de 04 (quatro) genes para β -lactamases, a quais estão relacionadas com a resistência a penicilinas ID BLASTn [gi|926465406|ALD19783.1](#) com 100% de identidade, cefalosporinas de 1ª e 2ª geração, ID BLASTn [gi|490383288|WP_004262799.1](#) com 99% de identidade e [gi|446804399|WP_000881655.1](#) com 100% de identidade, de 3ª e 4ª geração ID BLASTn [gi|1002048020|AMM70781.1](#) com 100% de identidade.

As proteínas de aquisição de ferro nos isolados PR01 e PR02 foram iutA (receptor de aerobactina), FhuB (ferrichrome permease) e FhuC (proteína de ligação a ATP), que são caracterizadas como receptores associados a captação de ferro (Cabrera et al., 2001). Também, são elementos responsáveis pelo rompimento da parede celular microbiana, consequentemente os queladores de ferro como as enterobactinas ficam liberados no espaço extracelular (West et al., 1987).

O gene FhuC já foi identificado anteriormente em algumas espécies de Enterobacteriaceae, como *Edwardsiella ictaluri* (Abdelhamed et al., 2016) e *Yersinia enterocolitica* biovar 1 (Kanaujia; Bajaj; Vlrdi, 2015). Em *E. ictaluri*, a proteína FhuC contribui para a aquisição de hidroxamate férrico (Abdelhamed et al., 2016). A

1 detecção de uma grande diversidade de genes em *P. rettgeri* que são responsáveis
2 pela produção de inúmeros elementos, como sideróforos e proteínas de membrana e
3 transporte que contribuem para a captação e transporte de ferro, indicam que estes
4 produtos possam ser relevantes na virulência e a rápida multiplicação desta espécie
5 bacteriana.

6 **Ilhas genômicas**

7 A grande diversidade de genes adquiridos por transferência horizontal em *P.*
8 *rettgeri* PR01 e *P. rettgeri* PR02, indicam uma alta resiliência da espécie com relação
9 a sobrevivência a ambientes com condições adversas e estressantes, como por
10 exemplo, na presença de agentes oxidantes, biocidas, além de ambientes com
11 limitações de nutrientes.

12 O sistema toxina-antitoxina representado por diferentes sequências genicas no
13 suplemento II em *P. rettgeri* PR01 e *P. rettgeri* PR02 encontrados nas ilhas genômicas
14 podem contribuir para a formação de biofilmes em superfícies abióticas e colonização
15 de tecidos do hospedeiro, como as vias urinárias, além da geração de células
16 persistentes em condições inóspitas, como observado para células que sobrevivem
17 na presença de drogas antimicrobianas (Page; Peti, 2016; Wang; Wood, 2011). Estas
18 suposições foram levantadas, levando-se em consideração que o sistema toxina-
19 antitoxina é um fator importante em *Escherichia coli* uropatogênica para a colonização
20 nicho-específica (vias urinárias), resistência ao estresse provocado por
21 ciprofloxacina, além da resistência a oxidação por nitrito de sódio e por outros
22 intermediários reativos do nitrogênio (Norton; Mulvey, 2012).

23

1 **Análise das sequências de inserção e transposons no mobiloma bacteriano**

2 Entre as sequências de inserções observadas, a presença do Tn3 em ambas
3 as linhagens PR01 e PR02 é referente a uma transposase de 1.845 pb. O transposon
4 Tn3 tem sido relacionado a presença do gene *bla*_{TEM-1D} (uma beta-lactamase class A)
5 em algumas espécies de bactérias da família Enterobacteriaceae (Sandner-Miranda
6 et al., 2016). As linhagens *P. rettgeri* PR01 e *P. rettgeri* PR02, caracterizadas neste
7 estudo transportam o gene *bla*_{TEM}, e provavelmente existe uma boa correlação ente
8 a presença do transposon Tn3 e a disseminação deste gene.

9 A contribuição das sequências de inserção para a plasticidade genética em
10 *Providencia* spp. ainda não foi estudada profundamente, entretanto pode-se estipular
11 que sua contribuição possa ser muito similar a observada para outras espécies de
12 bactérias da família Enterobacteriaceae. Em *E. coli* O157 (um sorotipo enterro-
13 hemorrágico), por exemplo, foi sugerido que os elementos IS possivelmente tenham
14 um papel na inativação e imobilização de fagos e plasmídeos recebidos pelas
15 linhagens bacterianas (Ooka et al., 2009).

16 Nas análises de CDs realizadas a presença de sequências de bacteriófagos
17 nos genomas bacterianos, principalmente em enterobactérias, pode ser resultado da
18 aquisição do material genético por transferência horizontal em nichos mais
19 específicos como o trato gastrointestinal de hospedeiros animais (Huddleston, 2014).
20 Esta suposição se baseia na evidencia sobre a existência de uma maior
21 disponibilidade de material genético viral no lume intestinal após a sua liberação
22 durante o ciclo lítico das bactérias neste nicho (Waller et al., 2014). Evidências
23 experimentais utilizando PCR quantitativo para DNA de bacteriófagos presente em
24 fezes humanas, indicam que estes elementos transportam uma grande quantidade
25 de genes de resistência, como por exemplo genes para β -lactamases *bla*_{TEM} e *bla*_{CTX-M-1}
26 (*Modi et al., 2014*) . Estas observações podem corroborar para dimensionar o
27 impacto dos genomas de bacteriófagos na circulação e veiculação de genes de
28 resistência a antibióticos pela população bacteriana (*Modi et al., 2014*). Por fim,
29 acredita-se que os bacteriófagos associados aos eventos de transdução possam
30 aumentar o aporte de mecanismos no resistoma da população bacteriana no trato
31 gastrointestinal principalmente entre as Enterobacteriaceae, assim ter um grande
32 impacto na evolução de patógenos emergentes, como a espécie *P. rettgeri*.

Nas análises de elementos transponíveis o transposon Tn21 detectado nas estirpes PR01 e PR02, está relacionado de maneira direta a resistência antibióticos, como os aminoglicosídeos, que pode ser observado na tabela 2, tal característica advém da presença do gene *2''-aminoglycoside nucleotidyl-transferase* ANT(2'') (Schmidt; Nucken; Henschke, 1988). O Tn21 também é relacionado ao gene *ampC*, que também se mostrou positivo nas estirpes conferindo resistência a penicilinas e cefalosporinas de 1ª e 2ª nas linhagens de *K. pneumoniae*, denominadas de LCT-KP214 e LCT-KP289 (Guo et al., 2014).

A transponase do Tn7 identificada nas estirpes é heteromérica, e se distingue das demais pela complexidade de sua transposição de inserção do sítio alvo, att Tn7, por requerer quatro proteínas, Tn7, TnsABC + D e dois DNA de substrato (Young et al., 2013). A presença de sequências no DNA em *P. rettgeri* PR01 e *P. rettgeri* PR02, relacionadas ao transposon Tn7 indica uma alta versatilidade destas linhagens na disseminação de genes de resistência a antibióticos para este micro-organismo. O *transposon* Tn7 é um elemento móvel particularmente sofisticado, e se desenvolveu em alguns micro-organismos Gram negativos, como *E. coli* e *Acidithiobacillus ferrooxidans*, que apresentam estilos de vida completamente diferentes (Oppon et al., 1998; Peters, Craig, 2001). O transposon Tn7 apresenta mecanismos alternativos para promover sua propagação entre a população bacteriana. Este elemento móvel pode-se mover para sítios de baixa frequência de inserção, como outros elementos moveis transponíveis, na medida em que transpõe para muitos sítios, sem a existência de uma sequência específica no DNA receptor. Ainda, este elemento pode se mover de preferência para a uma sequência de inserção para o Tn7 em determinados *replicons*, preferencialmente em plasmídeos conjugativos. Esta última contribui de maneira significativa para a dispersão do Tn7 entre as populações bacterianas, acarretando assim na aquisição de genes associados à resistência a antibióticos por muitas espécies de bactérias (Peters, Craig, 2001).

Este estudo evidenciou várias características moleculares analisadas após o sequenciamento parcial do genoma de dois isolados de *Providencia rettgeri* (PR01 e PR02). Estas características que estão relacionados a capacidade destas linhagens em suportar a presença de vários antibióticos, principalmente da classe dos β -lactâmicos de uso clínico como cefalosporinas e carbapenêmicos. A presença de uma

1 grande diversidade de bombas de efluxo, além de enzimas inativadoras de
2 aminoglicosídeos, β -lactamases que são expressas a partir do gene *bla*_{NDM} e *bla*_{TEM}
3 estão contribuindo para a rápida disseminação e acessão de *Providencia rettgeri* em
4 ambientes nosocomiais brasileiros, aqui em primeira instancia detectados em
5 isolados de hospitais Maranhenses. Verificou-se também que as linhagens de *P.*
6 *rettgeri* podem se adaptar a ambientes com caraterísticas diversas, principalmente no
7 tocante a captação de ferro e sistemas de auto-regulação, como o sistema toxina e
8 anti-toxina. Em adição, as análises parciais do genoma indicaram ainda que esta
9 espécie possui um genoma altamente dinâmico, composto por um mobiloma
10 caracterizado por vários segmentos gênicos relacionados a proteínas de mobilidade
11 e genes de bacteriófagos. Por fim, a disseminação destes elementos genéticos pode
12 contribuir para transmissão de genes associados à virulência e resistência bacteriana
13 a biocidas e drogas antibióticas entre outras especies de bactéria da família
14 Enterobacteriaceae.

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Tabela 1 Genes relacionados à enzimas -lactamases utilizados como alvo para a Reação em Cadeia da Polimerase

Gene alvo	Iniciadores para os genes de β -lactamase	Fragmento	Temperatura de anelamento °C	Referências
<i>bla_{KPC-2}</i>	F- TGTCACTGTATCGCCGTC R- TCAGTGCTCTACAGAAAACC	1011 pb	65	(Cunningham et al., 2013)
<i>bla_{NDM-1}</i>	F- CAGCACACTTCCTATCTC R- CCGCAACCATCCCCTCTT	984 pb	56	(Liu et al., 2012)
<i>bla_{TEM}</i>	F-GAGTATTCAACATTTCCGTGTC R-TAATCAGTGAGGCACCTATCTC	800 pb	56	(Khalilzadegan et al., 2015)
<i>bla_{SHV}</i>	F-AGCCGCTTGAGCAAATTAAAC R-ATCCCGCAGATAAATCACCAC	713 pb	56	(Dallenne et al., et al., 2010)

Tabela 2 Perfil de suscetibilidade das linhagens de *Providencia rettgeri* a diferentes antibióticos

Linhagens de <i>Providencia rettgeri</i>								
Antibiotico(S)	P(R)01	P(R)02	P12	P13	P18	P20	P21	P27
Amicacina	16 (S)	16 (S)	16 (S)	≥64 (R)	16 (S)	8 (S)	16 (S)	≤4 (S)
Ampicilina	≥32 (R)	≥32 (R)	≥32 (R)	≥32 (R)	≥32 (R)	≥32 (R)	≥32 (R)	≥16 (S)
Sulbactam	≥32 (R)	≥32 (R)	≥32 (R)	≥32 (R)	≥32 (R)	≥32 (R)	≥32 (R)	ND
Cefepima	≥64 (R)	16 (S)	≥64 (R)	4 (S)	≥64 (R)	≤1 (S)	≤1 (S)	8 (S)
Ceftazidima	≥64 (R)	≥64 (R)	≥64 (R)	≥64 (R)	≥64 (R)	≥64 (R)	≥64 (R)	≤0.5 (S)
Ceftriaxona	8 (S)	16 (S)	≥64 (R)	≥64 (R)	16 (S)	2 (S)	≥64 (R)	8 (S)
Cefuroxima	≥64 (R)	≥64 (R)	≥64 (R)	≥64 (R)	≥64 (R)	≥64 (R)	≥64 (R)	ND
Ciprofloxacina	≥4 (R)	1 (S)	≥4 (R)	≥4 (R)	1 (S)	≤0.25 (S)	≥4 (R)	>2 (S)
Ertapepenem	4 (R)	4 (R)	≥8 (R)	≥8 (R)	≥16 (R)	≥8 (R)	4 (R)	≤0.25 (S)
Gentamicina	≥16 (R)	≥16 (R)	≥16 (R)	≥16 (R)	≥16 (R)	≥16 (R)	1 (S)	>8 (S)
Imipenem	≥16 (R)	≥16 (R)	≥16 (R)	≥16 (R)	≥16 (R)	≥16 (R)	≥16 (R)	≤1 (S)
Meropenem	≥16 (R)	≥16 (R)	≥16 (R)	≥16 (R)	≥16 (R)	≥16 (R)	8 (S)	≤0.25 (S)
Pieracilina/Tazobactam	≥128 (R)	≥128 (R)	≥128 (R)	≥128 (R)	≥128 (R)	≥128 (R)	≥128 (R)	≤4/4 (R)

Tabela 3 Caracterização molecular dos mecanismos de resistência em linhagens de *Providencia rettgeri*

Marcadores de Resistências	Linhagens de <i>Providencia rettgeri</i>							
	p3	p11	p12	p13	p18	p20	p21	p27
<i>bla</i> _{NDM}	P	P	P	P	P	P	NC	NC
<i>bla</i> _{SHV}	N	N	N	N	N	N	N	N
<i>bla</i> _{KPC}	N	N	N	N	N	N	N	N
<i>bla</i> _{TEM}	P	P	P	P	P	P	P	P
<i>bla</i> _{AMPC}	P	P	P	P	P	P	P	P
Plasmídeos	FIIA5	FIIA5	FIIA5	FIIA5	FIIA5	FIIA5	FIIA5	FIIA5
Integrans	Classe 1 e 2	Classe 1 e 2	Classe 1 e 2	Classe 1 e 2	Classe 1 e 2	Classe 1 e 2	Classe 1 e 2	Classe 1 e 2

Tabela 4 Características da montagem do genoma das linhagens *Providencia rettgeri* PR01 e PR02 indicadas pelo RAST

Características	Linhagens	
	PR01	PR02
Tamanho	4.734,158	4.665,741
Conteúdo de GC	41.2	41.5
N50*	27355	102640
L50*	51	13
Número de sub-sistemas	367	510
Número de se sequências codificantes	4425	4477
Número de RNase	88	89

Figura 1- Detecção do gene blaNDM em linhagens de *Providencia rettgeri*, fragmento de 984 pb

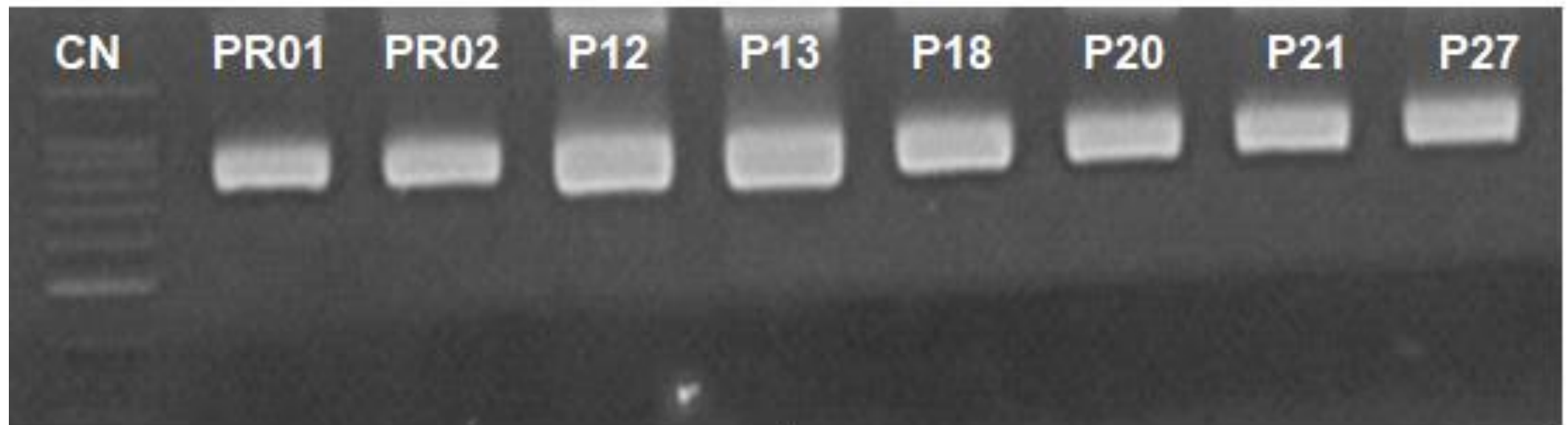


Figura 2-Análises de filogenia molecular das linhagens *Providencia rettgeri* PR01 e *Providencia rettgeri* PR02, pelo método de máxima verossimilhança. A porcentagem em que os taxa associados agrupados na árvore são mostrados ao lado dos ramos.

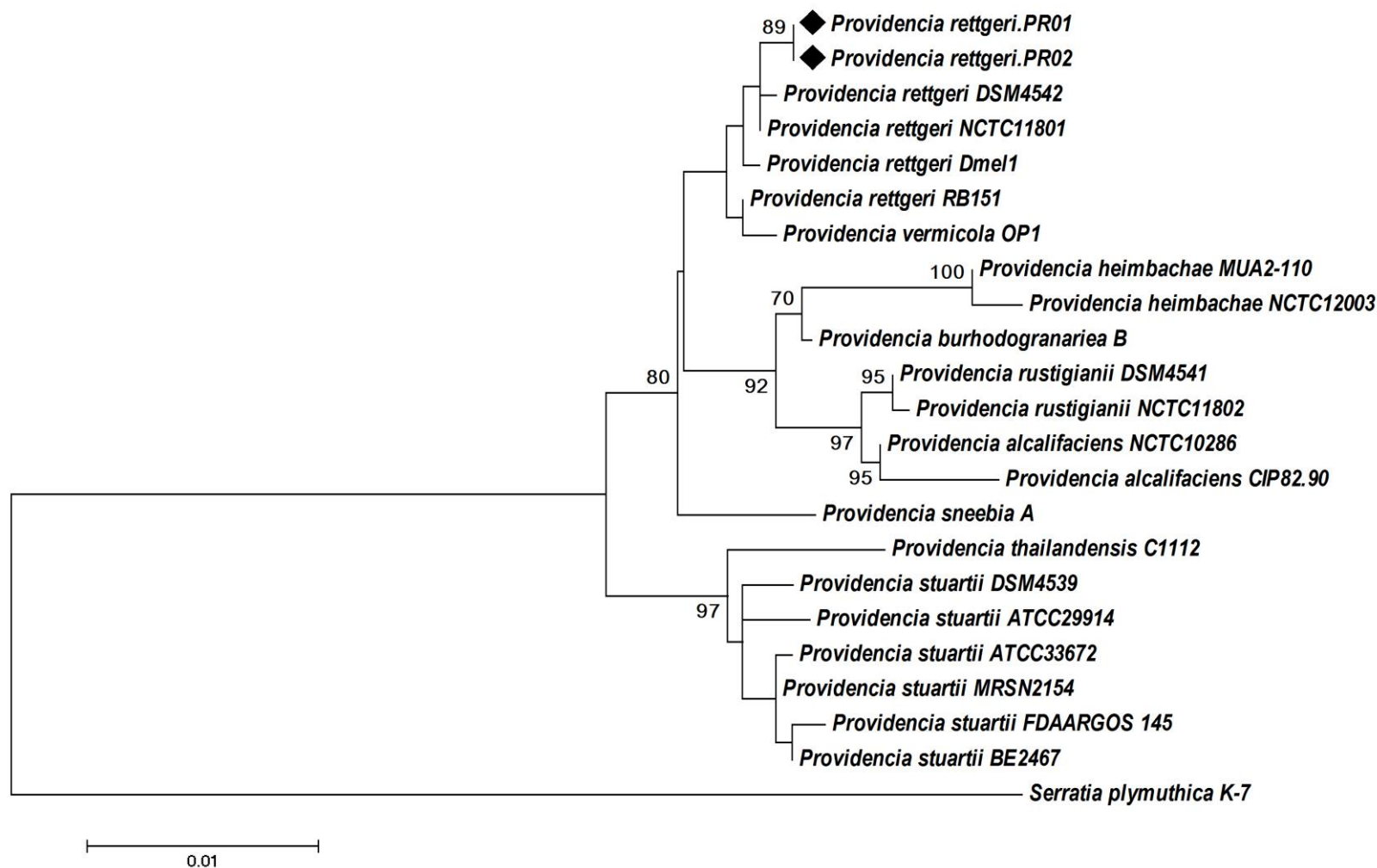


Figura 3-Perfil genético da linhagem *Providencia rettgeri* PR01

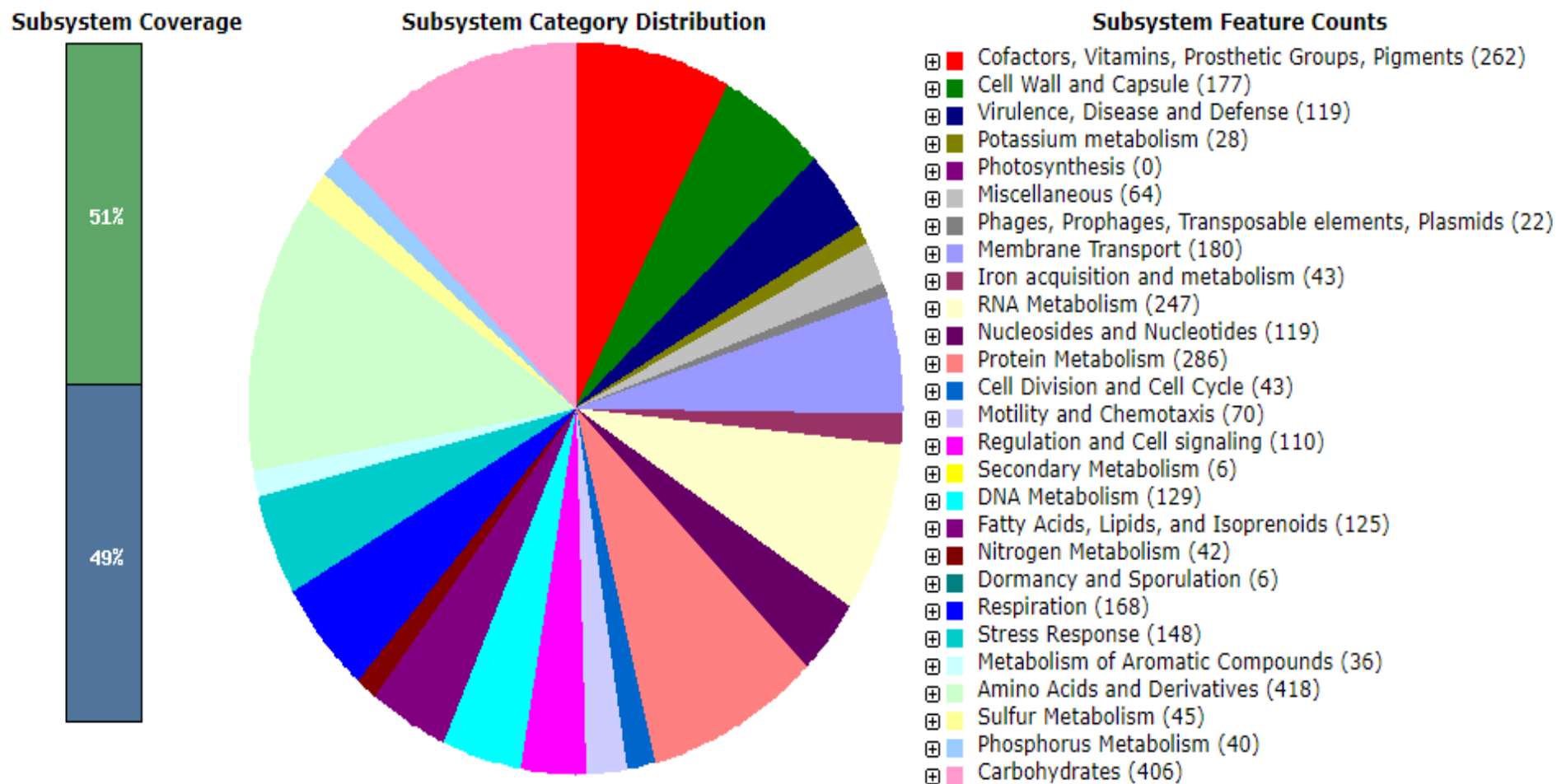


Figura 4-Perfil genético da linhagem *Providencia rettgeri* PR02

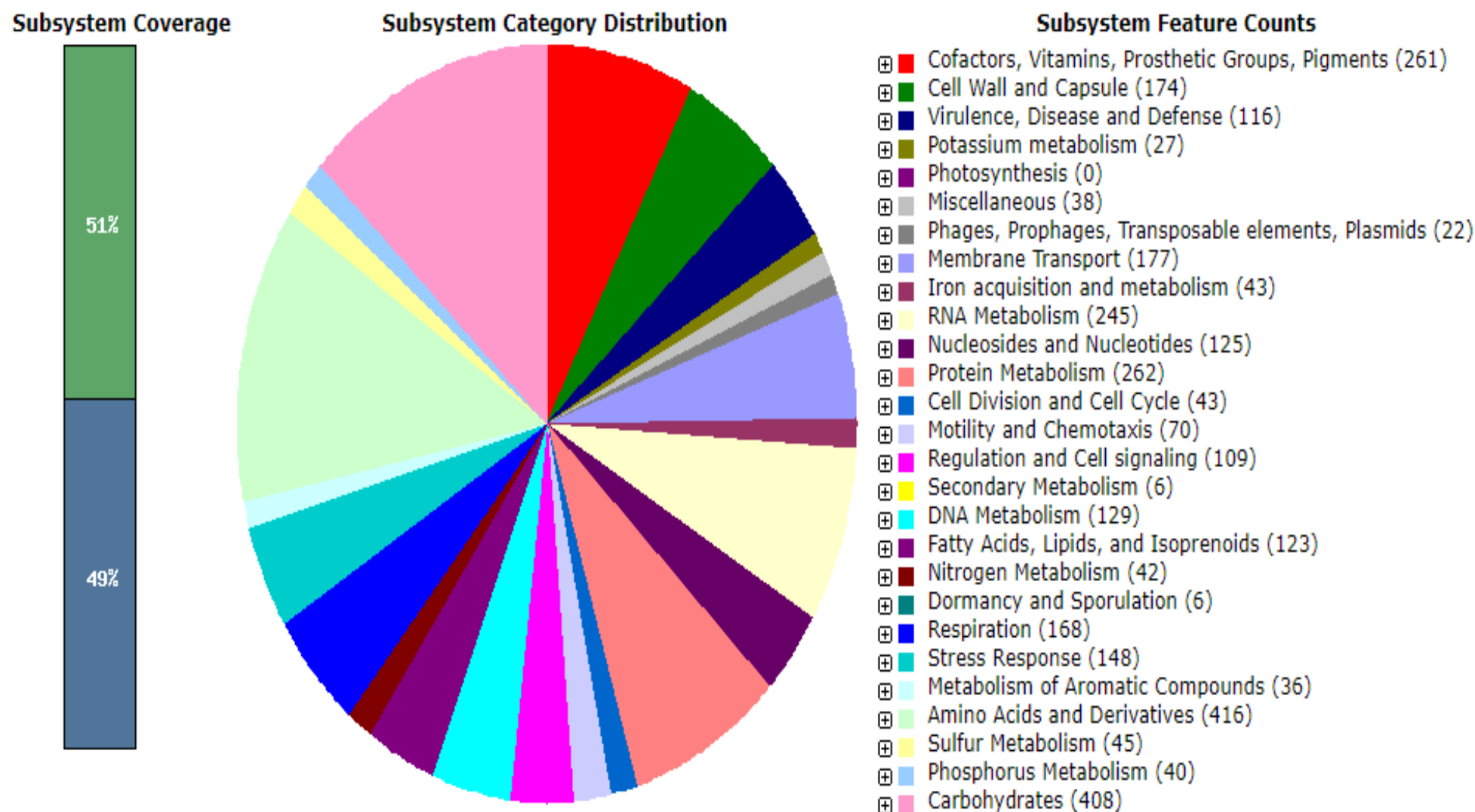


Figura 5-Perfil genético do subsistema de virulência e defesa da *Providencia rettgeri* PR02

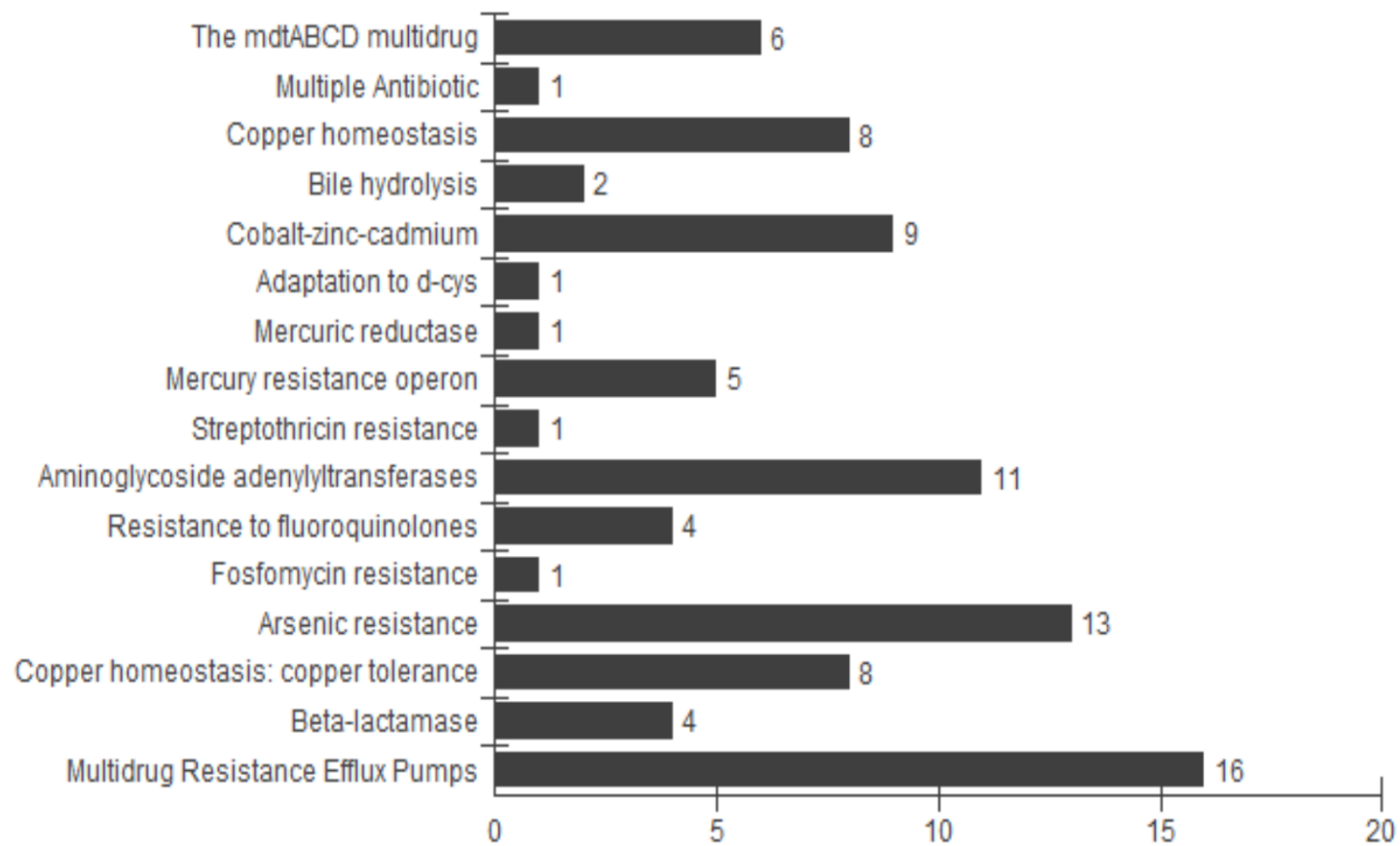


Figura 6-Perfil genético do subsistema de virulência e defesa da *Providencia rettgeri* PR02

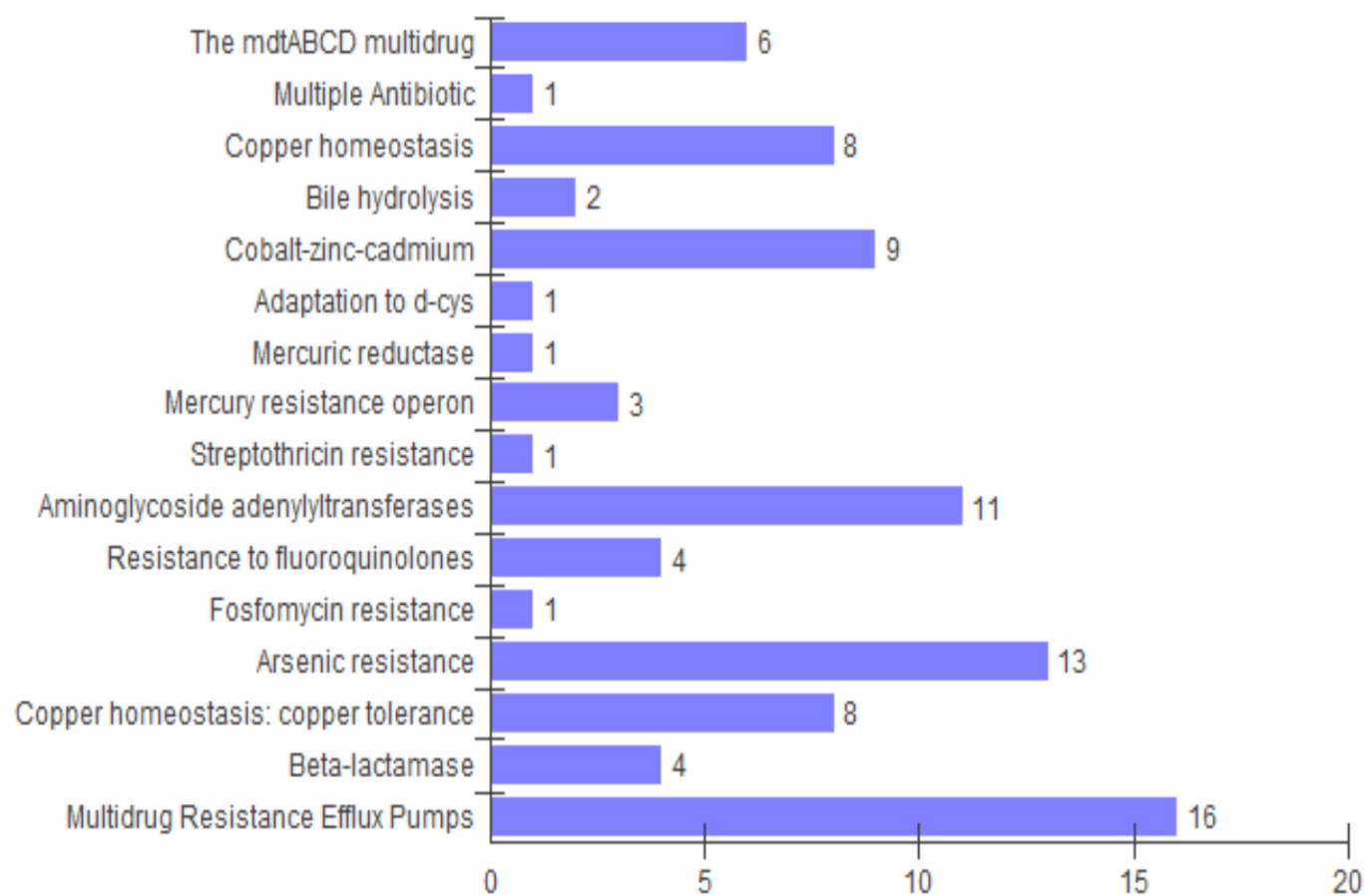


Figura 7-Perfil genético do subsistema de metabolismo e aquisição de ferro das linhagens de *Providencia rettgeri* PR01 e PR02

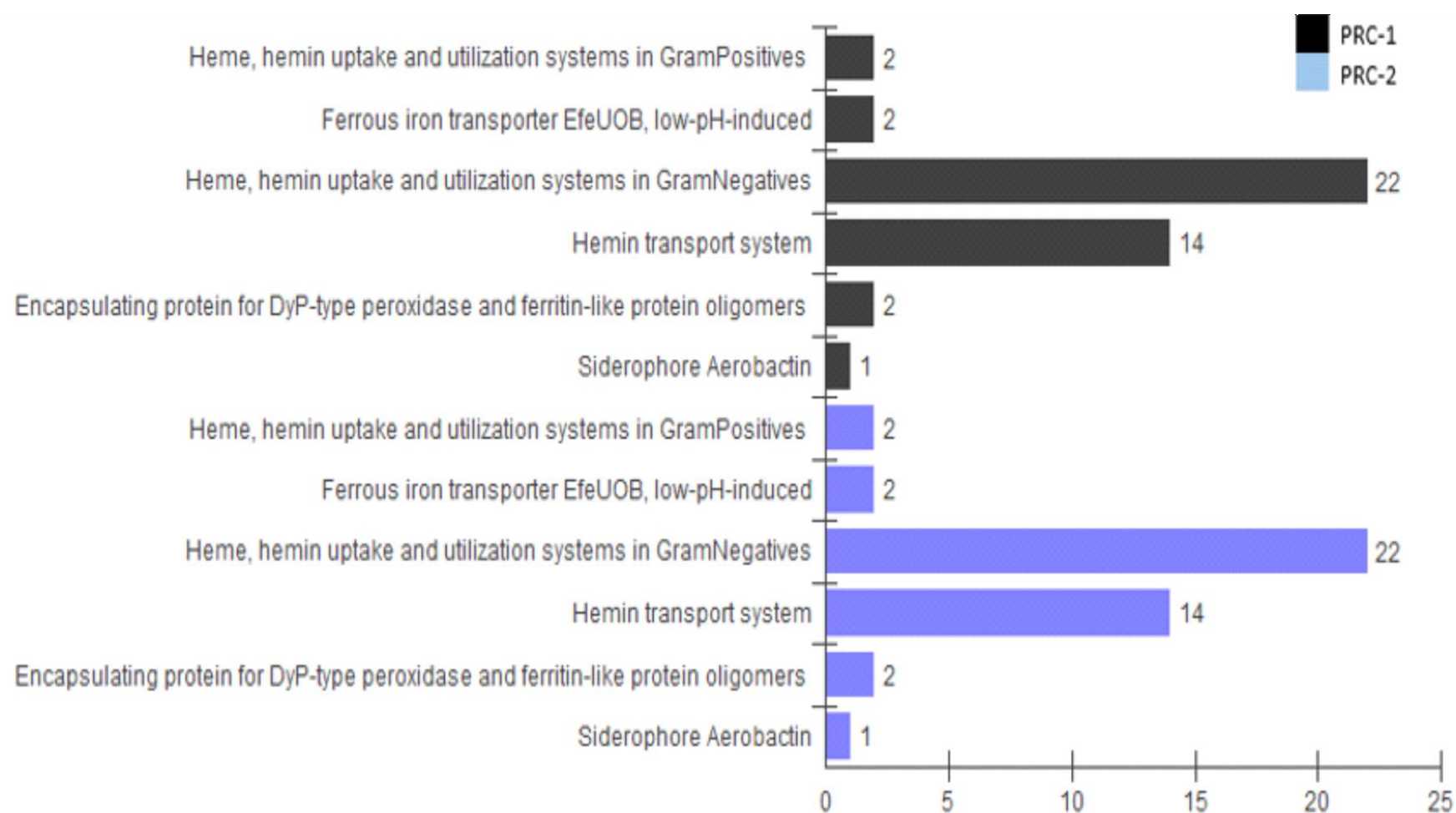


Figura 8-Perfil genético do subsistema apresentando fagos, pro-fagos, elementos móveis transponíveis e plasmídeos das linhagens de *Providencia rettgeri* PR01 e PR02

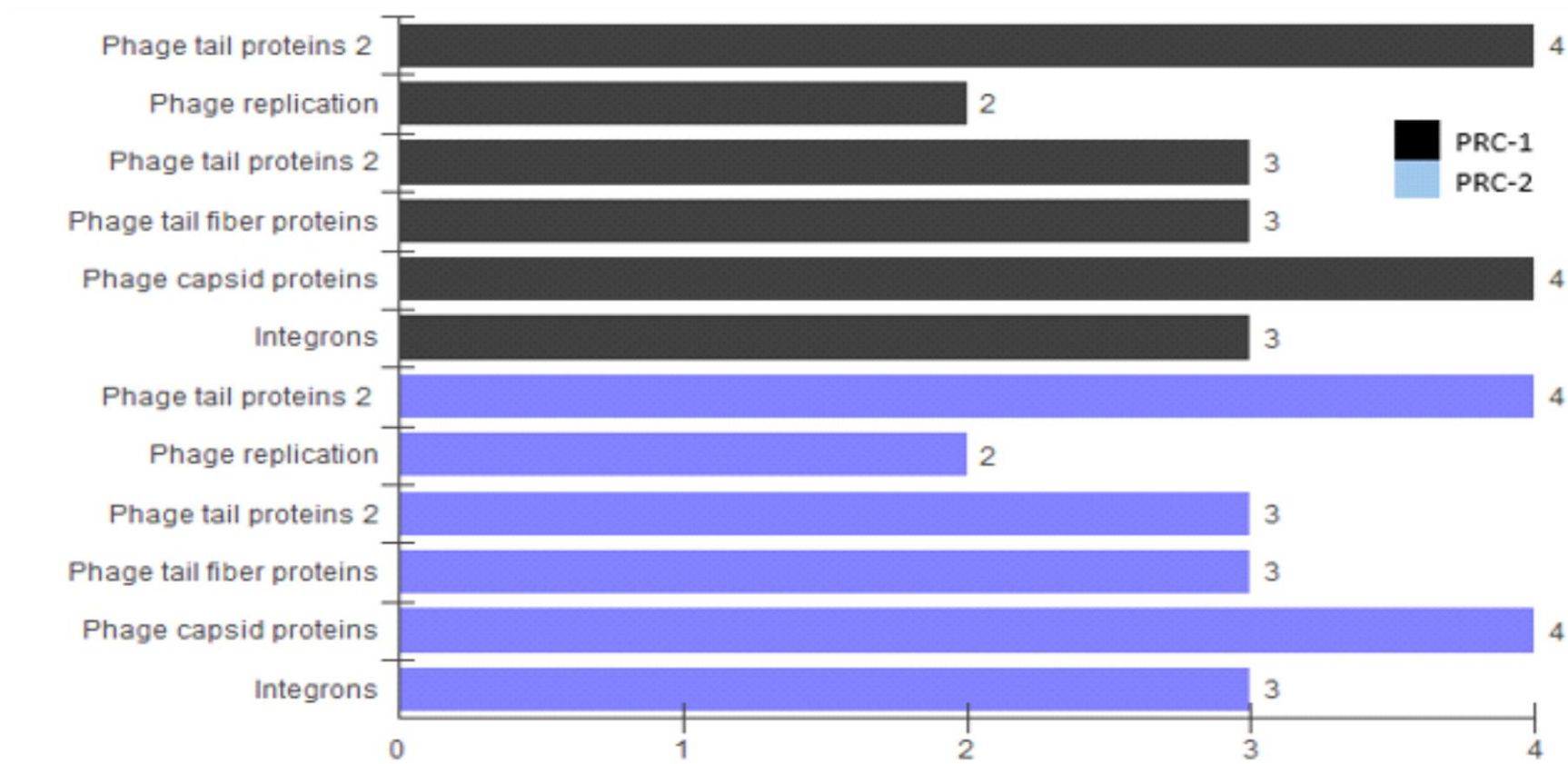


Figura 9-Perfil genético do subsistema de mobilidade e quimiotaxia das linhagens de *Providencia rettgeri* PR01 e PR02

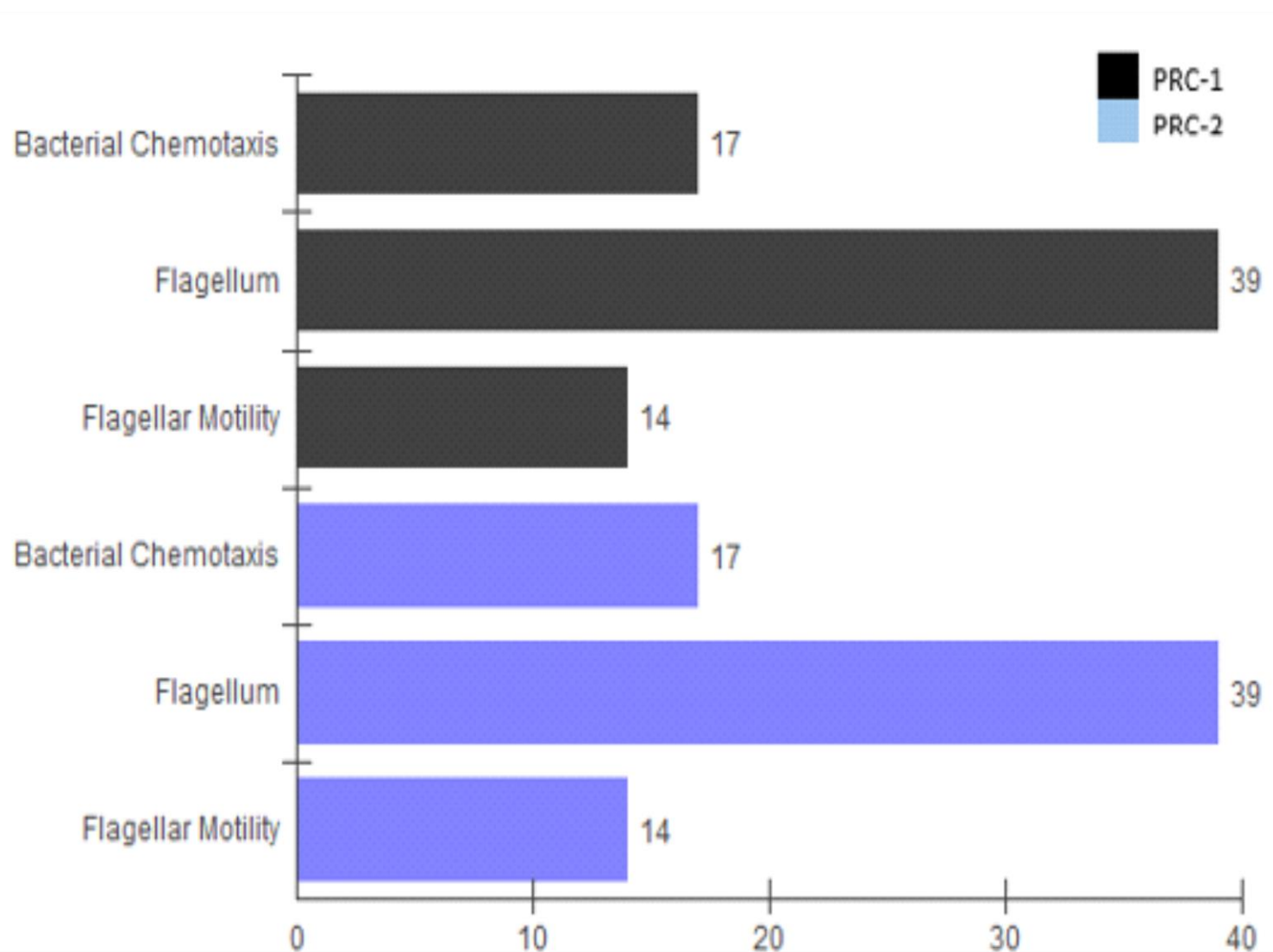


Figura 10- Predição de ilhas genômicas, ilhas de patogenicidade (PAIs) e de resistência a antibióticos (RIs) em *P. rettgeri* PR01 e *P. rettgeri* PR02

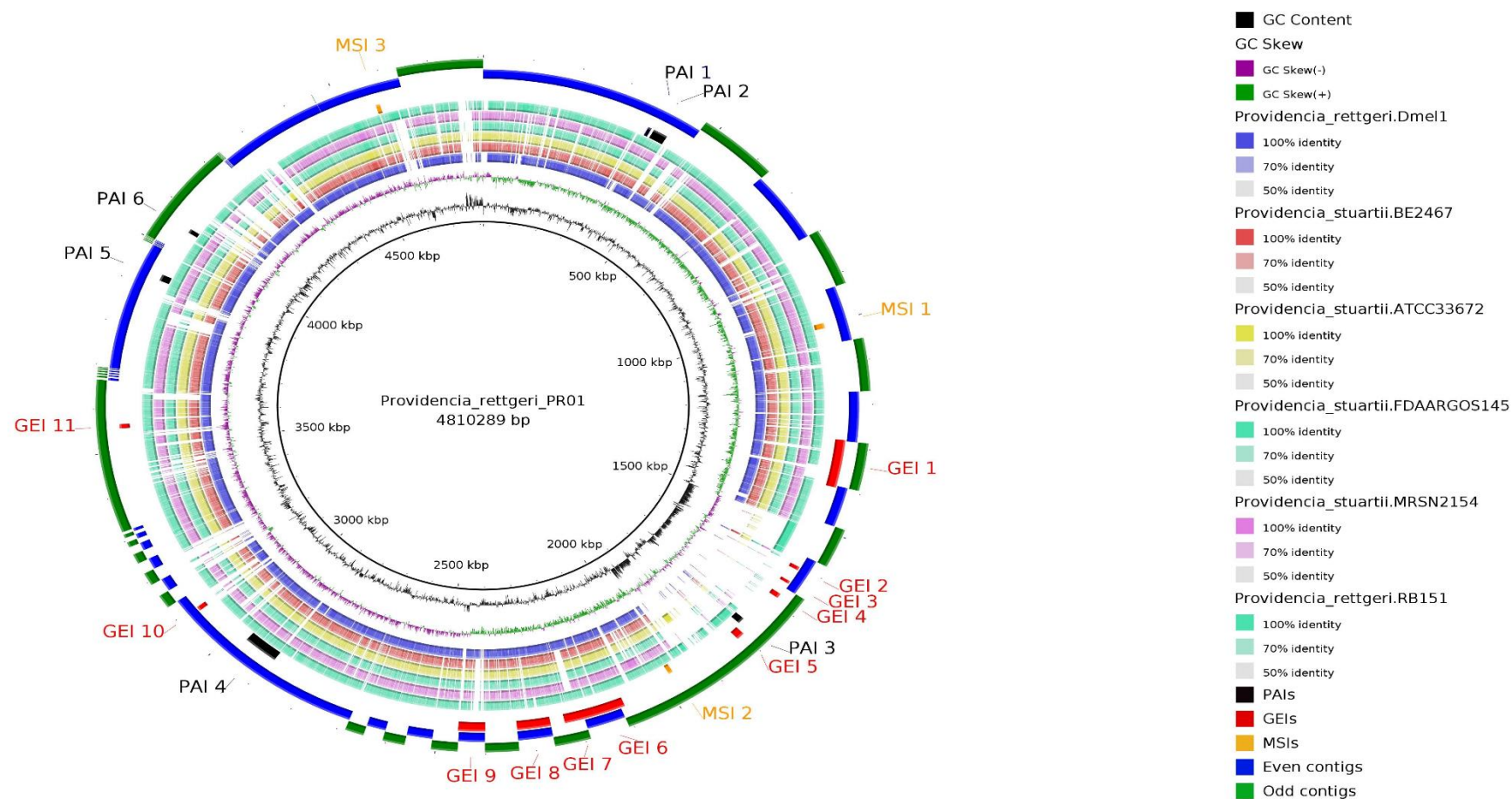


Figura 11- Ilha Genômica PR01

Mapa genômico das ilhas genômicas presentes em *Providencia rettgeri* PR01

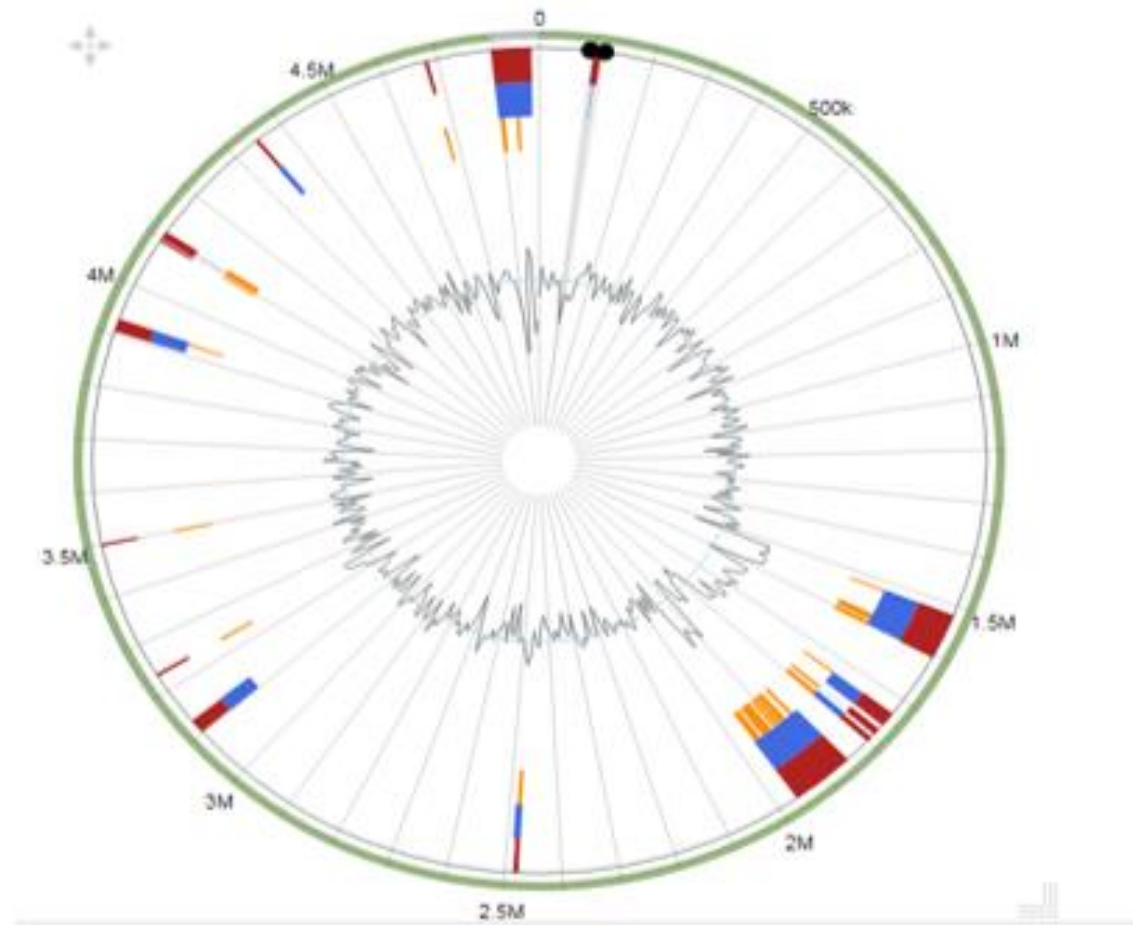


Figura 12- Ilha genômica PR02

Mapa genômico das ilhas genômicas presentes em *Providencia rettgeri* PR02

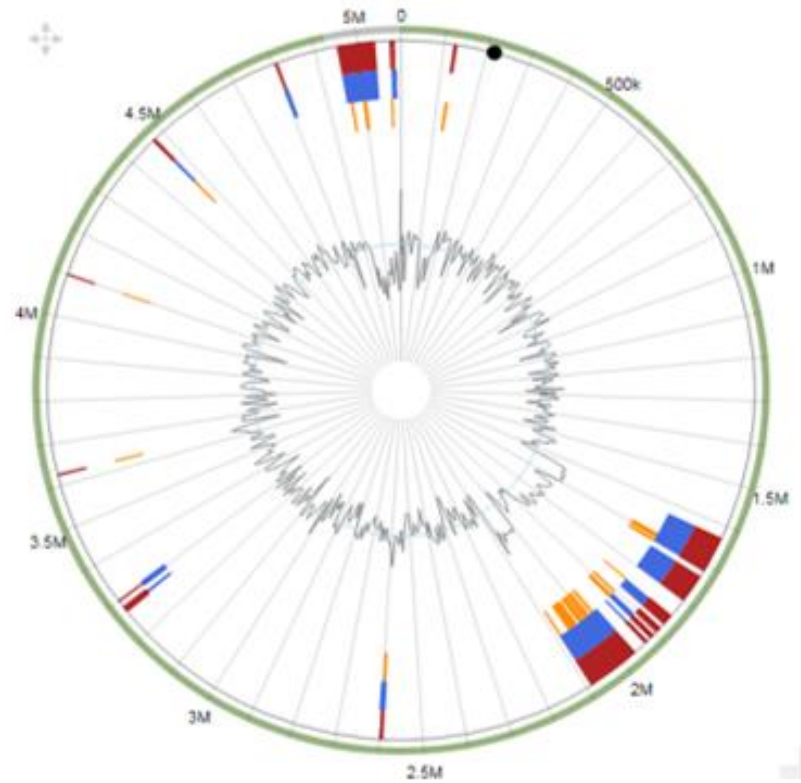
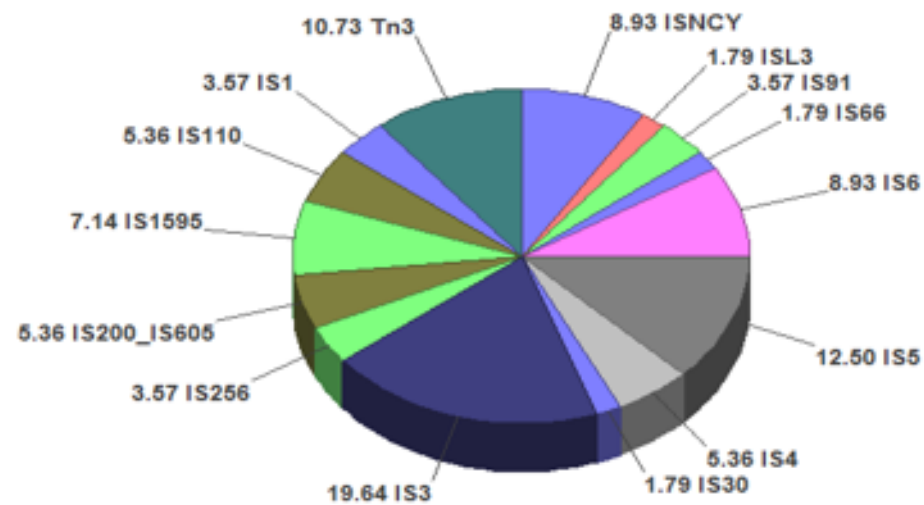


Figura 13- Sequência de Inserção PR01

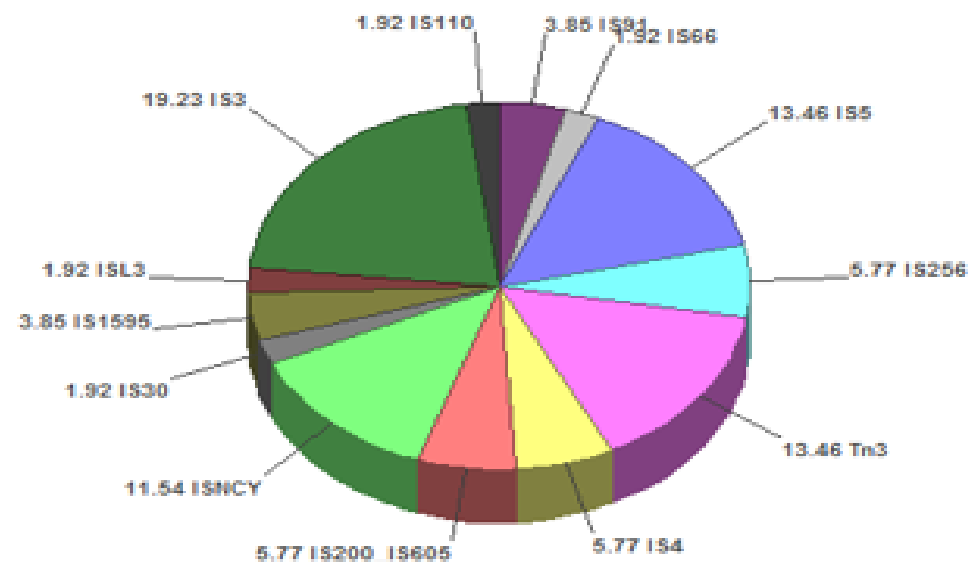
Sequências de Inserção Encontradas na Providencia PR01



Tn3 10.73%	IS110 5.36%	IS200_IS605 5.36%	IS3 19.64%	IS4 5.36%	IS6 8.93%	IS91 3.57%
IS1 3.57%	IS1595 7.14%	IS256 3.57%	IS30 1.79%	IS5 12.50%	IS66 1.79%	ISL3 1.79%
ISNCY 8.93%						

Figura 14-Sequência de Inserção PR02

Sequências de Inserção Encontradas na Providencia PR02



IS110 1,92%	IS1595 3,85%	IS200_IS605 5,77%	IS256 5,77%	IS66 1,92%
IS3 19,23%	IS30 1,92%	IS4 5,77%	IS5 13,46%	IS91 3,85%
ISL3 1,92%	ISNCY 11,54%	Tn3 13,46%		

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Suplemento I

Sub-sistemas PR01

Virulence, Disease and Defense

Sigla	Gene	Tamanho	Aminoácidos	Identidade	Sequenciamento	E. value
Tolerance to colicin E2						
CbrC-like	Colicin E2 tolerance protein CbrC-like protein	588bp	196aa	99%	gi 490381407 WP_004260923.1	8.00E-151
CreA	Conserved uncharacterized protein CreA	477bp	159aa	100%	gi 491044024 WP_004905686.1	2.00E-108
Colicin V and Bacteriocin Production Cluster						
R1	tRNA pseudouridine synthase A (EC 4.2.1.70)	816bp	272aa	100%	gi 490377681 WP_004257279.1	0.0
DedA	DedA protein	687bp	229aa	99%	gi 490377686 WP_004257284.1	5E-159
R3	Acetyl-coenzyme A carboxyl transferase beta chain (EC 6.4.1.2)	972bp	324aa	99%	gi 490377689 WP_004257287.1	0.0
R4	Dihydrofolate synthase (EC 6.3.2.12), Folylpolyglutamate synthase (EC 6.3.2.17)	1287bp	429aa	99%	gi 490377692 WP_004257290.1	0.0
R5	Dihydrofolate synthase (EC 6.3.2.12), Folylpolyglutamate synthase (EC 6.3.2.17)	1287bp	429aa	99%	gi 490377692 WP_004257290.1	0.0
DedD	DedD protein	642bp	214aa	100%	gi 739086890 WP_036957975.1	3E-147
*toxin	Colicin V production protein	504bp	168aa	100%	gi 490377706 WP_004257304.1	3E-115
PurF	Amidophosphoribosyltransferase (EC 2.4.2.14)	1518bp	506aa	100%	gi 490377709 WP_004257307.1	0.0
Mediator of hyperadherence YidE in Enterobacteria and its conserved region						
YidE	Mediator of hyperadherence YidE	1659bp	553aa	99%	gi 490383146 WP_004262657.1	0.0
YidR	Uncharacterized protein YidR	1317bp	439aa	99%	gi 291312036 EFE52489.1	0.0
YidQ	Outer membrane lipoprotein YidQ	243bp	81aa	98%	gi 291312035 EFE52488.1	6.00E-51
Mycobacterium virulence operon involved in protein synthesis (SSU ribosomal proteins)						
Rv0682	SSU ribosomal protein S12p (S23e)	375bp	125aa	100%	gi 490386408 WP_004265905.1	1E-84
Rv0683	SSU ribosomal protein S7p (S5e)	471bp	157aa	100%	gi 490382975 WP_004262486.1	4E-105
Rv0684	Translation elongation factor G	2127bp	709aa	100%	gi 490382972 WP_004262483.1	0.0

Rv0685	Translation elongation factor Tu	405bp	135aa	100%	gi 414098736 EKT60381.1	6E-92
Rv0685	Translation elongation factor Tu	138bp	46aa	98%	gi 391630004 EIS69838.1	2E-23
Rv0685	Translation elongation factor Tu	891bp	297aa	99%	gi 1042401681 OBY35966.1	0.0
Mycobacterium virulence operon involved in DNA transcription						
Rv0667	DNA-directed RNA polymerase beta subunit (EC 2.7.7.6)	4029bp	1343aa	100%	gi 490384883 WP_004264386.1	0.0
Rv0668	DNA-directed RNA polymerase beta' subunit (EC 2.7.7.6)	4224bp	1408aa	100%	gi 1042401688 OBY35973.1	0.0
Mycobacterium virulence operon possibly involved in quinolinate biosynthesis						
Rv1594	Quinolinate synthetase (EC 2.5.1.72)	1041bp	347aa	99%	gi 490377169 WP_004256768.1	0.0
Rv1595	L-aspartate oxidase (EC 1.4.3.16)	1596bp	532aa	99%	gi 739087991 WP_036959065.1	0.0
Rv1596	Quinolinate phosphoribosyltransferase [decarboxylating] (EC 2.4.2.19)	903bp	301aa	99%	gi 490382479 WP_004261992.1	0.0
Mycobacterium virulence operon involved in protein synthesis (LSU ribosomal proteins)						
Rv1641	Translation initiation factor 3	432bp	144aa	100%	gi 291311981 EFE52434.1	1E-95
Rv1641	Translation initiation factor 3	432bp	144aa	99%	gi 291311982 EFE52435.1	1E-94
Rv1642	LSU ribosomal protein L35p	198bp	66aa	100%	gi 490384195 WP_004263702.1	4E-40
Rv1643	LSU ribosomal protein L20p	357bp	119aa	100%	gi 490384208 WP_004263714.1	2E-75
Beta-lactamase						
BL	Beta-lactamase (EC 3.5.2.6)	861bp	287aa	100%	gi 1002048020 AMM70781.1	0.0
BL	Beta-lactamase (EC 3.5.2.6)	876bp	292aa	100%	gi 446804399 WP_000881655.1	0.0
BL	Beta-lactamase (EC 3.5.2.6)	1143bp	381aa	94%	gi 490383288 WP_004262799.1	0.0
bl	Beta-lactamase	813bp	271aa	100%	gi 926465406 ALD19783.1	0.0
Multiple Antibiotic Resistance MAR locus						
MarC	Multiple antibiotic resistance protein MarC	705bp	235aa	100%	gi 490384158 WP_004263665.1	2E-161
The mdtABCD multidrug resistance cluster						
BaeS	Sensory histidine kinase BaeS	1293bp	431aa	99%	gi 490373912 WP_004253515.1	0.0
BaeR	Response regulator BaeR	708bp	236aa	100%	gi 490373910 WP_004253513.1	8E-167

MdtA	Probable RND efflux membrane fusion protein	1245bp	415aa	99%	gi 490373920 WP_004253523.1	0.0
MdtA	Probable RND efflux membrane fusion protein	1152bp	384aa	99%	gi 489146363 WP_003056109.1	0.0
MdtB	Multidrug transporter MdtB	3102bp	1034aa	99%	gi 490373917 WP_004253520.1	0.0
MdtC	Multidrug transporter MdtC	3123bp	1041aa	99%	gi 490373914 WP_004253517.1	0.0
Multidrug Resistance Efflux Pumps						
CmeB	RND efflux system, inner membrane transporter CmeB	3159bp	1053aa	100%	gi 490380917 WP_004260435.1	0.0
*ToIC	RND efflux system, outer membrane lipoprotein CmeC	1419bp	473aa	99%	gi 414097180 EKT58835.1	0.0
*ToIC	RND efflux system, outer membrane lipoprotein CmeC	1389bp	463aa	100%	gi 739086918 WP_036958003.1	0.0
*ToIC	RND efflux system, outer membrane lipoprotein CmeC	1395bp	465aa	93%	gi 491044003 WP_004905665.1	0.0
*Reg	Transcription repressor of multidrug efflux pump acrAB operon, TetR (AcrR) family	651bp	217aa	99%	gi 490380911 WP_004260429.1	1E-150
*MATE_family_MDR_Pump	Multi antimicrobial extrusion protein (Na(+)/drug antiporter), MATE family of MDR efflux pumps	306bp	102aa	96%	gi 384481908 AFH95703.1	2E-39
*MATE_family_MDR_Pump	Multi antimicrobial extrusion protein (Na(+)/drug antiporter), MATE family of MDR efflux pumps	1029bp	343aa	99%	gi 491048771 WP_004910423.1	0.0
MFS	Multidrug-efflux transporter, major facilitator superfamily (MFS) (TC 2.A.1)	1227bp	409aa	99%	gi 490376269 WP_004255869.1	0.0
MacA	Macrolide-specific efflux protein MacA	1107bp	369aa	99%	gi 490376798 WP_004256397.1	0.0
MacA	Macrolide-specific efflux protein MacA	1182bp	394aa	100%	gi 414097178 EKT58833.1	0.0
MacB	Macrolide export ATP-binding/permease protein MacB (EC 3.6.3.-)	1944bp	648aa	99%	gi 490376796 WP_004256395.1	0.0
MacB	Macrolide export ATP-binding/permease protein MacB (EC 3.6.3.-)	1974bp	658aa	100%	gi 491046017 WP_004907673.1	0.0
MtrF	Multidrug efflux pump component MtrF	1572bp	524aa	99%	gi 490379478 WP_004258999.1	0.0
RND	Membrane fusion protein of RND family multidrug efflux pump	1188bp	396aa	100%	gi 490380914 WP_004260432.1	0.0

AcrB	RND multidrug efflux transporter, Acriflavin resistance protein	1095bp	365aa	77%	gi 739105285 WP_036975758.1	0.0
AcrB	RND multidrug efflux transporter, Acriflavin resistance protein	3105bp	1035aa	100%	gi 490381616 WP_004261131.1	0.0
Bile hydrolysis						
bsh	Choloylglycine hydrolase (EC 3.5.1.24)	1080bp	360aa	100%	gi 490382593 WP_004262105.1	0.0
DamX	DamX, an inner membrane protein involved in bile resistance	906bp	302aa	99%	gi 490383694 WP_004263203.1	0.0
Fosfomycin resistance						
FosA	Fosfomycin resistance protein FosA	414bp	138aa	100%	gi 490384164 WP_004263671.1	3E-98
Copper homeostasis						
*CIA	Lead, cadmium, zinc and mercury transporting ATPase (EC 3.6.3.3) (EC 3.6.3.5), Copper-translocating P-type ATPase (EC 3.6.3.4)	2925bp	975aa	98%	gi 739087196 WP_036958281.1	0.0
*CIA	Lead, cadmium, zinc and mercury transporting ATPase (EC 3.6.3.3) (EC 3.6.3.5), Copper-translocating P-type ATPase (EC 3.6.3.4)	2400bp	800aa	99%	gi 1042403368 OBY37647.1	0.0
BCO	Blue copper oxidase CueO precursor	1617bp	539aa	99%	gi 490381034 WP_004260551.1	0.0
*HL	Cytochrome c heme lyase subunit CcmF	1932bp	644aa	99%	gi 491051693 WP_004913344.1	0.0
*HL	Cytochrome c heme lyase subunit CcmL, Cytochrome c heme lyase subunit CcmH	1071bp	357aa	99%	gi 490382689 WP_004262201.1	0.0
CRD	Copper resistance protein D	897bp	299aa	99%	gi 490374296 WP_004253898.1	0.0
CopG	CopG protein	423bp	141aa	98%	gi 491043999 WP_004905661.1	2E-97
copCp	Copper resistance protein C precursor	387bp	129aa	100%	gi 490374293 WP_004253895.1	7E-86
Streptothricin resistance						
SatA	Streptothricin acetyltransferase, Streptomyces lavendulae type	525bp	175aa	100%	gi 446626810 WP_000704156.1	1E-125
Adaptation to d-cysteine						
DcyD	D-cysteine desulfhydrase (EC 4.4.1.15)	990bp	330aa	99%	gi 490374828 WP_004254429.1	0.0

Mercuric reductase						
MIR	Mercuric ion reductase (EC 1.16.1.1)	1686bp	562aa	100%	gi 446131441 WP_000209296.1	0.0
Resistance to fluoroquinolones						
parC	Topoisomerase IV subunit A (EC 5.99.1.-)	2256bp	752aa	99%	gi 490378099 WP_004257697.1	0.0
parE	Topoisomerase IV subunit B (EC 5.99.1.-)	1896bp	632aa	99%	gi 490378103 WP_004257701.1	0.0
gyrA	DNA gyrase subunit A (EC 5.99.1.3)	2616bp	872aa	99%	gi 491050539 WP_004912190.1	0.0
gyrB	DNA gyrase subunit B (EC 5.99.1.3)	2415bp	805aa	99%	gi 491044587 WP_004906247.1	0.0
Cobalt-zinc-cadmium resistance						
*CzcD	Cobalt-zinc-cadmium resistance protein	930bp	310aa	100%	gi 490382252 WP_004261765.1	0.0
*CzcA?	Cobalt-zinc-cadmium resistance protein	3078bp	1026aa	99%	gi 491044000 WP_004905662.1	0.0
*CusB/CzsB	CzcA, Cation efflux system protein CusA Probable Co/Zn/Cd efflux system membrane fusion protein	1095bp	365aa	62%	gi 1093685069 WP_070928012.1	6E-144
*CusB/CzsB	Probable Co/Zn/Cd efflux system membrane fusion protein	1110bp	370aa	99%	gi 490381614 WP_004261129.1	0.0
*CusB/CzsB	Cobalt/zinc/cadmium efflux RND transporter, membrane fusion protein, CzcB family	1269bp	423aa	99%	gi 491044001 WP_004905663.1	0.0
*CusA	Cobalt-zinc-cadmium resistance protein CzcA, Cation efflux system protein CusA	3078bp	1026aa	99%	gi 491044000 WP_004905662.1	0.0
*CzrR	DNA-binding heavy metal response regulator	684bp	228aa	97%	gi 491044004 WP_004905666.1	1E-153
cusF	Cation efflux system protein CusF precursor	342bp	114aa	97%	gi 491044002 WP_004905664.1	4E-74
*TR	Transcriptional regulator, MerR family	423bp	141aa	99%	gi 490374914 WP_004254515.1	2E-92
Mercury resistance operon						
MerR	Mercuric resistance operon regulatory protein	456bp	152aa	100%	gi 447089372 WP_001166628.1	4E-103
MerT	Mercuric transport protein, MerT	366bp	122aa	100%	gi 447217400 WP_001294656.1	8E-84
MerC	Mercuric transport protein, MerC	426bp	142aa	100%	gi 446445141 WP_000522996.1	3E-98
MerA	Mercuric ion reductase (EC 1.16.1.1)	1686bp	562aa	100%	gi 446131441 WP_000209296.1	0.0

MerD	Mercuric resistance operon coregulator	246bp	82aa	99%	gi 585339711 WP_024221342.1	2E-47
Copper homeostasis: copper tolerance						
ScsB	Membrane protein, suppressor for copper-sensitivity ScsB	2073bp	691aa	99%	gi 490377422 WP_004257021.1	0.0
ScsC	Secreted protein, suppressor for copper-sensitivity ScsC	732bp	244aa	99%	gi 490377417 WP_004257016.1	7E-169
ScsD	Membrane protein, suppressor for copper-sensitivity ScsD	501bp	167aa	100%	gi 491047306 WP_004908960.1	5E-115
CutF	Copper homeostasis protein CutF precursor, Lipoprotein NlpE involved in surface adhesion	657bp	219aa	99%	gi 490377815 WP_004257413.1	3E-156
CutE	Apolipoprotein N-acyltransferase (EC 2.3.1.-), Copper homeostasis protein CutE	1530bp	510aa	99%	gi 490377396 WP_004256995.1	0.0
CorC	Magnesium and cobalt efflux protein CorC	1287bp	429aa	99%	gi 1042400154 OBY34443.1	0.0
CorC	Magnesium and cobalt efflux protein CorC	909bp	303aa	99%	gi 291314305 EFE54758.1	0.0
CorC	Magnesium and cobalt efflux protein CorC	1368bp	456aa	100%	gi 491044300 WP_004905961.1	0.0
Aminoglycoside adenylyltransferases						
AadA1	Streptomycin 3"-O-adenylyltransferase (EC 2.7.7.47), Spectinomycin 9-O-adenylyltransferase	159bp	53aa	100%	gi 487942357 WP_002015823.1	3E-31
AadA1	Streptomycin 3"-O-adenylyltransferase (EC 2.7.7.47), Spectinomycin 9-O-adenylyltransferase	1014bp	338aa	100%	gi 678222195 AIM49564.1	0.0
AadA1	Streptomycin 3"-O-adenylyltransferase (EC 2.7.7.47), Spectinomycin 9-O-adenylyltransferase	792bp	264aa	100%	gi 501453940 WP_012477385.1	0.0
AadA1	Streptomycin 3"-O-adenylyltransferase (EC 2.7.7.47), Spectinomycin 9-O-adenylyltransferase	150bp	50aa	100%	gi 486167809 WP_001531151.1	5E-29
AadA1	Streptomycin 3"-O-adenylyltransferase (EC 2.7.7.47), Spectinomycin 9-O-adenylyltransferase	789bp	263aa	100%	gi 447129059 WP_001206315.1	0.0
AadA2	Spectinomycin 9-O-adenylyltransferase	765bp	255aa	99%	gi 490381153 WP_004260670.1	2E-178

AadA2	Spectinomycin 9-O-adenylyltransferase	159bp	53aa	100%	gi 487942357 WP_002015823.1	3E-31
AadA2	Spectinomycin 9-O-adenylyltransferase	1014bp	338aa	100%	gi 678222195 AIM49564.1	0.0
AadA2	Spectinomycin 9-O-adenylyltransferase	792bp	264aa	100%	gi 501453940 WP_012477385.1	0.0
AadA2	Spectinomycin 9-O-adenylyltransferase	150bp	50aa	100%	gi 486167809 WP_001531151.1	5E-29
AadA2	Spectinomycin 9-O-adenylyltransferase	789bp	263aa	100%	gi 447129059 WP_001206315.1	0.0
Arsenic resistance						
arsR	Arsenical resistance operon repressor	336bp	112aa	100%	gi 757599390 WP_042847035.1	4E-73
arsR	Arsenical resistance operon repressor	336bp	112aa	99%	gi 739097345 WP_036968128.1	6E-72
arsR	Arsenical resistance operon repressor	189bp	63aa	92%	gi 1116088436 WP_072144547.1	2E-33
arsD	Arsenical resistance operon trans-acting repressor ArsD	366bp	122aa	100%	gi 757599389 WP_042847034.1	2E-85
arsD	Arsenical resistance operon trans-acting repressor ArsD	366bp	122aa	99%	gi 739097347 WP_036968130.1	2E-84
arsA	Arsenical pump-driving ATPase (EC 3.6.3.16)	1752bp	584aa	99%	gi 827610366 KLO01850.1	0.0
arsA	Arsenical pump-driving ATPase (EC 3.6.3.16)	1572bp	584aa	99%	gi 739097348 WP_036968131.1	0.0
arsB	Arsenic efflux pump protein	1290bp	430aa	98%	gi 490357905 WP_004237678.1	0.0
arsB	Arsenic efflux pump protein	1290bp	430aa	99%	gi 739104700 WP_036975217.1	0.0
arsB	Arsenic efflux pump protein	1362bp	454aa	98%	gi 739104700 WP_036975217.1	0.0
arsC	Arsenate reductase (EC 1.20.4.1)	357bp	119aa	99%	gi 490373943 WP_004253545.1	1E-78
arsC	Arsenate reductase (EC 1.20.4.1)	417bp	139aa	91%	gi 749308148 WP_040132278.1	8E-88
arsC	Arsenate reductase (EC 1.20.4.1)	441bp	147aa	100%	gi 739097354 WP_036968137.1	2E-100

Iron acquisition and metabolism

Sigla	Gene	Tamaño	Aminoácidos	Identidad	Secuenciamento	E. value
Siderophore Aerobactin						
iutA	Aerobactin siderophore receptor iutA TonB-dependent siderophore receptor	2202bp	734aa	99%	gi 490378242 WP_004257840.1	0.0
Heme, hemin uptake and utilization systems in GramPositives						
*Heme_oxygenases	Hemin transport protein HmuS	1050bp	350aa	99%	gi 491051296 WP_004912947.1	0.0
*Heme_oxygenases	Hemin transport protein HmuS	1050bp	350aa	99%	gi 739087580 WP_036958663.1	0.0
Ferrous iron transporter EfeUOB, low-pH-induced						
*EfeU	Ferrous iron transport permease EfeU	834bp	278aa	99%	gi 490381107 WP_004260624.1	0.0
EfeB	Ferrous iron transport peroxidase EfeB	1275bp	425aa	99%	gi 490381112 WP_004260629.1	0.0
Encapsulating protein for DyP-type peroxidase and ferritin-like protein oligomers						
DyP	Predicted dye-decolorizing peroxidase (DyP), encapsulated subgroup	900bp	300aa	99%	gi 490385387 WP_004264885.1	0.0
Yfe	Predicted outer membrane lipoprotein YfeY	606bp	202aa	99%	gi 915327114 WP_050763802.1	4E-145
Hemin transport system						
HBP	Periplasmic hemin-binding protein	822bp	274aa	99%	gi 491051298 WP_004912949.1	0.0
*Rec	TonB-dependent hemin , ferrichrome receptor	2019bp	673aa	91%	gi 739087359 WP_036958444.1	0.0
*Rec	TonB-dependent hemin , ferrichrome receptor	2145bp	715aa	99%	gi 490384250 WP_004263756.1	0.0
hmuS	Hemin transport protein HmuS	1050bp	350aa	99%	gi 491051296 WP_004912947.1	0.0
hmuS	Hemin transport protein HmuS	1050bp	350aa	99%	gi 739087580 WP_036958663.1	0.0
ATPb	ABC-type hemin transport system, ATPase component	789bp	263aa	98%	gi 1172252176 WP_080641516.1	0.0

ATPb	ABC-type hemin transport system, ATPase component	768bp	256aa	99%	gi 1042402961 OBY37241.1	2E-178
PP_7	Hemin ABC transporter, permease protein	1002bp	334aa	99%	gi 491051302 WP_004912953.1	0.0
PP_7	Hemin ABC transporter, permease protein	1017bp	339aa	100%	gi 1042402962 OBY37242.1	0.0
TonB	Ferric siderophore transport system, periplasmic binding protein TonB	744bp	248aa	97%	gi 490382765 WP_004262277.1	1E-162
TonB	Ferric siderophore transport system, periplasmic binding protein TonB	966bp	322aa	98%	gi 490378848 WP_004258445.1	0.0
TonB	Ferric siderophore transport system, periplasmic binding protein TonB	720bp	240aa	99%	gi 739086610 WP_036957695.1	6E-173
TonB	Ferric siderophore transport system, periplasmic binding protein TonB	783bp	261aa	99%	gi 739087479 WP_036958564.1	3E-179
TonB	Ferric siderophore transport system, periplasmic binding protein TonB	756bp	252aa	97%	gi 490373999 WP_004253601.1	1E-165
Heme, hemin uptake and utilization systems in GramNegatives						
HBP	Periplasmic hemin-binding protein	822bp	274aa	99%	gi 491051298 WP_004912949.1	0.0
fhuC	Ferrichrome transport ATP-binding protein FhuC (TC 3.A.1.14.3)	783bp	261aa	100%	gi 490382157 WP_004261670.1	0.0
fhuC	Ferrichrome transport ATP-binding protein FhuC (TC 3.A.1.14.3)	786bp	262aa	100%	gi 757595221 WP_042843418.1	8E-180
FCR	TonB-dependent hemin , ferrichrome receptor	2019bp	673aa	91%	gi 739087359 WP_036958444.1	0.0
FCR	TonB-dependent hemin , ferrichrome receptor	2145bp	715aa	99%	gi 490384250 WP_004263756.1	0.0
FR	Flavin reductase (EC 1.5.1.30)	624bp	208aa	99%	gi 739087413 WP_036958498.1	7E-143
ETFb	Electron transfer flavoprotein, beta subunit	756bp	252aa	99%	gi 490378882 WP_004258479.1	6E-168
ParA	Paraquat-inducible protein A	1227bp	409aa	99%	gi 1102614758 WP_071585858.1	0.0
ParA	Paraquat-inducible protein A	1227bp	409aa	99%	gi 739086601 WP_036957686.1	0.0
ParB	Paraquat-inducible protein B	1650bp	550aa	99%	gi 490376536 WP_004256135.1	0.0

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ParB	Paraquat-inducible protein B	2340bp	780aa	99%	gi 490374304 WP_004253906.1	0.0
hmuS	Hemin transport protein HmuS	1050bp	350aa	99%	gi 491051296 WP_004912947.1	0.0
hmuS	Hemin transport protein HmuS	1050bp	350aa	99%	gi 739087580 WP_036958663.1	0.0
PP_7	Hemin ABC transporter, permease protein	1002bp	334aa	99%	gi 491051302 WP_004912953.1	0.0
PP_7	Hemin ABC transporter, permease protein	1017bp	339aa	100%	gi 1042402962 OBY37242.1	0.0
ATPb	ABC-type hemin transport system, ATPase component	789bp	263aa	98%	gi 1172252176 WP_08064151.6.1	0.0
ATPb	ABC-type hemin transport system, ATPase component	768bp	256aa	99%	gi 1042402961 OBY37241.1	2E-178
TonB	Ferric siderophore transport system, periplasmic binding protein TonB	744bp	248aa	97%	gi 490382765 WP_004262277.1	1E-162
TonB	Ferric siderophore transport system, periplasmic binding protein TonB	966bp	322aa	98%	gi 490378848 WP_004258445.1	0.0
TonB	Ferric siderophore transport system, periplasmic binding protein TonB	720bp	240aa	99%	gi 739086610 WP_036957695.1	6E-173
TonB	Ferric siderophore transport system, periplasmic binding protein TonB	783bp	261aa	99%	gi 739087479 WP_036958564.1	3E-179
TonB	Ferric siderophore transport system, periplasmic binding protein TonB	756bp	252aa	97%	gi 490373999 WP_004253601.1	1E-165

Phages, Prophages

Sigla	Gene	Tamanho	Aminoácido S	Identidade	Sequenciamento	E. value
Transposable elements - CBSS-203122.12.peg.188						
R1	TniB NTP-binding protein	510bp	170aa	100%	gi 213054790 ACJ39692.1	2E-120
R8	MG(2+) CHELATASE FAMILY PROTEIN ComM-related protein	1527bp	509aa	99%	gi 491044471 WP_004906131. 1	0.0
R9	MG(2+) CHELATASE FAMILY PROTEIN ComM-related protein	1527bp	509aa	99%	gi 491044471 WP_004906131. 1	0.0
Phages, Prophages, Transposable elements, Plasmids - Integrans						
IntI2	Integron integrase IntI2	537bp	179aa	100%	gi 445994041 WP_000071896. 1	8E-125
IntI2	Integron integrase IntI2	537bp	179aa	100%	gi 446419664 WP_000497519. 1	9E-75
IntIPac	Integron integrase IntIPac	867bp	289aa	100%	gi 487966541 WP_002039642. 1	0.0
Phages, Prophages - Phage tail proteins						
TP	Phage tail protein	444bp	148aa	100%	gi 1101502596 APC13989.1	2E-102
*Measure	Phage tail length tape-measure protein	2700bp	900aa	9800%	gi 873900395 WP_048606828. 1	0.0
*Tube	Phage major tail tube protein	516bp	172aa	100%	gi 1101998870 WP_07154873 1.1	8E-120
*Sheath	Phage tail sheath monomer	1530bp	510aa	100%	gi 1101998871 WP_07154873 2.1	0.0
Phages, Prophages - Phage replication						
*Rep	Phage replication protein	2424bp	808aa	93%	gi 1101998888 WP_07154874 9.1	0.0
*dPol_3	DNA polymerase III alpha subunit (EC 2.7.7.7)	3483bp	1161aa	100%	gi 739086905 WP_036957990. 1	0.0
Phages, Prophages - Phage tail proteins 2						
TMP1_1	Phage tape measure	447bp	149aa	99%	gi 873900393 WP_048606826. 1	1E-99
TSM	Phage tail sheath monomer	1530bp	510aa	100%	gi 1101998871 WP_07154873	0.0

TMP	Phage tail length tape-measure protein	2700bp	900aa	98%	2.1 gij 873900395 WP_048606828.1	0.0
Phages, Prophages Phage tail fiber proteins						
TFP	Phage tail fiber protein	1530bp	510aa	99%	gij 1101502589 APC13982.1	0.0
TFAP	Phage tail fiber assembly protein	645bp	215aa	100%	gij 1101502588 APC13981.1	2E-149
TFP2	Phage tail fibers	612bp	204aa	100%	gij 873900403 WP_048606836.1	2E-146
Phages, Prophages Phage capsid proteins						
*Generic	Phage capsid and scaffold	444bp	148aa	100%	gij 739091104 WP_036962066.1	3.00E-108
MajCap	Phage major capsid protein	1095bp	365aa	99%	gij 1101998881 WP_07154874.2.1	0.0
*Scaffolding	Phage capsid scaffolding protein	828bp	276aa	99%	gij 1101998882 WP_07154874.3.1	0.0
HCSP	Phage head completion-stabilization protein	477bp	159aa	100%	gij 1101998879 WP_07154874.0.1	4.0E-108

Motility and Chemotaxis

Sigla	Gene	Tamanho	Aminoácidos	Identidade	Sequenciamento	E. value
Bacterial Chemotaxis						
aer	Aerotaxis sensor receptor protein	1533bp	511aa	98%	gi 739086922 WP_036958007.1	0.0
tsr	Methyl-accepting chemotaxis protein I (serine chemoreceptor protein)	1653bp	551aa	98%	gi 490373823 WP_004253426.1	0.0
tsr	Methyl-accepting chemotaxis protein I (serine chemoreceptor protein)	1569bp	523aa	99%	gi 490376118 WP_004255718.1	0.0
tsr	Methyl-accepting chemotaxis protein I (serine chemoreceptor protein)	1602bp	534aa	99%	gi 490376121 WP_004255721.1	0.0
tsr	Methyl-accepting chemotaxis protein I (serine chemoreceptor protein)	1488bp	496aa	99%	gi 490378843 WP_004258440.1	0.0
*cheW-V	Positive regulator of CheA protein activity (CheW)	498bp	166aa	100%	gi 491048180 WP_004909833.1	3E-111
cheA	Signal transduction histidine kinase CheA (EC 2.7.3.-)	2058bp	686aa	99%	gi 739086771 WP_036957856.1	0.0
cheY	Chemotaxis regulator - transmits chemoreceptor signals to flagellar motor components CheY	393bp	131aa	100%	gi 490376105 WP_004255705.1	7E-85
cheZ	Chemotaxis response - phosphatase CheZ	639bp	213aa	99%	gi 104239971 OBY34011.1	8E-142
cheB	Chemotaxis response regulator protein-glutamate methylesterase CheB (EC 3.1.1.61)	1068bp	356aa	99%	gi 490376109 WP_004255709.1	0.0
fliG	Flagellar motor switch protein FliG	993bp	331aa	100%	gi 490376000 WP_004255600.1	0.0
fliM	Flagellar motor switch protein FliM	1008bp	336aa	99%	gi 490376019 WP_004255619.1	0.0
fliN	Flagellar motor switch protein FliN	417bp	139aa	100%	gi 490376021 WP_004255621.1	1E-90
malE	Maltose/maltodextrin ABC transporter, substrate binding periplasmic protein MalE	1191bp	397aa	92%	gi 1057035456 WP_068445917.1	0.0
dppA	Dipeptide-binding ABC transporter, periplasmic substrate-binding component (TC 3.A.1.5.2)	1608bp	536aa	100%	gi 490383128 WP_004262639.1	0.0
dppA	Dipeptide-binding ABC transporter, periplasmic substrate-binding component (TC 3.A.1.5.2)	1560bp	520aa	99%	gi 739086975 WP_036958060.1	0.0
cheR	Chemotaxis protein methyltransferase CheR (EC 2.1.1.80)	837bp	279aa	100%	gi 490376113 WP_004255713.1	0.0

Flagellum						
flgB	Flagellar basal-body rod protein FlgB	414bp	138aa	100%	gi 490376069 WP_004255669.1	2E-91
flgC	Flagellar basal-body rod protein FlgC	405bp	135aa	100%	gi 491048215 WP_004909868.1	3E-89
FlgD	Flagellar basal-body rod modification protein FlgD	858bp	286aa	99%	gi 490376064 WP_004255664.1	0.0
flgE	Flagellar hook protein FlgE	1221bp	407aa	96%	gi 873900357 WP_048606790.1	0.0
flgF	Flagellar basal-body rod protein FlgF	756bp	252aa	100%	gi 490376059 WP_004255659.1	2E-174
flgG	Flagellar basal-body rod protein FlgG	783bp	261aa	99%	gi 490376057 WP_004255657.1	0.0
FlgH	Flagellar L-ring protein FlgH	750bp	250aa	100%	gi 291314697 EFE55150.1	0.0
flgI	Flagellar P-ring protein FlgI	1104bp	368aa	100%	gi 490376051 WP_004255651.1	0.0
flgJ	Flagellar protein FlgJ [peptidoglycan hydrolase] (EC 3.2.1.-)	981bp	327aa	98%	gi 490376048 WP_004255648.1	0.0
flgK	Flagellar hook-associated protein FlgK	1644bp	548aa	99%	gi 739086767 WP_036957852.1	0.0
flgL	Flagellar hook-associated protein FlgL	936bp	312aa	98%	gi 491048236 WP_004909889.1	0.0
FliA	Flagellar biosynthesis protein FliA	2100bp	700aa	100%	gi 490376087 WP_004255687.1	0.0
fliE	Flagellar hook-basal body complex protein FliE	312bp	104aa	100%	gi 490375994 WP_004255594.1	4E-64
fliF	Flagellar M-ring protein FliF	1686bp	562aa	99%	gi 490375997 WP_004255597.1	0.0
fliG	Flagellar motor switch protein FliG	993bp	331aa	100%	gi 490376000 WP_004255600.1	0.0
fliH	Flagellar assembly protein FliH	729bp	243aa	99%	gi 490376003 WP_004255603.1	8E-177
FliI	Flagellum-specific ATP synthase FliI	1311bp	437aa	99%	gi 490376006 WP_004255606.1	0.0
fliK	Flagellar hook-length control protein FliK	1368bp	456aa	97%	gi 490376013 WP_004255613.1	0.0
fliJ	Flagellar protein FliJ	444bp	148aa	99%	gi 490376009 WP_004255609.1	3E-100
fliM	Flagellar motor switch protein FliM	1008bp	336aa	99%	gi 490376019 WP_004255619.1	0.0
fliN	Flagellar motor switch protein FliN	417bp	139aa	100%	gi 490376021 WP_004255621.1	1E-90
fliP	Flagellar biosynthesis protein FliP	744bp	248aa	99%	gi 739086765 WP_036957850.1	3E-167
fliQ	Flagellar biosynthesis protein FliQ	270bp	90aa	99%	gi 490376030 WP_004255630.1	2E-52
FliR	Flagellar biosynthesis protein FliR	780bp	260aa	99%	gi 490376033 WP_004255633.1	7E-180
*Flagellar-Regulation	RNA polymerase sigma factor RpoD	1854bp	618aa	100%	gi 1102614763 WP_071585863.1	0.0

*Flagellar-Regulation	Flagellar transcriptional activator FlhC	591bp	197aa	100%	gi 490376132 WP_004255732.1	1E-140
*Flagellar-Regulation	Flagellar biosynthesis protein FliZ	519bp	173aa	100%	gi 490376157 WP_004255757.1	4E-123
*Flagellar-Regulation	RNA polymerase sigma-54 factor RpoN	1446bp	482aa	99%	gi 490378708 WP_004258305.1	0.0
*Flagellar-Regulation	RNA polymerase sigma factor for flagellar operon	726bp	242aa	99%	gi 291314661 EFE55114.1	0.0
*Flagellin	Flagellar biosynthesis protein FliC	1170bp	390aa	87%	gi 1055836465 WP_067424616.1	0.0
fliD	Flagellar hook-associated protein FliD	1437bp	479aa	90%	gi 1057031755 WP_068442704.1	0.0
fliS	Flagellar biosynthesis protein FliS	399bp	133aa	100%	gi 491048316 WP_004909969.1	1E-86
fliT	Flagellar biosynthesis protein FliT	390bp	130aa	99%	gi 490375964 WP_004255564.1	2E-85
*fliO	Flagellar biosynthesis protein FliO	459bp	153aa	99%	gi 490376024 WP_004255624.1	7E-102
FlhB	Flagellar biosynthesis protein FlhB	1152bp	384aa	99%	gi 490376088 WP_004255688.1	0.0
MotA	Flagellar motor rotation protein MotA	954bp	318aa	99%	gi 291314724 EFE55177.1	0.0
MotB	Flagellar motor rotation protein MotB	918bp	306aa	99%	gi 490376126 WP_004255726.1	0.0
*flhD	Flagellar transcriptional activator FlhD	351bp	117aa	100%	gi 739086775 WP_036957860.1	3E-74
fliL	Flagellar biosynthesis protein FliL	480bp	160aa	99%	gi 490376015 WP_004255615.1	3E-107
Flagellar motility						
FlhA	Flagellar biosynthesis protein FlhA	2100bp	700aa	100%	gi 490376087 WP_004255687.1	0.0
fliM	Flagellar motor switch protein FliM	1008bp	336aa	99%	gi 490376019 WP_004255619.1	0.0
fliN	Flagellar motor switch protein FliN	417bp	139aa	100%	gi 490376021 WP_004255621.1	1E-90
cheY	Chemotaxis regulator - transmits chemoreceptor signals to flagellar motor components CheY	393bp	131aa	100%	gi 490376105 WP_004255705.1	7E-85
cheA	Signal transduction histidine kinase CheA (EC 2.7.3.-)	2058bp	686aa	99%	gi 739086771 WP_036957856.1	0.0
FlhB	Flagellar biosynthesis protein FlhB	1152bp	384aa	99%	gi 490376088 WP_004255688.1	0.0
FliI	Flagellum-specific ATP synthase FliI	1311bp	437aa	99%	gi 490376006 WP_004255606.1	0.0
FliR	Flagellar biosynthesis protein FliR	780bp	260aa	99%	gi 490376033 WP_004255633.1	7E-180

FlgD	Flagellar basal-body rod modification protein FlgD	858bp	286aa	99%	gi 490376064 WP_004255664.1	0.0
FlgH	Flagellar L-ring protein FlgH	750bp	250aa	100%	gi 291314697 EFE55150.1	0.0
MotA	Flagellar motor rotation protein MotA	954bp	318aa	99%	gi 291314724 EFE55177.1	0.0
MotB	Flagellar motor rotation protein MotB	918bp	306aa	99%	gi 490376126 WP_004255726.1	0.0
FliA	RNA polymerase sigma factor for flagellar operon	726bp	242aa	99%	gi 291314661 EFE55114.1	9E-167
RpoN	RNA polymerase sigma-54 factor RpoN	1446bp	482aa	99%	gi 490378708 WP_004258305.1	0.0

Sub-sistemas PR02

Virulence, Disease and Defense

Sigla	Gene	Tamanho	Aminoácidos	Identidade	Sequenciamento	E. value
Tolerance to colicin E2						
CbrC-like	Colicin E2 tolerance protein CbrC-like protein	588bp	196aa	99%	gi 490381407 WP_004260923.1	8.00E-151
CreA	Conserved uncharacterized protein CreA	477bp	159aa	100%	gi 491044024 WP_004905686.1	2.00E-108
Colicin V and Bacteriocin Production Cluster						
R1	tRNA pseudouridine synthase A (EC 4.2.1.70)	816bp	272aa	100%	gi 490377681 WP_004257279.1	0.0
DedA	DedA protein	687bp	229aa	99%	gi 490377686 WP_004257284.1	5E-159
R3	Acetyl-coenzyme A carboxyl transferase beta chain (EC 6.4.1.2)	972bp	324aa	99%	gi 490377689 WP_004257287.1	0.0
R4	Dihydrofolate synthase (EC 6.3.2.12), Folylpolyglutamate synthase (EC 6.3.2.17)	1287bp	429aa	99%	gi 490377692 WP_004257290.1	0.0
R5	Dihydrofolate synthase (EC 6.3.2.12), Folylpolyglutamate synthase (EC 6.3.2.17)	1287bp	429aa	99%	gi 490377692 WP_004257290.1	0.0
DedD	DedD protein	642bp	214aa	100%	gi 739086890 WP_036957975.1	3E-147
*toxin	Colicin V production protein	504bp	168aa	100%	gi 490377706 WP_004257304.1	3E-115
PurF	Amidophosphoribosyltransferase (EC 2.4.2.14)	1518bp	506aa	100%	gi 490377709 WP_004257307.1	0.0
Mediator of hyperadherence YidE in Enterobacteria and its conserved region						
YidE	Mediator of hyperadherence YidE	1659bp	553aa	99%	gi 490383146 WP_004262657.1	0.0
YidR	Uncharacterized protein YidR	1317bp	439aa	99%	gi 291312036 EFE52489.1	0.0
YidQ	Outer membrane lipoprotein YidQ	243bp	81aa	98%	gi 291312035 EFE52488.1	6.00E-51
Mycobacterium virulence operon involved in protein synthesis (SSU ribosomal proteins)						
Rv0682	SSU ribosomal protein S12p (S23e)	375bp	125aa	100%	gi 490386408 WP_004265905.1	1E-84
Rv0683	SSU ribosomal protein S7p (S5e)	471bp	157aa	100%	gi 490382975 WP_004262486.1	4E-105

Rv0684	Translation elongation factor G	2127bp	709aa	100%	gi 490382972 WP_004262483.1	0.0
Rv0685	Translation elongation factor Tu	1185bp	395aa	99%	gi 490384870 WP_004264373.1	0.0
Rv0685	Translation elongation factor Tu	138bp	46aa	98%	gi 391630004 EIS69838.1	2E-23
Mycobacterium virulence operon involved in DNA transcription						
Rv0667	DNA-directed RNA polymerase beta subunit (EC 2.7.7.6)	4029bp	1343aa	100%	gi 490384883 WP_004264386.1	0.0
Rv0668	DNA-directed RNA polymerase beta' subunit (EC 2.7.7.6)	4224bp	1408aa	100%	gi 1042401688 OBY35973.1	0.0
Mycobacterium virulence operon possibly involved in quinolinate biosynthesis						
Rv1594	Quinolinate synthetase (EC 2.5.1.72)	1041bp	347aa	99%	gi 490377169 WP_004256768.1	0.0
Rv1595	L-aspartate oxidase (EC 1.4.3.16)	1596bp	532aa	99%	gi 739087991 WP_036959065.1	0.0
Rv1596	Quinolinate phosphoribosyltransferase [decarboxylating] (EC 2.4.2.19)	903bp	301aa	99%	gi 490382479 WP_004261992.1	0.0
Mycobacterium virulence operon involved in protein synthesis (LSU ribosomal proteins)						
Rv1641	Translation initiation factor 3	432bp	144aa	100%	gi 291311981 EFE52434.1	1E-95
Rv1641	Translation initiation factor 3	432bp	144aa	99%	gi 291311982 EFE52435.1	1E-94
Rv1642	LSU ribosomal protein L35p	198bp	66aa	100%	gi 490384195 WP_004263702.1	4E-40
Rv1643	LSU ribosomal protein L20p	357bp	119aa	100%	gi 490384208 WP_004263714.1	2E-75
Beta-lactamase						
BL	Beta-lactamase (EC 3.5.2.6)	861bp	287aa	100%	gi 1002048020 AMM70781.1	0.0
BL	Beta-lactamase (EC 3.5.2.6)	876bp	292aa	100%	gi 446804399 WP_000881655.1	0.0
BL	Beta-lactamase (EC 3.5.2.6)	1143bp	381aa	94%	gi 490383288 WP_004262799.1	0.0
bl	Beta-lactamase	813bp	271aa	100%	gi 926465406 ALD19783.1	0.0
Multiple Antibiotic Resistance MAR locus						
MarC	Multiple antibiotic resistance protein MarC	705bp	235aa	100%	gi 490384158 WP_004263665.1	2E-161
The mdtABCD multidrug resistance cluster						
BaeS	Sensory histidine kinase BaeS	1293bp	431aa	99%	gi 490373912 WP_004253515.1	0.0
BaeR	Response regulator BaeR	708bp	236aa	100%	gi 490373910 WP_004253513.1	8E-167

MdtA	Probable RND efflux membrane fusion protein	1245bp	415aa	99%	gi 490373920 WP_004253523.1	0.0
MdtA	Probable RND efflux membrane fusion protein	1152bp	384aa	99%	gi 489146363 WP_003056109.1	0.0
MdtB	Multidrug transporter MdtB	3102bp	1034aa	99%	gi 490373917 WP_004253520.1	0.0
MdtC	Multidrug transporter MdtC	3123bp	1041aa	99%	gi 490373914 WP_004253517.1	0.0
Multidrug Resistance Efflux Pumps						
CmeB	RND efflux system, inner membrane transporter CmeB	3159bp	1053aa	100%	gi 490380917 WP_004260435.1	0.0
*TolC	RND efflux system, outer membrane lipoprotein CmeC	1419bp	473aa	99%	gi 414097180 EKT58835.1	0.0
*TolC	RND efflux system, outer membrane lipoprotein CmeC	1389bp	463aa	100%	gi 739086918 WP_036958003.1	0.0
*TolC	RND efflux system, outer membrane lipoprotein CmeC	1395bp	465aa	93%	gi 491044003 WP_004905665.1	0.0
*Reg	Transcription repressor of multidrug efflux pump acrAB operon, TetR (AcrR) family	651bp	217aa	99%	gi 490380911 WP_004260429.1	1E-150
*MATE_family_MDR_Pump	Multi antimicrobial extrusion protein (Na(+)/drug antiporter), MATE family of MDR efflux pumps	306bp	102aa	96%	gi 384481908 AFH95703.1	2E-39
*MATE_family_MDR_Pump	Multi antimicrobial extrusion protein (Na(+)/drug antiporter), MATE family of MDR efflux pumps	1029bp	343aa	99%	gi 491048771 WP_004910423.1	0.0
MFS	Multidrug-efflux transporter, major facilitator superfamily (MFS) (TC 2.A.1)	1227bp	409aa	99%	gi 490376269 WP_004255869.1	0.0
MacA	Macrolide-specific efflux protein MacA	1107bp	369aa	99%	gi 490376798 WP_004256397.1	0.0
MacA	Macrolide-specific efflux protein MacA	1182bp	394aa	100%	gi 414097178 EKT58833.1	0.0
MacB	Macrolide export ATP-binding/permease protein MacB (EC 3.6.3.-)	1944bp	648aa	99%	gi 490376796 WP_004256395.1	0.0
MacB	Macrolide export ATP-binding/permease protein MacB (EC 3.6.3.-)	1974bp	658aa	100%	gi 491046017 WP_004907673.1	0.0
MtrF	Multidrug efflux pump component MtrF	1572bp	524aa	99%	gi 490379478 WP_004258999.1	0.0
RND	Membrane fusion protein of RND family multidrug efflux pump	1188bp	396aa	100%	gi 490380914 WP_004260432.1	0.0

AcrB	RND multidrug efflux transporter, Acriflavin resistance protein	1095bp	365aa	77%	gi 739105285 WP_036975758.1	0.0
AcrB	RND multidrug efflux transporter, Acriflavin resistance protein	3105bp	1035aa	100%	gi 490381616 WP_004261131.1	0.0
Bile hydrolysis						
bsh	Choloylglycine hydrolase (EC 3.5.1.24)	1080bp	360aa	100%	gi 490382593 WP_004262105.1	0.0
DamX	DamX, an inner membrane protein involved in bile resistance	906bp	302aa	99%	gi 490383694 WP_004263203.1	0.0
Fosfomycin resistance						
FosA	Fosfomycin resistance protein FosA	414bp	138aa	100%	gi 490384164 WP_004263671.1	3E-98
Copper homeostasis						
*CIA	Lead, cadmium, zinc and mercury transporting ATPase (EC 3.6.3.3) (EC 3.6.3.5), Copper-translocating P-type ATPase (EC 3.6.3.4)	2925bp	975aa	98%	gi 739087196 WP_036958281.1	0.0
*CIA	Lead, cadmium, zinc and mercury transporting ATPase (EC 3.6.3.3) (EC 3.6.3.5), Copper-translocating P-type ATPase (EC 3.6.3.4)	2400bp	800aa	99%	gi 1042403368 OBY37647.1	0.0
BCO	Blue copper oxidase CueO precursor	1617bp	539aa	99%	gi 490381034 WP_004260551.1	0.0
*HL	Cytochrome c heme lyase subunit CcmF	1932bp	644aa	99%	gi 491051693 WP_004913344.1	0.0
*HL	Cytochrome c heme lyase subunit CcmL, Cytochrome c heme lyase subunit CcmH	1071bp	357aa	99%	gi 490382689 WP_004262201.1	0.0
CRD	Copper resistance protein D	897bp	299aa	99%	gi 490374296 WP_004253898.1	0.0
CopG	CopG protein	423bp	141aa	98%	gi 491043999 WP_004905661.1	2E-97
copCp	Copper resistance protein C precursor	387bp	129aa	100%	gi 490374293 WP_004253895.1	7E-86
Streptothricin resistance						
SatA	Streptothricin acetyltransferase, Streptomyces lavendulae type	525bp	175aa	100%	gi 446626810 WP_000704156.1	1E-125
Adaptation to d-cysteine						
DcyD	D-cysteine desulfhydrase (EC 4.4.1.15)	990bp	330aa	99%	gi 490374828 WP_004254429.1	0.0

Mercuric reductase						
MIR	Mercuric ion reductase (EC 1.16.1.1)	1686bp	562aa	100%	gi 446131441 WP_000209296.1	0.0
Resistance to fluoroquinolones						
parC	Topoisomerase IV subunit A (EC 5.99.1.-)	2256bp	752aa	99%	gi 490378099 WP_004257697.1	0.0
parE	Topoisomerase IV subunit B (EC 5.99.1.-)	1896bp	632aa	99%	gi 490378103 WP_004257701.1	0.0
gyrA	DNA gyrase subunit A (EC 5.99.1.3)	2616bp	872aa	99%	gi 491050539 WP_004912190.1	0.0
gyrB	DNA gyrase subunit B (EC 5.99.1.3)	2415bp	805aa	99%	gi 491044587 WP_004906247.1	0.0
Cobalt-zinc-cadmium resistance						
*CzcD	Cobalt-zinc-cadmium resistance protein	930bp	310aa	100%	gi 490382252 WP_004261765.1	0.0
*CzcA?	Cobalt-zinc-cadmium resistance protein	3078bp	1026aa	99%	gi 491044000 WP_004905662.1	0.0
*CusB/CzsB	CzcA, Cation efflux system protein CusA Probable Co/Zn/Cd efflux system membrane fusion protein	1095bp	365aa	62%	gi 1093685069 WP_070928012.1	6E-144
*CusB/CzsB	Probable Co/Zn/Cd efflux system membrane fusion protein	1110bp	370aa	99%	gi 490381614 WP_004261129.1	0.0
*CusB/CzsB	Cobalt/zinc/cadmium efflux RND transporter, membrane fusion protein, CzcB family	1269bp	423aa	99%	gi 491044001 WP_004905663.1	0.0
*CusA	Cobalt-zinc-cadmium resistance protein	3078bp	1026aa	99%	gi 491044000 WP_004905662.1	0.0
*CzrR	CzcA, Cation efflux system protein CusA DNA-binding heavy metal response regulator	684bp	228aa	97%	gi 491044004 WP_004905666.1	1E-153
cusF	Cation efflux system protein CusF precursor	342bp	114aa	97%	gi 491044002 WP_004905664.1	4E-74
*TR	Transcriptional regulator, MerR family	423bp	141aa	99%	gi 490374914 WP_004254515.1	2E-92
Mercury resistance operon						
MerC	Mercuric transport protein, MerC	426bp	142aa	100%	gi 446445141 WP_000522996.1	3E-98
MerA	Mercuric ion reductase (EC 1.16.1.1)	1686bp	562aa	100%	gi 446131441 WP_000209296.1	0.0
MerD	Mercuric resistance operon coregulator	246bp	82aa	99%	gi 585339711 WP_024221342.1	2E-47
Copper homeostasis: copper tolerance						
ScsB	Membrane protein, suppressor for copper-sensitivity ScsB	2073bp	691aa	99%	gi 490377422 WP_004257021.1	0.0

ScsC	Secreted protein, suppressor for copper-sensitivity ScsC	732bp	244aa	99%	gi 490377417 WP_004257016.1	7E-169
ScsD	Membrane protein, suppressor for copper-sensitivity ScsD	501bp	167aa	100%	gi 491047306 WP_004908960.1	5E-115
CutF	Copper homeostasis protein CutF precursor, Lipoprotein NlpE involved in surface adhesion	657bp	219aa	99%	gi 490377815 WP_004257413.1	3E-156
CutE	Apolipoprotein N-acyltransferase (EC 2.3.1.-), Copper homeostasis protein CutE	1530bp	510aa	99%	gi 490377396 WP_004256995.1	0.0
CorC	Magnesium and cobalt efflux protein CorC	1287bp	429aa	99%	gi 1042400154 OBY34443.1	0.0
CorC	Magnesium and cobalt efflux protein CorC	909bp	303aa	99%	gi 291314305 EFE54758.1	0.0
CorC	Magnesium and cobalt efflux protein CorC	1368bp	456aa	100%	gi 491044300 WP_004905961.1	0.0
Aminoglycoside adenylyltransferases						
AadA1	Streptomycin 3"-O-adenylyltransferase (EC 2.7.7.47), Spectinomycin 9-O-adenylyltransferase	159bp	53aa	100%	gi 487942357 WP_002015823.1	3E-31
AadA1	Streptomycin 3"-O-adenylyltransferase (EC 2.7.7.47), Spectinomycin 9-O-adenylyltransferase	1014bp	338aa	100%	gi 678222195 AIM49564.1	0.0
AadA1	Streptomycin 3"-O-adenylyltransferase (EC 2.7.7.47), Spectinomycin 9-O-adenylyltransferase	792bp	264aa	100%	gi 501453940 WP_012477385.1	0.0
AadA1	Streptomycin 3"-O-adenylyltransferase (EC 2.7.7.47), Spectinomycin 9-O-adenylyltransferase	150bp	50aa	100%	gi 486167809 WP_001531151.1	5E-29
AadA1	Streptomycin 3"-O-adenylyltransferase (EC 2.7.7.47), Spectinomycin 9-O-adenylyltransferase	789bp	263aa	100%	gi 447129059 WP_001206315.1	0.0
AadA2	Spectinomycin 9-O-adenylyltransferase	765bp	255aa	99%	gi 490381153 WP_004260670.1	2E-178
AadA2	Spectinomycin 9-O-adenylyltransferase	159bp	53aa	100%	gi 487942357 WP_002015823.1	3E-31
AadA2	Spectinomycin 9-O-adenylyltransferase	1014bp	338aa	100%	gi 678222195 AIM49564.1	0.0
AadA2	Spectinomycin 9-O-adenylyltransferase	792bp	264aa	100%	gi 501453940 WP_012477385.1	0.0
AadA2	Spectinomycin 9-O-adenylyltransferase	150bp	50aa	100%	gi 486167809 WP_001531151.1	5E-29

AadA2	Spectinomycin 9-O-adenylyltransferase	789bp	263aa	100%	gi 447129059 WP_001206315.1	0.0
Arsenic resistance						
arsR	Arsenical resistance operon repressor	336bp	112aa	100%	gi 757599390 WP_042847035.1	4E-73
arsR	Arsenical resistance operon repressor	336bp	112aa	99%	gi 739097345 WP_036968128.1	6E-72
arsR	Arsenical resistance operon repressor	189bp	63aa	92%	gi 1116088436 WP_072144547.1	2E-33
arsD	Arsenical resistance operon trans-acting repressor ArsD	366bp	122aa	100%	gi 757599389 WP_042847034.1	2E-85
arsD	Arsenical resistance operon trans-acting repressor ArsD	366bp	122aa	99%	gi 739097347 WP_036968130.1	2E-84
arsA	Arsenical pump-driving ATPase (EC 3.6.3.16)	1752bp	584aa	99%	gi 827610366 KLO01850.1	0.0
arsA	Arsenical pump-driving ATPase (EC 3.6.3.16)	1572bp	584aa	99%	gi 739097348 WP_036968131.1	0.0
arsB	Arsenic efflux pump protein	1290bp	430aa	98%	gi 490357905 WP_004237678.1	0.0
arsB	Arsenic efflux pump protein	1290bp	430aa	99%	gi 739104700 WP_036975217.1	0.0
arsB	Arsenic efflux pump protein	1362bp	454aa	98%	gi 739104700 WP_036975217.1	0.0
arsC	Arsenate reductase (EC 1.20.4.1)	357bp	119aa	99%	gi 490373943 WP_004253545.1	1E-78
arsC	Arsenate reductase (EC 1.20.4.1)	417bp	139aa	91%	gi 749308148 WP_040132278.1	8E-88
arsC	Arsenate reductase (EC 1.20.4.1)	441bp	147aa	100%	gi 739097354 WP_036968137.1	2E-100

Iron acquisition and metabolism

Sigla	Gene	Tamaño	Aminoácidos	Identidad	Secuenciamento	E. value
Siderophore Aerobactin						
iutA	Aerobactin siderophore receptor iutA TonB-dependent siderophore receptor	2202bp	734aa	99%	gi 490378242 WP_004257840.1	0.0
Heme, hemin uptake and utilization systems in GramPositives						
*Heme_oxygenase s	Hemin transport protein HmuS	1050bp	350aa	99%	gi 491051296 WP_004912947.1	0.0
*Heme_oxygenase s	Hemin transport protein HmuS	1050bp	350aa	99%	gi 739087580 WP_036958663.1	0.0
Ferrous iron transporter EfeUOB, low-pH-induced						
*EfeU	Ferrous iron transport permease EfeU	834bp	278aa	99%	gi 490381107 WP_004260624.1	0.0
EfeB	Ferrous iron transport peroxidase EfeB	1275bp	425aa	99%	gi 490381112 WP_004260629.1	0.0
Encapsulating protein for DyP-type peroxidase and ferritin-like protein oligomers						
DyP	Predicted dye-decolorizing peroxidase (DyP), encapsulated subgroup	900bp	300aa	99%	gi 490385387 WP_004264885.1	0.0
Yfe	Predicted outer membrane lipoprotein YfeY	606bp	202aa	99%	gi 915327114 WP_050763802.1	4E-145
Hemin transport system						
HBP	Periplasmic hemin-binding protein	822bp	274aa	99%	gi 491051298 WP_004912949.1	0.0
*Rec	TonB-dependent hemin , ferrichrome receptor	2019bp	673aa	91%	gi 739087359 WP_036958444.1	0.0
*Rec	TonB-dependent hemin , ferrichrome receptor	2145bp	715aa	99%	gi 490384250 WP_004263756.1	0.0
hmuS	Hemin transport protein HmuS	1050bp	350aa	99%	gi 491051296 WP_004912947.1	0.0
hmuS	Hemin transport protein HmuS	1050bp	350aa	99%	gi 739087580 WP_036958663.1	0.0

ATPb	ABC-type hemin transport system, ATPase component	789bp	263aa	98%	gi 1172252176 WP_080641516.1	0.0
ATPb	ABC-type hemin transport system, ATPase component	768bp	256aa	99%	gi 1042402961 OBY37241.1	2E-178
PP_7	Hemin ABC transporter, permease protein	1002bp	334aa	99%	gi 491051302 WP_004912953.1	0.0
PP_7	Hemin ABC transporter, permease protein	1017bp	339aa	100%	gi 1042402962 OBY37242.1	0.0
TonB	Ferric siderophore transport system, periplasmic binding protein TonB	744bp	248aa	97%	gi 490382765 WP_004262277.1	1E-162
TonB	Ferric siderophore transport system, periplasmic binding protein TonB	966bp	322aa	98%	gi 490378848 WP_004258445.1	0.0
TonB	Ferric siderophore transport system, periplasmic binding protein TonB	720bp	240aa	99%	gi 739086610 WP_036957695.1	6E-173
TonB	Ferric siderophore transport system, periplasmic binding protein TonB	783bp	261aa	99%	gi 739087479 WP_036958564.1	3E-179
TonB	Ferric siderophore transport system, periplasmic binding protein TonB	756bp	252aa	97%	gi 490373999 WP_004253601.1	1E-165
Heme, hemin uptake and utilization systems in GramNegatives						
HBP	Periplasmic hemin-binding protein	822bp	274aa	99%	gi 491051298 WP_004912949.1	0.0
fhuC	Ferrichrome transport ATP-binding protein FhuC (TC 3.A.1.14.3)	783bp	261aa	100%	gi 490382157 WP_004261670.1	0.0
fhuC	Ferrichrome transport ATP-binding protein FhuC (TC 3.A.1.14.3)	786bp	262aa	100%	gi 757595221 WP_042843418.1	8E-180
FCR	TonB-dependent hemin , ferrichrome receptor	2019bp	673aa	91%	gi 739087359 WP_036958444.1	0.0
FCR	TonB-dependent hemin , ferrichrome receptor	2145bp	715aa	99%	gi 490384250 WP_004263756.1	0.0
FR	Flavin reductase (EC 1.5.1.30)	624bp	208aa	99%	gi 739087413 WP_036958498.1	7E-143
ETFb	Electron transfer flavoprotein, beta subunit	756bp	252aa	99%	gi 490378882 WP_004258479.1	6E-168
ParA	Paraquat-inducible protein A	1227bp	409aa	99%	gi 1102614758 WP_071585858.1	0.0
ParA	Paraquat-inducible protein A	1227bp	409aa	99%	gi 739086601 WP_036957686.1	0.0

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ParB	Paraquat-inducible protein B	1650bp	550aa	99%	gi 490376536 WP_004256135.1	0.0
ParB	Paraquat-inducible protein B	2340bp	780aa	99%	gi 490374304 WP_004253906.1	0.0
hmuS	Hemin transport protein HmuS	1050bp	350aa	99%	gi 491051296 WP_004912947.1	0.0
hmuS	Hemin transport protein HmuS	1050bp	350aa	99%	gi 739087580 WP_036958663.1	0.0
PP_7	Hemin ABC transporter, permease protein	1002bp	334aa	99%	gi 491051302 WP_004912953.1	0.0
PP_7	Hemin ABC transporter, permease protein	1017bp	339aa	100%	gi 1042402962 OBY37242.1	0.0
ATPb	ABC-type hemin transport system, ATPase component	789bp	263aa	98%	gi 1172252176 WP_080641516.1	0.0
ATPb	ABC-type hemin transport system, ATPase component	768bp	256aa	99%	gi 1042402961 OBY37241.1	2E-178
TonB	Ferric siderophore transport system, periplasmic binding protein TonB	744bp	248aa	97%	gi 490382765 WP_004262277.1	1E-162
TonB	Ferric siderophore transport system, periplasmic binding protein TonB	966bp	322aa	98%	gi 490378848 WP_004258445.1	0.0
TonB	Ferric siderophore transport system, periplasmic binding protein TonB	720bp	240aa	99%	gi 739086610 WP_036957695.1	6E-173
TonB	Ferric siderophore transport system, periplasmic binding protein TonB	783bp	261aa	99%	gi 739087479 WP_036958564.1	3E-179
TonB	Ferric siderophore transport system, periplasmic binding protein TonB	756bp	252aa	97%	gi 490373999 WP_004253601.1	1E-165

Phages, Prophages

Sigla	Gene	Tamanho	Aminoácidos	Identidade	Sequenciamento	E. value
Transposable elements - CBSS-203122.12.peg.188						
R1	TniB NTP-binding protein	510bp	170aa	100%	gi 213054790 ACJ39692.1	2E-120
R8	MG(2+) CHELATASE FAMILY PROTEIN ComM-related protein	1527bp	509aa	99%	gi 491044471 WP_004906131.1	0.0
R9	MG(2+) CHELATASE FAMILY PROTEIN ComM-related protein	1527bp	509aa	99%	gi 491044471 WP_004906131.1	0.0
Phages, Prophages, Transposable elements, Plasmids - Integrans						
IntI2	Integron integrase IntI2	537bp	179aa	100%	gi 445994041 WP_000071896.1	8E-125
IntI2	Integron integrase IntI2	537bp	179aa	100%	gi 446419664 WP_000497519.1	9E-75
IntIPac	Integron integrase IntIPac	1014bp	338bp	100%	gi 471237918 EMR15449.1	0.0
Phages, Prophages - Phage tail proteins						
TP	Phage tail protein	429bp	143aa	100%	gi 739091102 WP_036962064.1	2E-98
*Measure	Phage tail length tape-measure protein	2700bp	900aa	98%	gi 873900395 WP_048606828.1	0.0
*Tube	Phage major tail tube protein	516bp	172aa	100%	gi 1101998870 WP_071548731.1	8E-120
*Sheath	Phage tail sheath monomer	1530bp	510aa	100%	gi 1101998871 WP_071548732.1	0.0
Phages, Prophages - Phage replication						
*Rep	Phage replication protein	2424bp	808aa	93%	gi 1101998888 WP_071548749.1	0.0
*dPol_3	DNA polymerase III alpha subunit (EC 2.7.7.7)	3483bp	1161aa	100%	gi 739086905 WP_036957990.1	0.0
Phages, Prophages - Phage tail proteins 2						
TMP1_1	Phage tape measure	447bp	149aa	99%	gi 873900393 WP_048606826.1	1E-99
TSM	Phage tail sheath monomer	1530bp	510aa	100%	gi 1101998871 WP_071548732	0.0

TMP	Phage tail length tape-measure protein	2700bp	900aa	98%	gi 873900395 WP_048606828.1	0.0
Phages, Prophages - Phage tail fiber proteins						
TFP	Phage tail fiber protein	1530bp	510aa	99%	gi 1101502589 APC13982.1	0.0
TFAP	Phage tail fiber assembly protein	645bp	215aa	100%	gi 1101502588 APC13981.1	2E-149
TFP2	Phage tail fibers	612bp	204aa	100%	gi 873900403 WP_048606836.1	2E-146
Phages, Prophages - Phage capsid proteins						
*Generic	Phage capsid and scaffold	444bp	148aa	100%	gi 739091104 WP_036962066.1	3.00E-108
MajCap	Phage major capsid protein	1095bp	365aa	99%	gi 1101998881 WP_071548742.1	0.0
*Scaffolding	Phage capsid scaffolding protein	828bp	276aa	99%	gi 1101998882 WP_071548743.1	0.0
HCSP	Phage head completion-stabilization protein	477bp	159aa	100%	gi 1101998879 WP_071548740.1	4.0E-108

Motility and Chemotaxis

Sigla	Gene	Tamanho	Aminoácidos	Identidade	Sequenciamento	E. value
Bacterial Chemotaxis						
aer	Aerotaxis sensor receptor protein	1533bp	511aa	98%	gi 739086922 WP_036958007.1	0.0
tsr	Methyl-accepting chemotaxis protein I (serine chemoreceptor protein)	1653bp	551aa	98%	gi 490373823 WP_004253426.1	0.0
tsr	Methyl-accepting chemotaxis protein I (serine chemoreceptor protein)	1569bp	523aa	99%	gi 490376118 WP_004255718.1	0.0
tsr	Methyl-accepting chemotaxis protein I (serine chemoreceptor protein)	1602bp	534aa	99%	gi 490376121 WP_004255721.1	0.0
tsr	Methyl-accepting chemotaxis protein I (serine chemoreceptor protein)	1488bp	496aa	99%	gi 490378843 WP_004258440.1	0.0
*cheW-V	Positive regulator of CheA protein activity (CheW)	498bp	166aa	100%	gi 491048180 WP_004909833.1	3E-111
cheA	Signal transduction histidine kinase CheA (EC 2.7.3.-)	2058bp	686aa	99%	gi 739086771 WP_036957856.1	0.0
cheY	Chemotaxis regulator - transmits chemoreceptor signals to flagellar motor components CheY	393bp	131aa	100%	gi 490376105 WP_004255705.1	7E-85
cheZ	Chemotaxis response - phosphatase CheZ	639bp	213aa	99%	gi 104239971 OBY34011.1	8E-142
cheB	Chemotaxis response regulator protein-glutamate methylesterase CheB (EC 3.1.1.61)	1068bp	356aa	99%	gi 490376109 WP_004255709.1	0.0
fliG	Flagellar motor switch protein FliG	993bp	331aa	100%	gi 490376000 WP_004255600.1	0.0
fliM	Flagellar motor switch protein FliM	1008bp	336aa	99%	gi 490376019 WP_004255619.1	0.0
fliN	Flagellar motor switch protein FliN	417bp	139aa	100%	gi 490376021 WP_004255621.1	1E-90
malE	Maltose/maltodextrin ABC transporter, substrate binding periplasmic protein MalE	1191bp	397aa	92%	gi 1057035456 WP_068445917.1	0.0
dppA	Dipeptide-binding ABC transporter, periplasmic substrate-binding component (TC 3.A.1.5.2)	1608bp	536aa	100%	gi 490383128 WP_004262639.1	0.0
dppA	Dipeptide-binding ABC transporter, periplasmic substrate-binding component (TC 3.A.1.5.2)	1560bp	520aa	99%	gi 739086975 WP_036958060.1	0.0

cheR	Chemotaxis protein methyltransferase CheR (EC 2.1.1.80)	837bp	279aa	100%	gi 490376113 WP_004255713.1	0.0
Flagellum						
flgB	Flagellar basal-body rod protein FlgB	414bp	138aa	100%	gi 490376069 WP_004255669.1	2E-91
flgC	Flagellar basal-body rod protein FlgC	405bp	135aa	100%	gi 491048215 WP_004909868.1	3E-89
FlgD	Flagellar basal-body rod modification protein FlgD	858bp	286aa	99%	gi 490376064 WP_004255664.1	0.0
flgE	Flagellar hook protein FlgE	1221bp	407aa	96%	gi 873900357 WP_048606790.1	0.0
flgF	Flagellar basal-body rod protein FlgF	756bp	252aa	100%	gi 490376059 WP_004255659.1	2E-174
flgG	Flagellar basal-body rod protein FlgG	783bp	261aa	99%	gi 490376057 WP_004255657.1	0.0
FlgH	Flagellar L-ring protein FlgH	750bp	250aa	100%	gi 291314697 EFE55150.1	0.0
flgI	Flagellar P-ring protein FlgI	1104bp	368aa	100%	gi 490376051 WP_004255651.1	0.0
flgJ	Flagellar protein FlgJ [peptidoglycan hydrolase] (EC 3.2.1.-)	981bp	327aa	98%	gi 490376048 WP_004255648.1	0.0
flgK	Flagellar hook-associated protein FlgK	1644bp	548aa	99%	gi 739086767 WP_036957852.1	0.0
flgL	Flagellar hook-associated protein FlgL	936bp	312aa	98%	gi 491048236 WP_004909889.1	0.0
FliA	Flagellar biosynthesis protein FliA	2100bp	700aa	100%	gi 490376087 WP_004255687.1	0.0
fliE	Flagellar hook-basal body complex protein FliE	312bp	104aa	100%	gi 490375994 WP_004255594.1	4E-64
fliF	Flagellar M-ring protein FliF	1686bp	562aa	99%	gi 490375997 WP_004255597.1	0.0
fliG	Flagellar motor switch protein FliG	993bp	331aa	100%	gi 490376000 WP_004255600.1	0.0
fliH	Flagellar assembly protein FliH	729bp	243aa	99%	gi 490376003 WP_004255603.1	8E-177
FliI	Flagellum-specific ATP synthase FliI	1311bp	437aa	99%	gi 490376006 WP_004255606.1	0.0
fliK	Flagellar hook-length control protein FliK	1368bp	456aa	97%	gi 490376013 WP_004255613.1	0.0
fliJ	Flagellar protein FliJ	444bp	148aa	99%	gi 490376009 WP_004255609.1	3E-100
fliM	Flagellar motor switch protein FliM	1008bp	336aa	99%	gi 490376019 WP_004255619.1	0.0
fliN	Flagellar motor switch protein FliN	417bp	139aa	100%	gi 490376021 WP_004255621.1	1E-90
fliP	Flagellar biosynthesis protein FliP	744bp	248aa	99%	gi 739086765 WP_036957850.1	3E-167
fliQ	Flagellar biosynthesis protein FliQ	270bp	90aa	99%	gi 490376030 WP_004255630.1	2E-52
FliR	Flagellar biosynthesis protein FliR	780bp	260aa	99%	gi 490376033 WP_004255633.1	7E-180

*Flagellar-Regulation	RNA polymerase sigma factor RpoD	1854bp	618aa	100%	gi 1102614763 WP_071585863.1	0.0
*Flagellar-Regulation	Flagellar transcriptional activator FlhC	591bp	197aa	100%	gi 490376132 WP_004255732.1	1E-140
*Flagellar-Regulation	Flagellar biosynthesis protein FliZ	519bp	173aa	100%	gi 490376157 WP_004255757.1	4E-123
*Flagellar-Regulation	RNA polymerase sigma-54 factor RpoN	1446bp	482aa	99%	gi 490378708 WP_004258305.1	0.0
*Flagellar-Regulation	RNA polymerase sigma factor for flagellar operon	726bp	242aa	99%	gi 291314661 EFE55114.1	0.0
*Flagellin	Flagellar biosynthesis protein FliC	1170bp	390aa	87%	gi 1055836465 WP_067424616.1	0.0
fliD	Flagellar hook-associated protein FliD	1437bp	479aa	90%	gi 1057031755 WP_068442704.1	0.0
fliS	Flagellar biosynthesis protein FliS	399bp	133aa	100%	gi 491048316 WP_004909969.1	1E-86
fliT	Flagellar biosynthesis protein FliT	390bp	130aa	99%	gi 490375964 WP_004255564.1	2E-85
*fliO	Flagellar biosynthesis protein FliO	459bp	153aa	99%	gi 490376024 WP_004255624.1	7E-102
FlhB	Flagellar biosynthesis protein FlhB	1152bp	384aa	99%	gi 490376088 WP_004255688.1	0.0
MotA	Flagellar motor rotation protein MotA	954bp	318aa	99%	gi 291314724 EFE55177.1	0.0
MotB	Flagellar motor rotation protein MotB	918bp	306aa	99%	gi 490376126 WP_004255726.1	0.0
*flhD	Flagellar transcriptional activator FlhD	351bp	117aa	100%	gi 739086775 WP_036957860.1	3E-74
fliL	Flagellar biosynthesis protein FliL	480bp	160aa	99%	gi 490376015 WP_004255615.1	3E-107
Flagellar motility						
FlhA	Flagellar biosynthesis protein FlhA	2100bp	700aa	100%	gi 490376087 WP_004255687.1	0.0
fliM	Flagellar motor switch protein FliM	1008bp	336aa	99%	gi 490376019 WP_004255619.1	0.0
fliN	Flagellar motor switch protein FliN	417bp	139aa	100%	gi 490376021 WP_004255621.1	1E-90
cheY	Chemotaxis regulator - transmits chemoreceptor signals to flagellar motor components CheY	393bp	131aa	100%	gi 490376105 WP_004255705.1	7E-85
cheA	Signal transduction histidine kinase CheA (EC 2.7.3.-)	2058bp	686aa	99%	gi 739086771 WP_036957856.1	0.0
FlhB	Flagellar biosynthesis protein FlhB	1152bp	384aa	99%	gi 490376088 WP_004255688.1	0.0

FliI	Flagellum-specific ATP synthase FliI	1311bp	437aa	99%	gi 490376006 WP_004255606.1	0.0
FliR	Flagellar biosynthesis protein FliR	780bp	260aa	99%	gi 490376033 WP_004255633.1	7E-180
FlgD	Flagellar basal-body rod modification protein FlgD	858bp	286aa	99%	gi 490376064 WP_004255664.1	0.0
FlgH	Flagellar L-ring protein FlgH	750bp	250aa	100%	gi 291314697 EFE55150.1	0.0
MotA	Flagellar motor rotation protein MotA	954bp	318aa	99%	gi 291314724 EFE55177.1	0.0
MotB	Flagellar motor rotation protein MotB	918bp	306aa	99%	gi 490376126 WP_004255726.1	0.0
FliA	RNA polymerase sigma factor for flagellar operon	726bp	242aa	99%	gi 291314661 EFE55114.1	9E-167
RpoN	RNA polymerase sigma-54 factor RpoN	1446bp	482aa	99%	gi 490378708 WP_004258305.1	0.0

Suplemento II

Localização das Ilhas genômicas PR01

Island start	Island end	Length	Method	Gene ID	Locus	Gene start	Gene end	Strand	Product
121511	130328	8817	Predicted by at least one method		Predicted_0402	121511	121912	1	hypothetical protein
121511	130328	8817	Predicted by at least one method	mrpA_4	Predicted_0403	121905	122438	1	Major MR/P fimbria protein precursor
121511	130328	8817	Predicted by at least one method		Predicted_0433	123815	124828	1	hypothetical protein
121511	130328	8817	Predicted by at least one method		Predicted_0435	126856	127881	-1	hypothetical protein
121511	130328	8817	Predicted by at least one method		Predicted_0435	127897	128157	-1	hypothetical protein
121511	130328	8817	Predicted by at least one method	papD_1	Predicted_0418	129618	130328	-1	Chaperone protein PapD precursor
162908	1715575	86488	Predicted by at least one method	hcpC_6	Predicted_0419	162908	163011	-1	Putative beta-lactamase HcpC precursor
162908	1715575	86488	Predicted by at least one method	rhsC	Predicted_0076	163184	163486	-1	Putative deoxyribonuclease RhsC
162908	1715575	86488	Predicted by at least one method		Predicted_0076	163500	163572	-1	hypothetical protein
162908	1715575	86488	Predicted by at least one method		Predicted_0076	163646	163670	-1	hypothetical protein
162908	1715575	86488	Predicted by at least one method		Predicted_0076	163677	163705	-1	hypothetical protein
162908	1715575	86488	Predicted by at least one method		Predicted_0076	163703	163876	-1	Zinc-binding domain of primase-helicase
162908	1715575	86488	Predicted by at least one method		Predicted_0076	163894	163933	-1	hypothetical protein
162908	1715575	86488	Predicted by at least one method		Predicted_0076	163943	163958	-1	hypothetical protein
162908	1715575	86488	Predicted by at least one method		Predicted_0075	163978	164063	-1	hypothetical protein
162908	1715575	86488	Predicted by at least one method		Predicted_0075	164071	164127	-1	hypothetical protein
162908	1715575	86488	Predicted by at least one method		Predicted_0075	164134	164156	-1	hypothetical protein
162908	1715575	86488	Predicted by at least one method		Predicted_0075	164157	164184	-1	hypothetical protein

7			one method		6	7	6		
162908	1715575	86488	Predicted by at least one method	yfbR_1	Predicted_0075	164183	164244	-1	5'-deoxynucleotidase YfbR
7			one method		5	9	4		
162908	1715575	86488	Predicted by at least one method		Predicted_0075	164251	164271	-1	hypothetical protein
7			one method		4	6	9		
162908	1715575	86488	Predicted by at least one method		Predicted_0075	164292	164537	-1	hypothetical protein
7			one method		3	1	4		
162908	1715575	86488	Predicted by at least one method		Predicted_0075	164546	164584	-1	hypothetical protein
7			one method		2	2	8		
162908	1715575	86488	Predicted by at least one method		Predicted_0075	164608	164663	-1	hypothetical protein
7			one method		1	1	5		
162908	1715575	86488	Predicted by at least one method		Predicted_0075	164670	164700	-1	hypothetical protein
7			one method		0	9	8		
162908	1715575	86488	Predicted by at least one method	hhaIM	Predicted_0074	164720	164882	-1	Modification methylase HhaI
7			one method		9	1	6		
162908	1715575	86488	Predicted by at least one method		Predicted_0074	164890	164920	-1	hypothetical protein
7			one method		8	4	6		
162908	1715575	86488	Predicted by at least one method		Predicted_0074	164928	164962	-1	hypothetical protein
7			one method		7	4	8		
162908	1715575	86488	Predicted by at least one method		Predicted_0074	164982	165017	-1	hypothetical protein
7			one method		6	0	3		
162908	1715575	86488	Predicted by at least one method		Predicted_0074	165024	165043	-1	hypothetical protein
7			one method		5	2	3		
162908	1715575	86488	Predicted by at least one method		Predicted_0074	165049	165076	-1	hypothetical protein
7			one method		4	1	6		
162908	1715575	86488	Predicted by at least one method		Predicted_0074	165077	165122	-1	hypothetical protein
7			one method		3	1	3		
162908	1715575	86488	Predicted by at least one method		Predicted_0074	165121	165160	-1	hypothetical protein
7			one method		2	3	2		
162908	1715575	86488	Predicted by at least one method	ascD_2	Predicted_0074	165166	165188	-1	CDP-6-deoxy-L-threo-D-glycero-4-hexulose-3-dehydrase reductase
7			one method		1	4	8		DNA methylase
162908	1715575	86488	Predicted by at least one method		Predicted_0074	165195	165283	-1	
7			one method		0	4	8		
162908	1715575	86488	Predicted by at least one method		Predicted_0073	165291	165341	-1	hypothetical protein
7			one method		9	2	2		
162908	1715575	86488	Predicted by at least one method		Predicted_0073	165350	165404	-1	hypothetical protein
7			one method		8	3	2		

162908 7	1715575	86488	Predicted by at least one method		Predicted_0073 7	165430 7	165464 2	-1	hypothetical protein
162908 7	1715575	86488	Predicted by at least one method		Predicted_0073 6	165470 4	165650 0	-1	von Willebrand factor type A domain
162908 7	1715575	86488	Predicted by at least one method		Predicted_0073 5	165658 5	165786 2	-1	hypothetical protein
162908 7	1715575	86488	Predicted by at least one method		Predicted_0073 4	165806 2	165907 2	-1	YqaJ-like viral recombinase domain
162908 7	1715575	86488	Predicted by at least one method		Predicted_0073 3	165913 5	166012 4	-1	RecT family
162908 7	1715575	86488	Predicted by at least one method	ssb_2	Predicted_0073 2	166021 9	166074 6	-1	Single-stranded DNA-binding protein
162908 7	1715575	86488	Predicted by at least one method		Predicted_0073 1	166081 0	166171 8	-1	hypothetical protein
162908 7	1715575	86488	Predicted by at least one method	cobS_3	Predicted_0073 0	166172 9	166269 7	-1	Aerobic cobaltochelatase subunit CobS
162908 7	1715575	86488	Predicted by at least one method		Predicted_0072 9	166291 7	166310 2	-1	hypothetical protein
162908 7	1715575	86488	Predicted by at least one method		Predicted_0072 8	166340 6	166372 6	-1	hypothetical protein
162908 7	1715575	86488	Predicted by at least one method		Predicted_0072 7	166402 0	166466 1	1	hypothetical protein
162908 7	1715575	86488	Predicted by at least one method		Predicted_0072 6	166478 4	166564 4	1	hypothetical protein
162908 7	1715575	86488	Predicted by at least one method		Predicted_0072 5	166568 3	166848 1	-1	conjugal transfer mating pair stabilization protein TraN
162908 7	1715575	86488	Predicted by at least one method		Predicted_0072 4	166859 7	166960 4	-1	TraU protein
162908 7	1715575	86488	Predicted by at least one method	yfgF	Predicted_0072 3	166960 1	167027 2	-1	Cyclic di-GMP phosphodiesterase YfgF
162908 7	1715575	86488	Predicted by at least one method		Predicted_0072 2	167026 9	167153 4	-1	Type-F conjugative transfer system pilin assembly protein
162908 7	1715575	86488	Predicted by at least one method		Predicted_0072 1	167149 7	167202 7	-1	Peptidase S26
162908 7	1715575	86488	Predicted by at least one method		Predicted_0072 0	167202 4	167234 1	-1	hypothetical protein
162908 7	1715575	86488	Predicted by at least one method		Predicted_0071 9	167235 6	167480 3	-1	F pilus assembly Type-IV secretion system for plasmid

									transfer
162908 7	1715575	86488	Predicted by at least one method	dsbC_3	Predicted_0071 8	167480 0	167550 7	-1	Thiol:disulfide interchange protein DsbC precursor
162908 7	1715575	86488	Predicted by at least one method		Predicted_0071 7	167565 6	168118 7	-1	hypothetical protein
162908 7	1715575	86488	Predicted by at least one method		Predicted_0071 6	168138 8	168177 1	-1	hypothetical protein
162908 7	1715575	86488	Predicted by at least one method		Predicted_0071 5	168178 4	168236 2	-1	Type IV conjugative transfer system lipoprotein (TraV)
162908 7	1715575	86488	Predicted by at least one method		Predicted_0071 4	168235 9	168367 5	-1	Bacterial conjugation TrbI-like protein
162908 7	1715575	86488	Predicted by at least one method		Predicted_0071 3	168367 2	168458 9	-1	TraK protein
162908 7	1715575	86488	Predicted by at least one method		Predicted_0071 2	168457 3	168519 9	-1	TraE protein
162908 7	1715575	86488	Predicted by at least one method		Predicted_0071 1	168562 2	168599 9	-1	hypothetical protein
162908 7	1715575	86488	Predicted by at least one method		Predicted_0071 0	168599 9	168614 2	-1	hypothetical protein
162908 7	1715575	86488	Predicted by at least one method		Predicted_0070 9	168632 3	168698 5	-1	hypothetical protein
162908 7	1715575	86488	Predicted by at least one method		Predicted_0070 8	168698 5	168736 2	-1	hypothetical protein
162908 7	1715575	86488	Predicted by at least one method		Predicted_0070 7	168737 2	168781 8	-1	hypothetical protein
162908 7	1715575	86488	Predicted by at least one method		Predicted_0070 6	168782 8	168845 7	-1	hypothetical protein
162908 7	1715575	86488	Predicted by at least one method		Predicted_0070 5	168841 4	168895 9	-1	hypothetical protein
162908 7	1715575	86488	Predicted by at least one method		Predicted_0070 4	168900 9	169087 4	-1	AAA-like domain
162908 7	1715575	86488	Predicted by at least one method		Predicted_0070 3	169087 1	169384 9	-1	Putative helicase
162908 7	1715575	86488	Predicted by at least one method		Predicted_0070 2	169401 7	169463 4	1	hypothetical protein
162908 7	1715575	86488	Predicted by at least one method	topB_1	Predicted_0070 1	169484 9	169704 1	1	DNA topoisomerase 3

162908 7	1715575	86488	Predicted by at least one method		Predicted_0070 0	169705 6	169754 4	1	hypothetical protein
162908 7	1715575	86488	Predicted by at least one method		Predicted_0069 9	169763 6	169791 4	-1	hypothetical protein
162908 7	1715575	86488	Predicted by at least one method		Predicted_0069 8	169809 3	169936 7	-1	Transposase DDE domain
162908 7	1715575	86488	Predicted by at least one method		Predicted_0069 7	169954 6	170016 3	-1	hypothetical protein
162908 7	1715575	86488	Predicted by at least one method		Predicted_0069 6	170025 7	170047 5	-1	hypothetical protein
162908 7	1715575	86488	Predicted by at least one method		Predicted_0069 5	170047 9	170074 2	-1	hypothetical protein
162908 7	1715575	86488	Predicted by at least one method		Predicted_0069 4	170068 1	170133 7	-1	hypothetical protein
162908 7	1715575	86488	Predicted by at least one method	ydiO	Predicted_0069 3	170133 7	170276 4	-1	putative BsuMI modification methylase subunit YdiO
162908 7	1715575	86488	Predicted by at least one method		Predicted_0069 2	170276 8	170326 8	-1	hypothetical protein
162908 7	1715575	86488	Predicted by at least one method		Predicted_0069 1	170327 7	170358 8	-1	hypothetical protein
162908 7	1715575	86488	Predicted by at least one method		Predicted_0069 0	170359 4	170410 0	-1	hypothetical protein
162908 7	1715575	86488	Predicted by at least one method		Predicted_0068 9	170409 3	170476 7	-1	hypothetical protein
162908 7	1715575	86488	Predicted by at least one method		Predicted_0068 8	170474 2	170502 3	-1	hypothetical protein
162908 7	1715575	86488	Predicted by at least one method		Predicted_0068 7	170501 6	170539 3	-1	hypothetical protein
162908 7	1715575	86488	Predicted by at least one method		Predicted_0068 6	170571 2	170585 5	1	hypothetical protein
162908 7	1715575	86488	Predicted by at least one method		Predicted_0068 5	170594 7	170658 2	1	hypothetical protein
162908 7	1715575	86488	Predicted by at least one method	korB	Predicted_0068 4	170695 6	170813 7	-1	Transcriptional repressor protein KorB
162908 7	1715575	86488	Predicted by at least one method	soj	Predicted_0068 3	170814 1	170892 6	-1	Sporulation initiation inhibitor protein Soj
162908 7	1715575	86488	Predicted by at least one method		Predicted_0068 2	170910 0	170941 1	-1	hypothetical protein

162908 7	1715575	86488	Predicted by at least one method		Predicted_0068 1	170951 0	170973 4	-1	hypothetical protein
162908 7	1715575	86488	Predicted by at least one method	bin3_2	Predicted_0068 0	170973 1	171046 8	-1	Putative transposon Tn552 DNA-invertase bin3
162908 7	1715575	86488	Predicted by at least one method		Predicted_0067 9	171057 5	171106 6	1	hypothetical protein
162908 7	1715575	86488	Predicted by at least one method	dhfrI	Predicted_0020 0	171263 2	171310 5	1	Dihydrofolate reductase type 1
162908 7	1715575	86488	Predicted by at least one method		Predicted_0020 1	171320 0	171372 4	1	TDP-fucosamine acetyltransferase
162908 7	1715575	86488	Predicted by at least one method	ant1_2	Predicted_0020 2	171378 2	171457 0	1	Streptomycin 3"- adenylyltransferase
162908 7	1715575	86488	Predicted by at least one method		Predicted_0020 3	171464 6	171514 3	1	hypothetical protein
162908 7	1715575	86488	Predicted by at least one method		Predicted_0020 4	171520 4	171557 5	1	hypothetical protein
169087 1	1705023	14152	Predicted by at least one method		Predicted_0070 4	168900 9	169087 4	-1	AAA-like domain
169087 1	1705023	14152	Predicted by at least one method		Predicted_0070 3	169087 1	169384 9	-1	Putative helicase
169087 1	1705023	14152	Predicted by at least one method		Predicted_0070 2	169401 7	169463 4	1	hypothetical protein
169087 1	1705023	14152	Predicted by at least one method	topB_1	Predicted_0070 1	169484 9	169704 1	1	DNA topoisomerase 3
169087 1	1705023	14152	Predicted by at least one method		Predicted_0070 0	169705 6	169754 4	1	hypothetical protein
169087 1	1705023	14152	Predicted by at least one method		Predicted_0069 9	169763 6	169791 4	-1	hypothetical protein
169087 1	1705023	14152	Predicted by at least one method		Predicted_0069 8	169809 3	169936 7	-1	Transposase DDE domain
169087 1	1705023	14152	Predicted by at least one method		Predicted_0069 7	169954 6	170016 3	-1	hypothetical protein
169087 1	1705023	14152	Predicted by at least one method		Predicted_0069 6	170025 7	170047 5	-1	hypothetical protein
169087 1	1705023	14152	Predicted by at least one method		Predicted_0069 5	170047 9	170074 2	-1	hypothetical protein
169087 1	1705023	14152	Predicted by at least one method		Predicted_0069 4	170068 1	170133 7	-1	hypothetical protein

169087 1	1705023	14152	Predicted by at least one method	ydiO	Predicted_0069 3	170133 7	170276 4	-1	putative BsuMI modification methylase subunit YdiO
169087 1	1705023	14152	Predicted by at least one method		Predicted_0069 2	170276 8	170326 8	-1	hypothetical protein
169087 1	1705023	14152	Predicted by at least one method		Predicted_0069 1	170327 7	170358 8	-1	hypothetical protein
169087 1	1705023	14152	Predicted by at least one method		Predicted_0069 0	170359 4	170410 0	-1	hypothetical protein
169087 1	1705023	14152	Predicted by at least one method		Predicted_0068 9	170409 3	170476 7	-1	hypothetical protein
169087 1	1705023	14152	Predicted by at least one method		Predicted_0068 8	170474 2	170502 3	-1	hypothetical protein
169087 1	1705023	14152	Predicted by at least one method		Predicted_0068 7	170501 6	170539 3	-1	hypothetical protein
170814 1	1717256	9115	Predicted by at least one method	soj	Predicted_0068 3	170814 1	170892 6	-1	Sporulation initiation inhibitor protein Soj
170814 1	1717256	9115	Predicted by at least one method		Predicted_0068 2	170910 0	170941 1	-1	hypothetical protein
170814 1	1717256	9115	Predicted by at least one method		Predicted_0068 1	170951 0	170973 4	-1	hypothetical protein
170814 1	1717256	9115	Predicted by at least one method	bin3_2	Predicted_0068 0	170973 1	171046 8	-1	Putative transposon Tn552 DNA-invertase bin3
170814 1	1717256	9115	Predicted by at least one method		Predicted_0067 9	171057 5	171106 6	1	hypothetical protein
170814 1	1717256	9115	Predicted by at least one method	dhfrI	Predicted_0020 0	171263 2	171310 5	1	Dihydrofolate reductase type 1
170814 1	1717256	9115	Predicted by at least one method		Predicted_0020 1	171320 0	171372 4	1	TDP-fucosamine acetyltransferase
170814 1	1717256	9115	Predicted by at least one method	ant1_2	Predicted_0020 2	171378 2	171457 0	1	Streptomycin 3"-adenylyltransferase
170814 1	1717256	9115	Predicted by at least one method		Predicted_0020 3	171464 6	171514 3	1	hypothetical protein
170814 1	1717256	9115	Predicted by at least one method		Predicted_0020 4	171520 4	171557 5	1	hypothetical protein
170814 1	1717256	9115	Predicted by at least one method		Predicted_0020 5	171604 8	171725 6	1	Transposase DDE domain
170814 1	1717256	9115	Predicted by at least one method		Predicted_0020 6	171725 3	171783 1	1	hypothetical protein

173036 4	1799760	69396	Predicted by at least one method	Predicted_0021 8	172973 5	173036 7	-1	hypothetical protein
173036 4	1799760	69396	Predicted by at least one method	Predicted_0021 9	173036 4	173092 4	-1	hypothetical protein
173036 4	1799760	69396	Predicted by at least one method	Predicted_0022 0	173126 7	173175 5	-1	hypothetical protein
173036 4	1799760	69396	Predicted by at least one method	Predicted_0022 1	173177 8	173210 4	-1	hypothetical protein
173036 4	1799760	69396	Predicted by at least one method	Predicted_0022 2	173209 1	173252 2	-1	hypothetical protein
173036 4	1799760	69396	Predicted by at least one method	Predicted_0022 3	173252 9	173278 3	-1	hypothetical protein
173036 4	1799760	69396	Predicted by at least one method	Predicted_0022 4	173278 0	173333 7	-1	hypothetical protein
173036 4	1799760	69396	Predicted by at least one method	Predicted_0022 5	173333 4	173379 8	-1	ASCH domain
173036 4	1799760	69396	Predicted by at least one method	Predicted_0022 6	173379 5	173409 7	-1	hypothetical protein
173036 4	1799760	69396	Predicted by at least one method	Predicted_0022 7	173442 4	173547 3	1	hypothetical protein
173036 4	1799760	69396	Predicted by at least one method	Predicted_0022 8	173562 5	173671 0	1	hypothetical protein
173036 4	1799760	69396	Predicted by at least one method	Predicted_0022 9	173672 8	173705 7	-1	hypothetical protein
173036 4	1799760	69396	Predicted by at least one method	Predicted_0023 0	173705 4	173782 7	-1	hypothetical protein
173036 4	1799760	69396	Predicted by at least one method	Predicted_0023 1	173781 4	173839 2	-1	hypothetical protein
173036 4	1799760	69396	Predicted by at least one method	Predicted_0023 2	173850 1	173902 5	-1	hypothetical protein
173036 4	1799760	69396	Predicted by at least one method	Predicted_0023 3	173902 5	173939 9	-1	hypothetical protein
173036 4	1799760	69396	Predicted by at least one method	Predicted_0023 4	173941 4	174031 9	-1	Modification methylase HpaII
173036 4	1799760	69396	Predicted by at least one method	Predicted_0023 5	174057 4	174076 2	-1	hypothetical protein
173036 4	1799760	69396	Predicted by at least one method	Predicted_0023 6	174077 4	174117 8	-1	hypothetical protein

173036 4	1799760	69396	Predicted by at least one method	nrdH_1	Predicted_0023 7	174119 2	174152 1	-1	Glutaredoxin-like NrdH	protein
173036 4	1799760	69396	Predicted by at least one method		Predicted_0023 8	174169 9	174201 0	-1	hypothetical protein	
173036 4	1799760	69396	Predicted by at least one method		Predicted_0023 9	174206 8	174244 2	-1	hypothetical protein	
173036 4	1799760	69396	Predicted by at least one method		Predicted_0024 0	174252 1	174320 1	-1	hypothetical protein	
173036 4	1799760	69396	Predicted by at least one method	dinG_1	Predicted_0024 1	174324 2	174530 2	-1	putative ATP-dependent helicase DinG	
173036 4	1799760	69396	Predicted by at least one method		Predicted_0024 2	174540 9	174571 7	-1	hypothetical protein	
173036 4	1799760	69396	Predicted by at least one method		Predicted_0024 3	174573 5	174601 3	-1	hypothetical protein	
173036 4	1799760	69396	Predicted by at least one method	virB	Predicted_0024 4	174629 0	174730 9	-1	Virulence transcriptional activator VirB	regulon
173036 4	1799760	69396	Predicted by at least one method	parA_1	Predicted_0024 5	174731 2	174855 9	-1	Plasmid partition protein A	
173036 4	1799760	69396	Predicted by at least one method		Predicted_0024 6	174877 5	174905 3	-1	hypothetical protein	
173036 4	1799760	69396	Predicted by at least one method		Predicted_0024 7	174911 0	174945 4	-1	hypothetical protein	
173036 4	1799760	69396	Predicted by at least one method		Predicted_0024 8	174956 0	174982 0	-1	hypothetical protein	
173036 4	1799760	69396	Predicted by at least one method		Predicted_0024 9	174985 1	175008 7	-1	hypothetical protein	
173036 4	1799760	69396	Predicted by at least one method		Predicted_0025 0	175008 4	175035 3	-1	hypothetical protein	
173036 4	1799760	69396	Predicted by at least one method		Predicted_0025 1	175040 4	175060 7	-1	hypothetical protein	
173036 4	1799760	69396	Predicted by at least one method		Predicted_0025 2	175067 1	175111 1	-1	hypothetical protein	
173036 4	1799760	69396	Predicted by at least one method		Predicted_0025 3	175132 0	175152 3	-1	hypothetical protein	
173036 4	1799760	69396	Predicted by at least one method		Predicted_0025 4	175153 3	175186 8	-1	hypothetical protein	
173036 4	1799760	69396	Predicted by at least one method		Predicted_0025 5	175191 4	175214 1	-1	hypothetical protein	

173036 4	1799760	69396	Predicted by at least one method		Predicted_0025 6	175216 6	175239 9	-1	hypothetical protein
173036 4	1799760	69396	Predicted by at least one method		Predicted_0025 7	175256 4	175308 2	-1	hypothetical protein
173036 4	1799760	69396	Predicted by at least one method		Predicted_0025 8	175325 3	175355 2	-1	hypothetical protein
173036 4	1799760	69396	Predicted by at least one method		Predicted_0025 9	175362 8	175429 3	-1	hypothetical protein
173036 4	1799760	69396	Predicted by at least one method		Predicted_0026 0	175436 3	175474 0	-1	hypothetical protein
173036 4	1799760	69396	Predicted by at least one method	ascD_1	Predicted_0026 1	175494 6	175520 3	-1	CDP-6-deoxy-L-threo-D- glycero-4-hexulose-3- dehydrase reductase hypothetical protein
173036 4	1799760	69396	Predicted by at least one method		Predicted_0026 2	175538 5	175568 1	-1	hypothetical protein
173036 4	1799760	69396	Predicted by at least one method		Predicted_0026 3	175588 4	175630 3	-1	hypothetical protein
173036 4	1799760	69396	Predicted by at least one method		Predicted_0026 4	175642 0	175676 7	-1	hypothetical protein
173036 4	1799760	69396	Predicted by at least one method		Predicted_0026 5	175700 0	175760 8	1	hypothetical protein
173036 4	1799760	69396	Predicted by at least one method		Predicted_0026 6	175748 6	175797 4	-1	hypothetical protein
173036 4	1799760	69396	Predicted by at least one method		Predicted_0026 7	175804 3	175847 4	-1	hypothetical protein
173036 4	1799760	69396	Predicted by at least one method		Predicted_0026 8	175948 8	176051 3	1	hypothetical protein
173036 4	1799760	69396	Predicted by at least one method		Predicted_0026 9	176053 0	176106 0	1	hypothetical protein
173036 4	1799760	69396	Predicted by at least one method		Predicted_0027 0	176109 0	176165 3	1	Transglycosylase SLT domain
173036 4	1799760	69396	Predicted by at least one method		Predicted_0027 1	176164 6	176217 9	1	hypothetical protein
173036 4	1799760	69396	Predicted by at least one method	flhC_1	Predicted_0027 2	176217 6	176269 1	1	Flagellar transcriptional regulator FlhC
173036 4	1799760	69396	Predicted by at least one method		Predicted_0027 3	176269 1	176311 0	1	hypothetical protein
173036 4	1799760	69396	Predicted by at least one method		Predicted_0027 4	176325 5	176400 1	1	hypothetical protein

173036 4	1799760	69396	Predicted by at least one method	Predicted_0027 5	176405 4	176676 8	-1	conjugal transfer mating pair stabilization protein TraN
173036 4	1799760	69396	Predicted by at least one method	Predicted_0027 6	176696 1	176713 7	1	hypothetical protein
173036 4	1799760	69396	Predicted by at least one method	Predicted_0027 7	176719 0	177034 8	-1	TraG-like protein, N-terminal region
173036 4	1799760	69396	Predicted by at least one method	Predicted_0027 8	177036 3	177179 9	-1	Conjugative relaxosome accessory transposon protein
173036 4	1799760	69396	Predicted by at least one method	Predicted_0027 9	177182 1	177288 2	-1	hypothetical protein
173036 4	1799760	69396	Predicted by at least one method	Predicted_0028 0	177286 0	177319 8	-1	hypothetical protein
173036 4	1799760	69396	Predicted by at least one method	Predicted_0028 1	177344 8	177362 1	-1	hypothetical protein
173036 4	1799760	69396	Predicted by at least one method	Predicted_0028 2	177368 8	177390 0	-1	hypothetical protein
173036 4	1799760	69396	Predicted by at least one method	Predicted_0028 3	177394 3	177445 2	-1	hypothetical protein
173036 4	1799760	69396	Predicted by at least one method	Predicted_0028 4	177449 0	177487 9	-1	hypothetical protein
173036 4	1799760	69396	Predicted by at least one method	Predicted_0028 5	177491 5	177517 5	-1	hypothetical protein
173036 4	1799760	69396	Predicted by at least one method	Predicted_0028 6	177521 9	177544 6	-1	hypothetical protein
173036 4	1799760	69396	Predicted by at least one method	Predicted_0028 7	177545 0	177576 4	-1	hypothetical protein
173036 4	1799760	69396	Predicted by at least one method	Predicted_0028 8	177582 5	177631 0	-1	hypothetical protein
173036 4	1799760	69396	Predicted by at least one method	Predicted_0028 9	177640 9	177818 7	-1	von Willebrand factor type A domain
173036 4	1799760	69396	Predicted by at least one method	Predicted_0029 0	177823 9	177852 0	-1	hypothetical protein
173036 4	1799760	69396	Predicted by at least one method	Predicted_0029 1	177857 3	177983 8	-1	hypothetical protein
173036 4	1799760	69396	Predicted by at least one method	Predicted_0029 2	177993 8	178099 9	-1	Aerobic cobaltochelatase subunit CobS
173036 4	1799760	69396	Predicted by at least one method	Predicted_0029 3	178106 6	178144 0	-1	hypothetical protein

173036	1799760	69396	Predicted by at least one method	Predicted_0029	178180	178292	-1	YqaJ-like viral recombinase domain
4				4	8	3		
173036	1799760	69396	Predicted by at least one method	Predicted_0029	178298	178452	-1	recombination and repair protein RecT
4				5	3	7		
173036	1799760	69396	Predicted by at least one method	Predicted_0029	178461	178530	-1	Single-strand binding protein family
4				6	3	2		
173036	1799760	69396	Predicted by at least one method	Predicted_0029	178577	178620	-1	hypothetical protein
4				7	0	7		
173036	1799760	69396	Predicted by at least one method	Predicted_0029	178628	178689	-1	hypothetical protein
4				8	6	4		
173036	1799760	69396	Predicted by at least one method	Predicted_0029	178740	178815	1	hypothetical protein
4				9	4	3		
173036	1799760	69396	Predicted by at least one method	Predicted_0030	178829	178882	1	Sel1 repeat
4				0	7	7		
173036	1799760	69396	Predicted by at least one method	Predicted_0030	178887	178996	-1	TraU protein
4				1	8	3		
173036	1799760	69396	Predicted by at least one method	Predicted_0030	178996	179121	-1	Type-F conjugative transfer system pilin assembly protein
4				2	0	9		
173036	1799760	69396	Predicted by at least one method	Predicted_0030	179121	179173	-1	signal peptidase I
4				3	3	7		
173036	1799760	69396	Predicted by at least one method	Predicted_0030	179175	179203	-1	hypothetical protein
4				4	5	0		
173036	1799760	69396	Predicted by at least one method	Predicted_0030	179202	179451	-1	AAA-like domain
4				5	7	6		
173036	1799760	69396	Predicted by at least one method	Predicted_0030	179454	179528	-1	putative thiol:disulfide interchange protein DsbC precursor
4				6	0	3		
173036	1799760	69396	Predicted by at least one method	Predicted_0030	179569	179622	-1	Type IV conjugative transfer system lipoprotein (TraV)
4				7	1	4		
173036	1799760	69396	Predicted by at least one method	Predicted_0030	179622	179761	-1	Bacterial conjugation TrbI-like protein
4				8	7	5		
173036	1799760	69396	Predicted by at least one method	Predicted_0030	179762	179871	-1	TraK protein
4				9	3	7		
173036	1799760	69396	Predicted by at least one method	Predicted_0031	179888	179947	-1	TraE protein
4				0	6	6		
173036	1799760	69396	Predicted by at least one method	Predicted_0031	179947	179976	-1	TraL protein
4				1	3	0		
184302	1848130	5101	Predicted by at least one method	Predicted_0036	184302	184474	-1	hypothetical protein

9			one method		4	9	4		
184302	1848130	5101	Predicted by at least		Predicted_0036	184486	184547	-1	hypothetical protein
9			one method		5	9	7		
184302	1848130	5101	Predicted by at least		Predicted_0036	184555	184600	-1	hypothetical protein
9			one method		6	0	2		
184302	1848130	5101	Predicted by at least		Predicted_0036	184601	184648	-1	hypothetical protein
9			one method		7	5	2		
184302	1848130	5101	Predicted by at least		Predicted_0036	184657	184721	-1	hypothetical protein
9			one method		8	1	8		
184302	1848130	5101	Predicted by at least		Predicted_0036	184725	184762	-1	hypothetical protein
9			one method		9	6	4		
184302	1848130	5101	Predicted by at least		Predicted_0037	184762	184813	-1	hypothetical protein
9			one method		0	1	0		
184555	1871329	25779	Predicted by at least		Predicted_0036	184555	184600	-1	hypothetical protein
0			one method		6	0	2		
184555	1871329	25779	Predicted by at least		Predicted_0036	184601	184648	-1	hypothetical protein
0			one method		7	5	2		
184555	1871329	25779	Predicted by at least		Predicted_0036	184657	184721	-1	hypothetical protein
0			one method		8	1	8		
184555	1871329	25779	Predicted by at least		Predicted_0036	184725	184762	-1	hypothetical protein
0			one method		9	6	4		
184555	1871329	25779	Predicted by at least		Predicted_0037	184762	184813	-1	hypothetical protein
0			one method		0	1	0		
184555	1871329	25779	Predicted by at least	rdgC_1	Predicted_0037	184818	184942	-1	Recombination-associated
0			one method		1	4	2		protein RdgC
184555	1871329	25779	Predicted by at least	dam_1	Predicted_0037	184941	185024	-1	DNA adenine methylase
0			one method		2	9	0		
184555	1871329	25779	Predicted by at least		Predicted_0037	185029	185093	-1	hypothetical protein
0			one method		3	5	6		
184555	1871329	25779	Predicted by at least	dnaN_1	Predicted_0037	185116	185243	-1	DNA polymerase III subunit
0			one method		4	3	1		beta
184555	1871329	25779	Predicted by at least		Predicted_0037	185258	185291	-1	hypothetical protein
0			one method		5	0	2		
184555	1871329	25779	Predicted by at least		Predicted_0037	185290	185337	-1	hypothetical protein
0			one method		6	9	0		
184555	1871329	25779	Predicted by at least		Predicted_0037	185342	185394	-1	hypothetical protein
0			one method		7	5	6		
184555	1871329	25779	Predicted by at least		Predicted_0037	185405	185447	-1	hypothetical protein

0			one method		8	3	8		
184555	1871329	25779	Predicted by at least		Predicted_0037	185464	185493	1	hypothetical protein
0			one method		9	0	9		
184555	1871329	25779	Predicted by at least		Predicted_0038	185497	185530	-1	hypothetical protein
0			one method		0	8	4		
184555	1871329	25779	Predicted by at least		Predicted_0038	185533	185581	-1	hypothetical protein
0			one method		1	5	7		
184555	1871329	25779	Predicted by at least		Predicted_0038	185591	185611	1	hypothetical protein
0			one method		2	5	5		
184555	1871329	25779	Predicted by at least		Predicted_0038	185623	185658	-1	hypothetical protein
0			one method		3	8	8		
184555	1871329	25779	Predicted by at least	ssb_1	Predicted_0038	185665	185717	-1	Single-stranded DNA-binding protein
0			one method		4	6	7		
184555	1871329	25779	Predicted by at least		Predicted_0038	185742	185785	-1	hypothetical protein
0			one method		5	6	7		
184555	1871329	25779	Predicted by at least		Predicted_0038	185786	185815	-1	hypothetical protein
0			one method		6	6	0		
184555	1871329	25779	Predicted by at least	hflK_1	Predicted_0038	185826	185918	-1	Modulator of FtsH protease HflK
0			one method		7	1	1		
184555	1871329	25779	Predicted by at least		Predicted_0038	185920	185963	-1	NfeD-like C-terminal, partner-binding
0			one method		8	5	6		
184555	1871329	25779	Predicted by at least		Predicted_0038	185992	186008	-1	hypothetical protein
0			one method		9	7	8		
184555	1871329	25779	Predicted by at least		Predicted_0039	186013	186104	-1	hypothetical protein
0			one method		0	9	4		
184555	1871329	25779	Predicted by at least		Predicted_0039	186104	186153	-1	hypothetical protein
0			one method		1	1	5		
184555	1871329	25779	Predicted by at least		Predicted_0039	186157	186220	-1	hypothetical protein
0			one method		2	0	8		
184555	1871329	25779	Predicted by at least		Predicted_0039	186220	186295	-1	hypothetical protein
0			one method		3	5	7		
184555	1871329	25779	Predicted by at least		Predicted_0039	186324	186350	-1	hypothetical protein
0			one method		4	7	4		
184555	1871329	25779	Predicted by at least		Predicted_0039	186381	186588	-1	AAA-like domain
0			one method		5	9	2		
184555	1871329	25779	Predicted by at least		Predicted_0039	186587	186727	-1	hypothetical protein
0			one method		6	9	0		
184555	1871329	25779	Predicted by at least	tnpR_1	Predicted_0039	186734	186792	-1	Transposon gamma-delta

0			one method		7	7	8		resolvase
184555	1871329	25779	Predicted by at least		Predicted_0039	186811	186846	1	hypothetical protein
0			one method		8	3	3		
184555	1871329	25779	Predicted by at least	higA_1	Predicted_0039	186846	186876	1	Antitoxin HigA
0			one method		9	5	4		
184555	1871329	25779	Predicted by at least		Predicted_0040	186890	186992	1	hypothetical protein
0			one method		0	5	1		
184555	1871329	25779	Predicted by at least		Predicted_0040	186997	187037	-1	hypothetical protein
0			one method		1	2	3		
184555	1871329	25779	Predicted by at least		Predicted_0040	187080	187132	-1	LexA repressor
0			one method		2	5	9		
188035	1892698	12339	Predicted by at least	comR_1	Predicted_0041	188035	188094	-1	HTH-type transcriptional repressor ComR
9			one method		7	9	0		
188035	1892698	12339	Predicted by at least		Predicted_0041	188108	188166	-1	hypothetical protein
9			one method		8	9	1		
188035	1892698	12339	Predicted by at least		Predicted_0041	188179	188248	-1	hypothetical protein
9			one method		9	7	9		
188035	1892698	12339	Predicted by at least	tus_1	Predicted_0042	188254	188342	-1	DNA replication terminus site-binding protein
9			one method		0	5	6		
188035	1892698	12339	Predicted by at least		Predicted_0042	188383	188582	-1	Tn3 transposase DDE domain
9			one method		1	3	1		
188035	1892698	12339	Predicted by at least		Predicted_0042	188587	188609	1	hypothetical protein
9			one method		2	0	4		
188035	1892698	12339	Predicted by at least		Predicted_0017	188808	188847	1	Transposase
9			one method		9	7	0		
188035	1892698	12339	Predicted by at least		Predicted_0018	188923	188996	1	Tn3 transposase DDE domain
9			one method		0	2	0		
188035	1892698	12339	Predicted by at least	aacA4_1	Predicted_0052	189135	189195	-1	Aminoglycoside N(6')-acetyltransferase type 1
9			one method		7	3	8		
188035	1892698	12339	Predicted by at least	tnpR_2	Predicted_0052	189214	189269	-1	Transposon Tn3 resolvase
9			one method		6	1	8		
188923	1896303	7071	Predicted by at least		Predicted_0018	188923	188996	1	Tn3 transposase DDE domain
2			one method		0	2	0		
188923	1896303	7071	Predicted by at least	aacA4_1	Predicted_0052	189135	189195	-1	Aminoglycoside N(6')-acetyltransferase type 1
2			one method		7	3	8		
188923	1896303	7071	Predicted by at least	tnpR_2	Predicted_0052	189214	189269	-1	Transposon Tn3 resolvase
2			one method		6	1	8		
188923	1896303	7071	Predicted by at least		Predicted_0052	189286	189586	1	Tn3 transposase DDE domain

2			one method		5	0	5		
188923	1896303	7071	Predicted by at least		Predicted_0052	189621	189630	1	hypothetical protein
2			one method		4	4	3		
188923	1896303	7071	Predicted by at least	dmlR_1	Predicted_0052	189628	189715	-1	HTH-type transcriptional
2			one method		3	1	0		regulator DmlR
190358	1915816	12231	Predicted by at least		Predicted_0051	190358	190380	1	hypothetical protein
5			one method		6	5	9		
190358	1915816	12231	Predicted by at least		Predicted_0051	190435	190464	1	Transposase
5			one method		5	7	1		
190358	1915816	12231	Predicted by at least		Predicted_0051	190467	190549	1	Integrase core domain
5			one method		4	1	5		
190358	1915816	12231	Predicted by at least		Predicted_0051	190588	190652	-1	DNA replication terminus site-
5			one method		3	5	6		binding protein
190358	1915816	12231	Predicted by at least		Predicted_0051	190661	190671	-1	hypothetical protein
5			one method		2	5	0		
190358	1915816	12231	Predicted by at least		Predicted_0051	190678	190701	-1	hypothetical protein
5			one method		1	6	3		
190358	1915816	12231	Predicted by at least	pir	Predicted_0051	190701	190785	-1	PI protein
5			one method		0	6	2		
190358	1915816	12231	Predicted by at least		Predicted_0050	190918	190967	-1	hypothetical protein
5			one method		9	8	9		
190358	1915816	12231	Predicted by at least		Predicted_0050	190969	191007	-1	Plasmid stability protein
5			one method		8	0	0		
190358	1915816	12231	Predicted by at least	parM	Predicted_0050	191007	191103	-1	Plasmid segregation protein
5			one method		7	5	4		ParM
190358	1915816	12231	Predicted by at least		Predicted_0050	191133	191214	-1	hypothetical protein
5			one method		6	1	3		
190358	1915816	12231	Predicted by at least		Predicted_0050	191246	191288	1	hypothetical protein
5			one method		5	7	6		
190358	1915816	12231	Predicted by at least		Predicted_0050	191322	191368	-1	hypothetical protein
5			one method		4	2	0		
190358	1915816	12231	Predicted by at least		Predicted_0050	191416	191471	1	Transglycosylase SLT domain
5			one method		3	7	5		
190358	1915816	12231	Predicted by at least		Predicted_0050	191473	191525	1	hypothetical protein
5			one method		2	8	9		
190358	1915816	12231	Predicted by at least		Predicted_0050	191525	191581	1	transcriptional activator FlhC
5			one method		1	6	6		
190435	1910070	5713	Predicted by at least		Predicted_0051	190435	190464	1	Transposase

7			one method		5	7	1	
190435	1910070	5713	Predicted by at least		Predicted_0051	190467	190549	1
7			one method		4	1	5	
190435	1910070	5713	Predicted by at least		Predicted_0051	190588	190652	-1
7			one method		3	5	6	
190435	1910070	5713	Predicted by at least		Predicted_0051	190661	190671	-1
7			one method		2	5	0	
190435	1910070	5713	Predicted by at least		Predicted_0051	190678	190701	-1
7			one method		1	6	3	
190435	1910070	5713	Predicted by at least	pir	Predicted_0051	190701	190785	-1
7			one method		0	6	2	
190435	1910070	5713	Predicted by at least		Predicted_0050	190918	190967	-1
7			one method		9	8	9	
190435	1910070	5713	Predicted by at least		Predicted_0050	190969	191007	-1
7			one method		8	0	0	
192318	1932903	9717	Predicted by at least		Predicted_0049	192318	192459	-1
6			one method		6	6	5	
192318	1932903	9717	Predicted by at least		Predicted_0049	192461	192564	-1
6			one method		5	1	2	
192318	1932903	9717	Predicted by at least		Predicted_0049	192587	192765	-1
6			one method		4	8	3	
192318	1932903	9717	Predicted by at least		Predicted_0049	192771	192890	-1
6			one method		3	8	5	
192318	1932903	9717	Predicted by at least	cobS_2	Predicted_0049	192900	193001	-1
6			one method		2	4	1	
192318	1932903	9717	Predicted by at least		Predicted_0049	193006	193043	-1
6			one method		1	5	3	
192318	1932903	9717	Predicted by at least		Predicted_0049	193047	193156	-1
6			one method		0	1	5	
192318	1932903	9717	Predicted by at least		Predicted_0048	193161	193290	-1
6			one method		9	4	3	
196168	2089481	12779	Predicted by at least		Predicted_0045	196168	196437	1
3		8	one method		8	3	0	
196168	2089481	12779	Predicted by at least		Predicted_0045	196440	196643	1
3		8	one method		7	2	2	
196168	2089481	12779	Predicted by at least		Predicted_0045	196643	196769	1
3		8	one method		6	6	5	
196168	2089481	12779	Predicted by at least	mrr_1	Predicted_0045	196770	196870	1

3		8	one method		5	8	6		
196168	2089481	12779	Predicted by at least		Predicted_0045	196887	197142	-1	hypothetical protein
3		8	one method		4	7	9		
196168	2089481	12779	Predicted by at least		Predicted_0045	197157	197204	1	hypothetical protein
3		8	one method		3	8	5		
196168	2089481	12779	Predicted by at least		Predicted_0045	197204	197249	1	hypothetical protein
3		8	one method		2	2	1		
196168	2089481	12779	Predicted by at least		Predicted_0045	197253	197296	1	hypothetical protein
3		8	one method		1	0	1		
196168	2089481	12779	Predicted by at least		Predicted_0045	197302	197322	-1	Prophage CP4-57 regulatory protein (AlpA)
3		8	one method		0	0	9		GTP-binding protein EngB
196168	2089481	12779	Predicted by at least	engB_1	Predicted_0044	197330	197429	1	
3		8	one method		9	7	6		
196168	2089481	12779	Predicted by at least		Predicted_0044	197440	197485	1	hypothetical protein
3		8	one method		8	6	2		
196168	2089481	12779	Predicted by at least		Predicted_0044	197489	197553	-1	hypothetical protein
3		8	one method		7	4	8		
196168	2089481	12779	Predicted by at least		Predicted_0044	197598	197691	1	hypothetical protein
3		8	one method		6	5	7		
196168	2089481	12779	Predicted by at least		Predicted_0044	197714	197747	-1	Resolvase, N terminal domain
3		8	one method		5	2	1		
196168	2089481	12779	Predicted by at least		Predicted_0044	197763	198062	1	Tn3 transposase DDE domain
3		8	one method		4	9	6		
196168	2089481	12779	Predicted by at least		Predicted_0044	198089	198243	1	DNA-dependent helicase II
3		8	one method		3	8	3		
196168	2089481	12779	Predicted by at least		Predicted_0044	198252	198360	-1	hypothetical protein
3		8	one method		2	9	5		
196168	2089481	12779	Predicted by at least		Predicted_0044	198365	198421	-1	hypothetical protein
3		8	one method		1	9	3		
196168	2089481	12779	Predicted by at least		Predicted_0044	198427	198523	-1	hypothetical protein
3		8	one method		0	2	4		
196168	2089481	12779	Predicted by at least		Predicted_0043	198532	198625	-1	Restriction endonuclease
3		8	one method		9	4	9		
196168	2089481	12779	Predicted by at least		Predicted_0043	198627	198684	-1	hypothetical protein
3		8	one method		8	7	6		
196168	2089481	12779	Predicted by at least		Predicted_0043	198683	198718	-1	hypothetical protein
3		8	one method		7	3	3		
196168	2089481	12779	Predicted by at least	sipT	Predicted_0043	198717	198791	-1	Signal peptidase I T

3		8	one method		6	0	9		
196168	2089481	12779	Predicted by at least		Predicted_0043	198800	198842	-1	hypothetical protein
3		8	one method		5	0	2		
196168	2089481	12779	Predicted by at least		Predicted_0043	198841	198942	-1	hypothetical protein
3		8	one method		4	2	2		
196168	2089481	12779	Predicted by at least		Predicted_0043	198944	199026	-1	hypothetical protein
3		8	one method		3	3	1		
196168	2089481	12779	Predicted by at least	xerC_1	Predicted_0043	199027	199122	-1	Tyrosine recombinase XerC
3		8	one method		2	5	8		
196168	2089481	12779	Predicted by at least		Predicted_0043	199152	199179	1	hypothetical protein
3		8	one method		1	3	2		
196168	2089481	12779	Predicted by at least		Predicted_0043	199183	199208	1	hypothetical protein
3		8	one method		0	1	5		
196168	2089481	12779	Predicted by at least		Predicted_0042	199218	199264	1	hypothetical protein
3		8	one method		9	2	0		
196168	2089481	12779	Predicted by at least		Predicted_0042	199296	199310	1	small toxic polypeptide
3		8	one method		8	1	4		
196168	2089481	12779	Predicted by at least		Predicted_0042	199319	199357	-1	hypothetical protein
3		8	one method		7	1	1		
196168	2089481	12779	Predicted by at least	mazF_1	Predicted_0042	199360	199393	-1	mRNA interferase MazF
3		8	one method		6	2	4		
196168	2089481	12779	Predicted by at least	mazE	Predicted_0042	199393	199417	-1	Antitoxin MazE
3		8	one method		5	4	9		
196168	2089481	12779	Predicted by at least	parA_2	Predicted_0042	199448	199511	1	Chromosome partitioning protein ParA
3		8	one method		4	9	2		
196168	2089481	12779	Predicted by at least		Predicted_0042	199523	199546	1	ParG
3		8	one method		3	7	4		
196168	2089481	12779	Predicted by at least		Predicted_0434	199758	199781	1	hypothetical protein
3		8	one method		2	0	9		
196168	2089481	12779	Predicted by at least	xerC_6	Predicted_0434	199844	199898	-1	Tyrosine recombinase XerC
3		8	one method		3	8	4		
196168	2089481	12779	Predicted by at least		Predicted_0119	200098	200148	-1	hypothetical protein
3		8	one method		4	3	0		
196168	2089481	12779	Predicted by at least		Predicted_0119	200148	200197	-1	hypothetical protein
3		8	one method		3	3	1		
196168	2089481	12779	Predicted by at least		Predicted_0119	200206	200240	1	hypothetical protein
3		8	one method		2	8	3		
196168	2089481	12779	Predicted by at least		Predicted_0119	200241	200288	-1	hypothetical protein

3		8	one method		1	8	8		
196168	2089481	12779	Predicted by at least		Predicted_0119	200288	200325	-1	hypothetical protein
3		8	one method		0	1	2		
196168	2089481	12779	Predicted by at least		Predicted_0118	200326	200345	-1	hypothetical protein
3		8	one method		9	3	7		
196168	2089481	12779	Predicted by at least		Predicted_0118	200345	200370	-1	hypothetical protein
3		8	one method		8	7	8		
196168	2089481	12779	Predicted by at least		Predicted_0118	200379	200434	-1	hypothetical protein
3		8	one method		7	8	6		
196168	2089481	12779	Predicted by at least		Predicted_0118	200450	200485	1	hypothetical protein
3		8	one method		6	9	9		
196168	2089481	12779	Predicted by at least	higA_4	Predicted_0118	200486	200516	1	Antitoxin HigA
3		8	one method		5	4	6		
196168	2089481	12779	Predicted by at least		Predicted_0118	200519	200548	-1	hypothetical protein
3		8	one method		4	3	6		
196168	2089481	12779	Predicted by at least	hupB_1	Predicted_0118	200557	200584	-1	DNA-binding protein HU-beta
3		8	one method		3	4	6		
196168	2089481	12779	Predicted by at least	yncB	Predicted_0118	200590	200643	-1	Endonuclease precursor YncB
3		8	one method		2	4	1		
196168	2089481	12779	Predicted by at least		Predicted_0118	200644	200663	-1	hypothetical protein
3		8	one method		1	8	6		
196168	2089481	12779	Predicted by at least		Predicted_0118	200666	200751	-1	hypothetical protein
3		8	one method		0	2	9		
196168	2089481	12779	Predicted by at least		Predicted_0117	200750	200773	-1	hypothetical protein
3		8	one method		9	6	3		
196168	2089481	12779	Predicted by at least		Predicted_0117	200773	200825	-1	hypothetical protein
3		8	one method		8	6	4		
196168	2089481	12779	Predicted by at least		Predicted_0117	200825	200869	-1	hypothetical protein
3		8	one method		7	1	7		
196168	2089481	12779	Predicted by at least		Predicted_0117	200869	200905	-1	hypothetical protein
3		8	one method		6	7	6		
196168	2089481	12779	Predicted by at least		Predicted_0117	200911	200954	-1	hypothetical protein
3		8	one method		5	3	1		
196168	2089481	12779	Predicted by at least	bdbD_2	Predicted_0117	200957	201043	-1	Disulfide bond formation protein D precursor
3		8	one method		4	5	5		
196168	2089481	12779	Predicted by at least	sppA_1	Predicted_0117	201045	201141	-1	Putative signal peptide
3		8	one method		3	1	0		
196168	2089481	12779	Predicted by at least		Predicted_0117	201141	201226	-1	peptidase SppA hypothetical protein

3		8	one method	2	0	4		
196168	2089481	12779	Predicted by at least	Predicted_0117	201226	201256	-1	hypothetical protein
3		8	one method	1	9	2		
196168	2089481	12779	Predicted by at least	Predicted_0117	201257	201304	-1	hypothetical protein
3		8	one method	0	3	6		
196168	2089481	12779	Predicted by at least	Predicted_0116	201314	201366	-1	hypothetical protein
3		8	one method	9	3	4		
196168	2089481	12779	Predicted by at least	Predicted_0116	201366	201418	-1	hypothetical protein
3		8	one method	8	7	8		
196168	2089481	12779	Predicted by at least	Predicted_0116	201419	201480	-1	hypothetical protein
3		8	one method	7	3	1		
196168	2089481	12779	Predicted by at least	Predicted_0116	201507	201618	1	hypothetical protein
3		8	one method	6	0	5		
196168	2089481	12779	Predicted by at least	Predicted_0116	201620	201663	1	hypothetical protein
3		8	one method	5	3	7		
196168	2089481	12779	Predicted by at least	Predicted_0116	201704	201732	-1	hypothetical protein
3		8	one method	4	7	8		
196168	2089481	12779	Predicted by at least	Predicted_0116	201767	201873	-1	hypothetical protein
3		8	one method	3	4	8		
196168	2089481	12779	Predicted by at least	Predicted_0116	201875	201904	-1	hypothetical protein
3		8	one method	2	9	6		
196168	2089481	12779	Predicted by at least	Predicted_0116	201905	201959	-1	hypothetical protein
3		8	one method	1	1	0		
196168	2089481	12779	Predicted by at least	Predicted_0116	202008	202107	1	hypothetical protein
3		8	one method	0	9	2		
196168	2089481	12779	Predicted by at least	Predicted_0115	202108	202138	1	hypothetical protein
3		8	one method	9	9	2		
196168	2089481	12779	Predicted by at least	Predicted_0115	202138	202180	1	H-NS histone family
3		8	one method	8	4	3		
196168	2089481	12779	Predicted by at least	Predicted_0115	202186	202241	-1	transcriptional activator FlhC
3		8	one method	7	3	4		
196168	2089481	12779	Predicted by at least	Predicted_0115	202241	202307	-1	hypothetical protein
3		8	one method	6	1	6		
196168	2089481	12779	Predicted by at least	Predicted_0115	202303	202356	-1	Transglycosylase SLT domain
3		8	one method	5	0	3		
196168	2089481	12779	Predicted by at least	Predicted_0115	202356	202383	-1	DNA-binding transcriptional
3		8	one method	4	3	5		regulator Nlp
196168	2089481	12779	Predicted by at least	Predicted_0115	202455	202491	1	hypothetical protein

3		8	one method		3	1	3		
196168	2089481	12779	Predicted by at least		Predicted_0115	202495	202856	-1	TraG-like protein, N-terminal
3		8	one method		2	1	5		region
196168	2089481	12779	Predicted by at least		Predicted_0115	202857	203001	-1	Conjugative relaxosome
3		8	one method		1	8	1		accessory transposon protein
196168	2089481	12779	Predicted by at least		Predicted_0115	203001	203104	-1	hypothetical protein
3		8	one method		0	3	1		
196168	2089481	12779	Predicted by at least		Predicted_0114	203116	203267	-1	DNA helicase IV
3		8	one method		9	4	5		
196168	2089481	12779	Predicted by at least		Predicted_0114	203268	203296	-1	hypothetical protein
3		8	one method		8	4	2		
196168	2089481	12779	Predicted by at least	smc_2	Predicted_0114	203296	203400	-1	Chromosome partition protein
3		8	one method		7	2	2		Smc
196168	2089481	12779	Predicted by at least	tus_3	Predicted_0114	203415	203503	-1	DNA replication terminus site-
3		8	one method		6	5	0		binding protein
196168	2089481	12779	Predicted by at least		Predicted_0114	203535	203584	-1	hypothetical protein
3		8	one method		5	7	8		
196168	2089481	12779	Predicted by at least	dnaQ_1	Predicted_0114	203584	203671	-1	DNA polymerase III subunit
3		8	one method		4	5	4		epsilon
196168	2089481	12779	Predicted by at least	xerD_2	Predicted_0114	203671	203755	-1	Tyrosine recombinase XerD
3		8	one method		3	9	2		
196168	2089481	12779	Predicted by at least		Predicted_0114	203773	203826	-1	hypothetical protein
3		8	one method		2	2	8		
196168	2089481	12779	Predicted by at least		Predicted_0114	203826	203854	-1	hypothetical protein
3		8	one method		1	1	8		
196168	2089481	12779	Predicted by at least		Predicted_0114	203856	203888	-1	hypothetical protein
3		8	one method		0	7	4		
196168	2089481	12779	Predicted by at least		Predicted_0113	203911	203971	1	hypothetical protein
3		8	one method		9	0	2		
196168	2089481	12779	Predicted by at least		Predicted_0113	203972	204018	-1	hypothetical protein
3		8	one method		8	8	0		
196168	2089481	12779	Predicted by at least		Predicted_0113	204034	204064	-1	hypothetical protein
3		8	one method		7	7	6		
196168	2089481	12779	Predicted by at least		Predicted_0113	204094	204121	-1	hypothetical protein
3		8	one method		6	1	3		
196168	2089481	12779	Predicted by at least		Predicted_0113	204151	204244	-1	Transposase DDE domain
3		8	one method		5	7	9		
196168	2089481	12779	Predicted by at least		Predicted_0113	204253	204286	1	hypothetical protein

3		8	one method		4	8	4		
196168	2089481	12779	Predicted by at least	dinB_3	Predicted_0113	204290	204417	-1	DNA polymerase IV
3		8	one method		3	0	7		
196168	2089481	12779	Predicted by at least	lexA_3	Predicted_0113	204418	204461	-1	LexA repressor
3		8	one method		2	3	1		
196168	2089481	12779	Predicted by at least		Predicted_0113	204470	204497	-1	Helix-turn-helix domain
3		8	one method		1	7	9		
196168	2089481	12779	Predicted by at least		Predicted_0113	204510	204545	1	hypothetical protein
3		8	one method		0	4	4		
196168	2089481	12779	Predicted by at least		Predicted_0112	204547	204586	1	Transposase DDE domain
3		8	one method		9	5	4		
196168	2089481	12779	Predicted by at least		Predicted_0112	204593	204648	1	hypothetical protein
3		8	one method		8	2	3		
196168	2089481	12779	Predicted by at least		Predicted_0112	204670	204694	-1	hypothetical protein
3		8	one method		7	5	4		
196168	2089481	12779	Predicted by at least		Predicted_0112	204700	204783	-1	Integrase core domain
3		8	one method		6	7	7		
196168	2089481	12779	Predicted by at least		Predicted_0112	204783	204814	-1	Transposase
3		8	one method		5	4	5		
196168	2089481	12779	Predicted by at least	tmrB	Predicted_0112	204819	204849	-1	Tunicamycin resistance protein
3		8	one method		4	8	1		
196168	2089481	12779	Predicted by at least	yokD	Predicted_0112	204850	204936	-1	SPBc2 prophage-derived aminoglycoside N(3')-acetyltransferase-like protein YokD
3		8	one method		3	4	4		
196168	2089481	12779	Predicted by at least	bla_1	Predicted_0112	204950	205036	-1	Beta-lactamase TEM precursor
3		8	one method		2	6	6		
196168	2089481	12779	Predicted by at least	tnpR_4	Predicted_0112	205054	205110	-1	Transposon Tn3 resolvase
3		8	one method		1	9	6		
196168	2089481	12779	Predicted by at least		Predicted_0112	205127	205427	1	Tn3 transposase DDE domain
3		8	one method		0	0	8		
196168	2089481	12779	Predicted by at least		Predicted_0009	205667	205821	1	Putative transposase
3		8	one method		7	8	9		
196168	2089481	12779	Predicted by at least	trpF	Predicted_0009	205877	205924	-1	N-(5'-phosphoribosyl)anthranilate isomerase
3		8	one method		8	5	5		
196168	2089481	12779	Predicted by at least		Predicted_0009	205935	205990	1	hypothetical protein
3		8	one method		9	5	6		

196168 3	2089481	12779 8	Predicted by at least one method	blaNDM -1	Predicted_0010 0	205978 7	206059 9	-1	Beta-lactamase precursor	NDM-1
196168 3	2089481	12779 8	Predicted by at least one method		Predicted_0010 1	206070 0	206171 0	-1	Integrase core domain	
196168 3	2089481	12779 8	Predicted by at least one method	aphA_1	Predicted_0010 2	206187 7	206265 6	-1	Aminoglycoside phosphotransferase	3'-
196168 3	2089481	12779 8	Predicted by at least one method		Predicted_0010 3	206276 2	206299 2	-1	hypothetical protein	
196168 3	2089481	12779 8	Predicted by at least one method		Predicted_0010 4	206298 2	206342 5	-1	Transposase	
196168 3	2089481	12779 8	Predicted by at least one method		Predicted_0010 5	206371 6	206597 7	-1	Tn3 transposase DDE domain	
196168 3	2089481	12779 8	Predicted by at least one method	hin	Predicted_0010 6	206598 0	206661 2	-1	DNA-invertase hin	
196168 3	2089481	12779 8	Predicted by at least one method	ttgC	Predicted_0010 7	206666 8	206815 5	-1	Toluene efflux pump outer membrane protein TtgC precursor	
196168 3	2089481	12779 8	Predicted by at least one method		Predicted_0010 8	206814 2	206861 5	-1	Cytochrome C'	
196168 3	2089481	12779 8	Predicted by at least one method	lolD_1	Predicted_0010 9	206864 4	206935 4	-1	Lipoprotein-releasing system ATP-binding protein LolD	
196168 3	2089481	12779 8	Predicted by at least one method		Predicted_0011 0	206935 9	207056 1	-1	FtsX-like permease family	
196168 3	2089481	12779 8	Predicted by at least one method	mdtA_1	Predicted_0011 1	207055 8	207170 9	-1	Multidrug resistance protein MdtA precursor	
196168 3	2089481	12779 8	Predicted by at least one method	slmA_1	Predicted_0011 2	207170 6	207231 4	-1	Nucleoid occlusion factor SlmA	
196168 3	2089481	12779 8	Predicted by at least one method		Predicted_0011 3	207239 5	207306 9	-1	hypothetical protein	
196168 3	2089481	12779 8	Predicted by at least one method		Predicted_0011 4	207362 4	207397 1	1	hypothetical protein	
196168 3	2089481	12779 8	Predicted by at least one method		Predicted_0011 5	207401 6	207704 2	-1	Tn3 transposase DDE domain	
196168 3	2089481	12779 8	Predicted by at least one method	bin3_1	Predicted_0011 6	207720 4	207784 8	1	Putative transposon Tn552 DNA-invertase bin3	
196168 3	2089481	12779 8	Predicted by at least one method	betT_1	Predicted_0011 7	207803 2	207836 1	1	High-affinity choline transport protein	

196168 3	2089481	12779 8	Predicted by at least one method	merR	Predicted_0011 8	207835 1	207880 6	-1	Mercuric resistance operon regulatory protein
196168 3	2089481	12779 8	Predicted by at least one method		Predicted_0011 9	207887 8	207924 3	1	MerT mercuric transport protein
196168 3	2089481	12779 8	Predicted by at least one method	merP	Predicted_0012 0	207925 9	207953 4	1	Mercuric transport protein periplasmic component precursor
196168 3	2089481	12779 8	Predicted by at least one method	merC	Predicted_0012 1	207956 2	207998 7	1	Mercuric resistance protein MerC
196168 3	2089481	12779 8	Predicted by at least one method	merA	Predicted_0012 2	208002 6	208171 1	1	Mercuric reductase
196168 3	2089481	12779 8	Predicted by at least one method		Predicted_0012 3	208172 9	208209 4	1	zinc-responsive transcriptional regulator
196168 3	2089481	12779 8	Predicted by at least one method		Predicted_0012 4	208209 1	208232 7	1	MerE protein
196168 3	2089481	12779 8	Predicted by at least one method		Predicted_0012 5	208239 3	208260 5	1	hypothetical protein
196168 3	2089481	12779 8	Predicted by at least one method		Predicted_0012 6	208286 1	208368 2	1	hypothetical protein
196168 3	2089481	12779 8	Predicted by at least one method		Predicted_0012 7	208414 2	208463 6	1	hypothetical protein
196168 3	2089481	12779 8	Predicted by at least one method	folP_1	Predicted_0012 8	208474 2	208558 1	-1	Dihydropteroate synthase
196168 3	2089481	12779 8	Predicted by at least one method	emrE_1	Predicted_0012 9	208557 5	208592 2	-1	Multidrug transporter EmrE
196168 3	2089481	12779 8	Predicted by at least one method	ant1_1	Predicted_0013 0	208608 6	208686 5	-1	Streptomycin 3"- adenylyltransferase
196168 3	2089481	12779 8	Predicted by at least one method	dfrA	Predicted_0013 1	208728 5	208778 2	-1	Dihydrofolate reductase
196168 3	2089481	12779 8	Predicted by at least one method	xerD_1	Predicted_0013 2	208792 7	208894 0	1	Tyrosine recombinase XerD
196168 3	2089481	12779 8	Predicted by at least one method		Predicted_0013 3	208892 7	208948 1	-1	hypothetical protein
197302 0	1980626	7606	Predicted by at least one method		Predicted_0045 0	197302 0	197322 9	-1	Prophage CP4-57 regulatory protein (AlpA)
197302 0	1980626	7606	Predicted by at least one method	engB_1	Predicted_0044 9	197330 7	197429 6	1	GTP-binding protein EngB
197302	1980626	7606	Predicted by at least		Predicted_0044	197440	197485	1	hypothetical protein

0			one method		8	6	2		
197302	1980626	7606	Predicted by at least		Predicted_0044	197489	197553	-1	hypothetical protein
0			one method		7	4	8		
197302	1980626	7606	Predicted by at least		Predicted_0044	197598	197691	1	hypothetical protein
0			one method		6	5	7		
197302	1980626	7606	Predicted by at least		Predicted_0044	197714	197747	-1	Resolvase, N terminal domain
0			one method		5	2	1		
197302	1980626	7606	Predicted by at least		Predicted_0044	197763	198062	1	Tn3 transposase DDE domain
0			one method		4	9	6		
198532	1990261	4937	Predicted by at least		Predicted_0043	198532	198625	-1	Restriction endonuclease
4			one method		9	4	9		
198532	1990261	4937	Predicted by at least		Predicted_0043	198627	198684	-1	hypothetical protein
4			one method		8	7	6		
198532	1990261	4937	Predicted by at least		Predicted_0043	198683	198718	-1	hypothetical protein
4			one method		7	3	3		
198532	1990261	4937	Predicted by at least	sipT	Predicted_0043	198717	198791	-1	Signal peptidase I T
4			one method		6	0	9		
198532	1990261	4937	Predicted by at least		Predicted_0043	198800	198842	-1	hypothetical protein
4			one method		5	0	2		
198532	1990261	4937	Predicted by at least		Predicted_0043	198841	198942	-1	hypothetical protein
4			one method		4	2	2		
198532	1990261	4937	Predicted by at least		Predicted_0043	198944	199026	-1	hypothetical protein
4			one method		3	3	1		
199152	2001971	10448	Predicted by at least		Predicted_0043	199152	199179	1	hypothetical protein
3			one method		1	3	2		
199152	2001971	10448	Predicted by at least		Predicted_0043	199183	199208	1	hypothetical protein
3			one method		0	1	5		
199152	2001971	10448	Predicted by at least		Predicted_0042	199218	199264	1	hypothetical protein
3			one method		9	2	0		
199152	2001971	10448	Predicted by at least		Predicted_0042	199296	199310	1	small toxic polypeptide
3			one method		8	1	4		
199152	2001971	10448	Predicted by at least		Predicted_0042	199319	199357	-1	hypothetical protein
3			one method		7	1	1		
199152	2001971	10448	Predicted by at least	mazF_1	Predicted_0042	199360	199393	-1	mRNA interferase MazF
3			one method		6	2	4		
199152	2001971	10448	Predicted by at least	mazE	Predicted_0042	199393	199417	-1	Antitoxin MazE
3			one method		5	4	9		
199152	2001971	10448	Predicted by at least	parA_2	Predicted_0042	199448	199511	1	Chromosome partitioning

3			one method		4	9	2		protein ParA
199152	2001971	10448	Predicted by at least		Predicted_0042	199523	199546	1	ParG
3			one method		3	7	4		
199152	2001971	10448	Predicted by at least		Predicted_0434	199758	199781	1	hypothetical protein
3			one method		2	0	9		
199152	2001971	10448	Predicted by at least	xerC_6	Predicted_0434	199844	199898	-1	Tyrosine recombinase XerC
3			one method		3	8	4		
199152	2001971	10448	Predicted by at least		Predicted_0119	200098	200148	-1	hypothetical protein
3			one method		4	3	0		
199152	2001971	10448	Predicted by at least		Predicted_0119	200148	200197	-1	hypothetical protein
3			one method		3	3	1		
200288	2009541	6660	Predicted by at least		Predicted_0119	200241	200288	-1	hypothetical protein
1			one method		1	8	8		
200288	2009541	6660	Predicted by at least		Predicted_0119	200288	200325	-1	hypothetical protein
1			one method		0	1	2		
200288	2009541	6660	Predicted by at least		Predicted_0118	200326	200345	-1	hypothetical protein
1			one method		9	3	7		
200288	2009541	6660	Predicted by at least		Predicted_0118	200345	200370	-1	hypothetical protein
1			one method		8	7	8		
200288	2009541	6660	Predicted by at least		Predicted_0118	200379	200434	-1	hypothetical protein
1			one method		7	8	6		
200288	2009541	6660	Predicted by at least		Predicted_0118	200450	200485	1	hypothetical protein
1			one method		6	9	9		
200288	2009541	6660	Predicted by at least	higA_4	Predicted_0118	200486	200516	1	Antitoxin HigA
1			one method		5	4	6		
200288	2009541	6660	Predicted by at least		Predicted_0118	200519	200548	-1	hypothetical protein
1			one method		4	3	6		
200288	2009541	6660	Predicted by at least	hupB_1	Predicted_0118	200557	200584	-1	DNA-binding protein HU-beta
1			one method		3	4	6		
200288	2009541	6660	Predicted by at least	yncB	Predicted_0118	200590	200643	-1	Endonuclease precursor YncB
1			one method		2	4	1		
200288	2009541	6660	Predicted by at least		Predicted_0118	200644	200663	-1	hypothetical protein
1			one method		1	8	6		
200288	2009541	6660	Predicted by at least		Predicted_0118	200666	200751	-1	hypothetical protein
1			one method		0	2	9		
200288	2009541	6660	Predicted by at least		Predicted_0117	200750	200773	-1	hypothetical protein
1			one method		9	6	3		
200288	2009541	6660	Predicted by at least		Predicted_0117	200773	200825	-1	hypothetical protein

1			one method		8	6	4			
200288	2009541	6660	Predicted by at least		Predicted_0117	200825	200869	-1	hypothetical protein	
1			one method		7	1	7			
200288	2009541	6660	Predicted by at least		Predicted_0117	200869	200905	-1	hypothetical protein	
1			one method		6	7	6			
200288	2009541	6660	Predicted by at least		Predicted_0117	200911	200954	-1	hypothetical protein	
1			one method		5	3	1			
201045	2024913	14462	Predicted by at least	sppA_1	Predicted_0117	201045	201141	-1	Putative signal peptide	
1			one method		3	1	0		peptidase SppA	
201045	2024913	14462	Predicted by at least		Predicted_0117	201141	201226	-1	hypothetical protein	
1			one method		2	0	4			
201045	2024913	14462	Predicted by at least		Predicted_0117	201226	201256	-1	hypothetical protein	
1			one method		1	9	2			
201045	2024913	14462	Predicted by at least		Predicted_0117	201257	201304	-1	hypothetical protein	
1			one method		0	3	6			
201045	2024913	14462	Predicted by at least		Predicted_0116	201314	201366	-1	hypothetical protein	
1			one method		9	3	4			
201045	2024913	14462	Predicted by at least		Predicted_0116	201366	201418	-1	hypothetical protein	
1			one method		8	7	8			
201045	2024913	14462	Predicted by at least		Predicted_0116	201419	201480	-1	hypothetical protein	
1			one method		7	3	1			
201045	2024913	14462	Predicted by at least		Predicted_0116	201507	201618	1	hypothetical protein	
1			one method		6	0	5			
201045	2024913	14462	Predicted by at least		Predicted_0116	201620	201663	1	hypothetical protein	
1			one method		5	3	7			
201045	2024913	14462	Predicted by at least		Predicted_0116	201704	201732	-1	hypothetical protein	
1			one method		4	7	8			
201045	2024913	14462	Predicted by at least		Predicted_0116	201767	201873	-1	hypothetical protein	
1			one method		3	4	8			
201045	2024913	14462	Predicted by at least		Predicted_0116	201875	201904	-1	hypothetical protein	
1			one method		2	9	6			
201045	2024913	14462	Predicted by at least		Predicted_0116	201905	201959	-1	hypothetical protein	
1			one method		1	1	0			
201045	2024913	14462	Predicted by at least		Predicted_0116	202008	202107	1	hypothetical protein	
1			one method		0	9	2			
201045	2024913	14462	Predicted by at least		Predicted_0115	202108	202138	1	hypothetical protein	
1			one method		9	9	2			
201045	2024913	14462	Predicted by at least		Predicted_0115	202138	202180	1	H-NS histone family	

1			one method	8	4	3		
201045	2024913	14462	Predicted by at least	Predicted_0115	202186	202241	-1	transcriptional activator FlhC
1			one method	7	3	4		
201045	2024913	14462	Predicted by at least	Predicted_0115	202241	202307	-1	hypothetical protein
1			one method	6	1	6		
201045	2024913	14462	Predicted by at least	Predicted_0115	202303	202356	-1	Transglycosylase SLT domain
1			one method	5	0	3		
201045	2024913	14462	Predicted by at least	Predicted_0115	202356	202383	-1	DNA-binding transcriptional
1			one method	4	3	5		regulator Nlp
201045	2024913	14462	Predicted by at least	Predicted_0115	202455	202491	1	hypothetical protein
1			one method	3	1	3		
203001	2060599	30586	Predicted by at least	Predicted_0115	203001	203104	-1	hypothetical protein
3			one method	0	3	1		
203001	2060599	30586	Predicted by at least	Predicted_0114	203116	203267	-1	DNA helicase IV
3			one method	9	4	5		
203001	2060599	30586	Predicted by at least	Predicted_0114	203268	203296	-1	hypothetical protein
3			one method	8	4	2		
203001	2060599	30586	Predicted by at least	Predicted_0114	203296	203400	-1	Chromosome partition protein
3			one method	7	2	2		Smc
203001	2060599	30586	Predicted by at least	Predicted_0114	203415	203503	-1	DNA replication terminus site-
3			one method	6	5	0		binding protein
203001	2060599	30586	Predicted by at least	Predicted_0114	203535	203584	-1	hypothetical protein
3			one method	5	7	8		
203001	2060599	30586	Predicted by at least	Predicted_0114	203584	203671	-1	DNA polymerase III subunit
3			one method	4	5	4		epsilon
203001	2060599	30586	Predicted by at least	Predicted_0114	203671	203755	-1	Tyrosine recombinase XerD
3			one method	3	9	2		
203001	2060599	30586	Predicted by at least	Predicted_0114	203773	203826	-1	hypothetical protein
3			one method	2	2	8		
203001	2060599	30586	Predicted by at least	Predicted_0114	203826	203854	-1	hypothetical protein
3			one method	1	1	8		
203001	2060599	30586	Predicted by at least	Predicted_0114	203856	203888	-1	hypothetical protein
3			one method	0	7	4		
203001	2060599	30586	Predicted by at least	Predicted_0113	203911	203971	1	hypothetical protein
3			one method	9	0	2		
203001	2060599	30586	Predicted by at least	Predicted_0113	203972	204018	-1	hypothetical protein
3			one method	8	8	0		
203001	2060599	30586	Predicted by at least	Predicted_0113	204034	204064	-1	hypothetical protein

3			one method		7	7	6		
203001	2060599	30586	Predicted by at least one method		Predicted_0113	204094	204121	-1	hypothetical protein
3			one method		6	1	3		
203001	2060599	30586	Predicted by at least one method		Predicted_0113	204151	204244	-1	Transposase DDE domain
3			one method		5	7	9		
203001	2060599	30586	Predicted by at least one method		Predicted_0113	204253	204286	1	hypothetical protein
3			one method		4	8	4		
203001	2060599	30586	Predicted by at least one method	dinB_3	Predicted_0113	204290	204417	-1	DNA polymerase IV
3			one method		3	0	7		
203001	2060599	30586	Predicted by at least one method	lexA_3	Predicted_0113	204418	204461	-1	LexA repressor
3			one method		2	3	1		
203001	2060599	30586	Predicted by at least one method		Predicted_0113	204470	204497	-1	Helix-turn-helix domain
3			one method		1	7	9		
203001	2060599	30586	Predicted by at least one method		Predicted_0113	204510	204545	1	hypothetical protein
3			one method		0	4	4		
203001	2060599	30586	Predicted by at least one method		Predicted_0112	204547	204586	1	Transposase DDE domain
3			one method		9	5	4		
203001	2060599	30586	Predicted by at least one method		Predicted_0112	204593	204648	1	hypothetical protein
3			one method		8	2	3		
203001	2060599	30586	Predicted by at least one method		Predicted_0112	204670	204694	-1	hypothetical protein
3			one method		7	5	4		
203001	2060599	30586	Predicted by at least one method		Predicted_0112	204700	204783	-1	Integrase core domain
3			one method		6	7	7		
203001	2060599	30586	Predicted by at least one method		Predicted_0112	204783	204814	-1	Transposase
3			one method		5	4	5		
203001	2060599	30586	Predicted by at least one method	tmrB	Predicted_0112	204819	204849	-1	Tunicamycin resistance protein
3			one method		4	8	1		
203001	2060599	30586	Predicted by at least one method	yokD	Predicted_0112	204850	204936	-1	SPBc2 prophage-derived aminoglycoside N(3')-acetyltransferase-like protein YokD
3			one method		3	4	4		
203001	2060599	30586	Predicted by at least one method	bla_1	Predicted_0112	204950	205036	-1	Beta-lactamase TEM precursor
3			one method		2	6	6		
203001	2060599	30586	Predicted by at least one method	tnpR_4	Predicted_0112	205054	205110	-1	Transposon Tn3 resolvase
3			one method		1	9	6		
203001	2060599	30586	Predicted by at least one method		Predicted_0112	205127	205427	1	Tn3 transposase DDE domain
3			one method		0	0	8		
203001	2060599	30586	Predicted by at least one method		Predicted_0009	205667	205821	1	Putative transposase

3			one method		7	8	9		
203001 3	2060599	30586	Predicted by at least one method	trpF	Predicted_0009 8	205877 5	205924 5	-1	N-(5'-phosphoribosyl)anthranilate isomerase
203001 3	2060599	30586	Predicted by at least one method		Predicted_0009 9	205935 5	205990 6	1	hypothetical protein
203001 3	2060599	30586	Predicted by at least one method	blaNDM -1	Predicted_0010 0	205978 7	206059 9	-1	Beta-lactamase precursor
208286 1	2089481	6620	Predicted by at least one method		Predicted_0012 6	208286 1	208368 2	1	hypothetical protein
208286 1	2089481	6620	Predicted by at least one method		Predicted_0012 7	208414 2	208463 6	1	hypothetical protein
208286 1	2089481	6620	Predicted by at least one method	folP_1	Predicted_0012 8	208474 2	208558 1	-1	Dihydropteroate synthase
208286 1	2089481	6620	Predicted by at least one method	emrE_1	Predicted_0012 9	208557 5	208592 2	-1	Multidrug transporter EmrE
208286 1	2089481	6620	Predicted by at least one method	ant1_1	Predicted_0013 0	208608 6	208686 5	-1	Streptomycin 3"-adenylyltransferase
208286 1	2089481	6620	Predicted by at least one method	dfrA	Predicted_0013 1	208728 5	208778 2	-1	Dihydrofolate reductase
208286 1	2089481	6620	Predicted by at least one method	xerD_1	Predicted_0013 2	208792 7	208894 0	1	Tyrosine recombinase XerD
208286 1	2089481	6620	Predicted by at least one method		Predicted_0013 3	208892 7	208948 1	-1	hypothetical protein
259114 2	2599539	8397	Predicted by at least one method		Predicted_0063 7	259114 2	259415 0	-1	Tn3 transposase DDE domain
259114 2	2599539	8397	Predicted by at least one method	tnpR_3	Predicted_0063 8	259431 4	259488 6	1	Transposon Tn3 resolvase
259114 2	2599539	8397	Predicted by at least one method	fdtC	Predicted_0063 9	259500 6	259545 8	1	dTDP-3-amino-3,6-dideoxy-alpha-D-galactopyranose 3-N-acetyltransferase
259114 2	2599539	8397	Predicted by at least one method	hapE	Predicted_0064 0	259549 0	259702 2	-1	4-hydroxyacetophenone monooxygenase
259114 2	2599539	8397	Predicted by at least one method		Predicted_0064 1	259701 9	259728 8	-1	PQ loop repeat
259114 2	2599539	8397	Predicted by at least one method	ntdC	Predicted_0064 2	259729 0	259830 6	-1	Glucose-6-phosphate 3-dehydrogenase
259114	2599539	8397	Predicted by at least	ntdB	Predicted_0064	259830	259913	-1	Kanosamine-6-phosphate

2			one method		3	3	6		phosphatase
259114	2599539	8397	Predicted by at least one method	ntdA_1	Predicted_0064	259912	259953	-1	3-oxo-glucose-6-phosphate:glutamate aminotransferase
2					4	0	9		
259431	2599539	5225	Predicted by at least one method	tnpR_3	Predicted_0063	259431	259488	1	Transposon Tn3 resolvase
4					8	4	6		
259431	2599539	5225	Predicted by at least one method	fdtC	Predicted_0063	259500	259545	1	dTDP-3-amino-3,6-dideoxy-alpha-D- galactopyranose 3-N-acetyltransferase
4					9	6	8		
259431	2599539	5225	Predicted by at least one method	hapE	Predicted_0064	259549	259702	-1	4-hydroxyacetophenone monooxygenase
4					0	0	2		
259431	2599539	5225	Predicted by at least one method		Predicted_0064	259701	259728	-1	PQ loop repeat
4					1	9	8		
259431	2599539	5225	Predicted by at least one method	ntdC	Predicted_0064	259729	259830	-1	Glucose-6-phosphate 3-dehydrogenase
4					2	0	6		
259431	2599539	5225	Predicted by at least one method	ntdB	Predicted_0064	259830	259913	-1	Kanosamine-6-phosphate phosphatase
4					3	3	6		
259431	2599539	5225	Predicted by at least one method	ntdA_1	Predicted_0064	259912	259953	-1	3-oxo-glucose-6-phosphate:glutamate aminotransferase
4					4	0	9		
326772	3284238	16512	Predicted by at least one method	ppnK_2	Predicted_0230	326772	326862	1	putative inorganic polyphosphate/ATP-NAD kinase
6					2	6	5		
326772	3284238	16512	Predicted by at least one method	recN	Predicted_0230	326871	327037	1	DNA repair protein RecN
6					3	2	3		
326772	3284238	16512	Predicted by at least one method	bamE	Predicted_0230	327048	327087	1	Outer membrane protein assembly factor BamE precursor
6					4	6	8		
326772	3284238	16512	Predicted by at least one method	pasI	Predicted_0230	327105	327135	-1	Persistence and stress-resistance antitoxin PasI
6					5	2	7		
326772	3284238	16512	Predicted by at least one method	pasT	Predicted_0230	327134	327177	-1	Persistence and stress-resistance toxin PasT
6					6	1	5		
326772	3284238	16512	Predicted by at least one method	smpB	Predicted_0230	327192	327241	1	SsrA-binding protein
6					7	9	1		
326772	3284238	16512	Predicted by at least one method	intA_4	Predicted_0230	327298	327419	1	Prophage CP4-57 integrase
6					9	9	7		
326772	3284238	16512	Predicted by at least one method		Predicted_0231	327424	327629	1	hypothetical protein
6					0	3	7		

326772 6	3284238	16512	Predicted by at least one method		Predicted_0231 1	327639 9	327767 0	-1	hypothetical protein
326772 6	3284238	16512	Predicted by at least one method	era_2	Predicted_0312 9	327907 3	327994 5	1	GTPase Era
326772 6	3284238	16512	Predicted by at least one method	klcA_3	Predicted_0312 8	328003 7	328046 2	1	Antirestriction protein KlcA
326772 6	3284238	16512	Predicted by at least one method		Predicted_0312 7	328047 6	328090 7	1	hypothetical protein
326772 6	3284238	16512	Predicted by at least one method	xerD_4	Predicted_0312 6	328099 3	328200 3	-1	Tyrosine recombinase XerD
326772 6	3284238	16512	Predicted by at least one method	xerC_4	Predicted_0312 5	328199 6	328296 1	-1	Tyrosine recombinase XerC
326772 6	3284238	16512	Predicted by at least one method	xerC_3	Predicted_0312 4	328296 1	328423 8	-1	Tyrosine recombinase XerC
329683 3	3302010	5177	Predicted by at least one method		Predicted_0433 4	329683 3	329710 8	-1	InsA C-terminal domain
329683 3	3302010	5177	Predicted by at least one method		Predicted_0433 5	329722 2	329734 1	1	hypothetical protein
329683 3	3302010	5177	Predicted by at least one method		Predicted_0221 0	329913 6	329924 3	-1	hypothetical protein
329683 3	3302010	5177	Predicted by at least one method		Predicted_0220 9	329927 9	329954 5	-1	hypothetical protein
329683 3	3302010	5177	Predicted by at least one method		Predicted_0220 8	329953 5	330059 3	-1	hypothetical protein
329683 3	3302010	5177	Predicted by at least one method	hcpA_5	Predicted_0220 7	330061 1	330109 0	-1	Major exported protein
329683 3	3302010	5177	Predicted by at least one method	intA_3	Predicted_0220 6	330157 6	330201 0	1	Prophage CP4-57 integrase
362354 9	3629520	5971	Predicted by at least one method		Predicted_0086 0	362354 9	362392 0	-1	hypothetical protein
362354 9	3629520	5971	Predicted by at least one method		Predicted_0086 1	362397 9	362446 1	-1	hypothetical protein
362354 9	3629520	5971	Predicted by at least one method		Predicted_0086 2	362464 7	362504 2	-1	hypothetical protein
362354 9	3629520	5971	Predicted by at least one method		Predicted_0086 3	362514 1	362562 6	-1	hypothetical protein
362354 9	3629520	5971	Predicted by at least one method	repA	Predicted_0086 4	362614 5	362701 7	-1	Replication protein RepA

362354 9	3629520	5971	Predicted by at least one method		Predicted_0086 5	362732 1	362762 6	-1	hypothetical protein
362354 9	3629520	5971	Predicted by at least one method		Predicted_0086 6	362762 3	362794 0	-1	hypothetical protein
362354 9	3629520	5971	Predicted by at least one method		Predicted_0086 7	362806 0	362862 6	-1	hypothetical protein
362354 9	3629520	5971	Predicted by at least one method	higA_2	Predicted_0086 8	362910 4	362952 0	1	Antitoxin HigA
409709 9	4102094	4995	Predicted by at least one method		Predicted_0111 4	409709 9	409778 5	-1	hypothetical protein
409709 9	4102094	4995	Predicted by at least one method		Predicted_0111 3	409778 9	410112 1	-1	ATP-dependent helicase HepA
409709 9	4102094	4995	Predicted by at least one method		Predicted_0111 2	410115 9	410209 4	-1	hypothetical protein
446867 2	4477529	8857	Predicted by at least one method	arsD_2	Predicted_0379 6	446867 2	446903 7	-1	Arsenical resistance operon trans-acting repressor ArsD
446867 2	4477529	8857	Predicted by at least one method	arsR_3	Predicted_0379 7	446908 5	446942 0	-1	Arsenical resistance operon repressor
446867 2	4477529	8857	Predicted by at least one method		Predicted_0379 8	446990 3	447064 6	-1	hypothetical protein
446867 2	4477529	8857	Predicted by at least one method		Predicted_0379 9	447064 8	447118 4	-1	hypothetical protein
446867 2	4477529	8857	Predicted by at least one method		Predicted_0342 1	447261 3	447293 9	1	hypothetical protein
446867 2	4477529	8857	Predicted by at least one method		Predicted_0342 0	447293 6	447401 5	1	hypothetical protein
446867 2	4477529	8857	Predicted by at least one method		Predicted_0341 9	447401 9	447496 3	-1	hypothetical protein
446867 2	4477529	8857	Predicted by at least one method	thyA_2	Predicted_0341 8	447496 7	447594 4	-1	Thymidylate synthase
446867 2	4477529	8857	Predicted by at least one method	intA_5	Predicted_0341 7	447625 5	447752 9	-1	Prophage CP4-57 integrase
446908 5	4475944	6859	Predicted by at least one method	arsR_3	Predicted_0379 7	446908 5	446942 0	-1	Arsenical resistance operon repressor
446908 5	4475944	6859	Predicted by at least one method		Predicted_0379 8	446990 3	447064 6	-1	hypothetical protein
446908 5	4475944	6859	Predicted by at least one method		Predicted_0379 9	447064 8	447118 4	-1	hypothetical protein

446908 5	4475944	6859	Predicted by at least one method		Predicted_0342 1	447261 3	447293 9	1	hypothetical protein
446908 5	4475944	6859	Predicted by at least one method		Predicted_0342 0	447293 6	447401 5	1	hypothetical protein
446908 5	4475944	6859	Predicted by at least one method		Predicted_0341 9	447401 9	447496 3	-1	hypothetical protein
446908 5	4475944	6859	Predicted by at least one method	thyA_2	Predicted_0341 8	447496 7	447594 4	-1	Thymidylate synthase
480332 0	4810555	7235	Predicted by at least one method		Predicted_0245 4	480271 6	480332 7	-1	Phage tail protein (Tail_P2_I)
480332 0	4810555	7235	Predicted by at least one method		Predicted_0245 5	480332 0	480423 1	-1	Baseplate J-like protein
480332 0	4810555	7235	Predicted by at least one method		Predicted_0245 6	480423 4	480457 5	-1	Gene 25-like lysozyme
480332 0	4810555	7235	Predicted by at least one method		Predicted_0245 7	480457 2	480519 5	-1	Phage-related baseplate assembly protein
480332 0	4810555	7235	Predicted by at least one method		Predicted_0245 8	480520 2	480558 2	-1	hypothetical protein
480332 0	4810555	7235	Predicted by at least one method		Predicted_0245 9	480567 1	480628 8	-1	Phage virion morphogenesis family
480332 0	4810555	7235	Predicted by at least one method		Predicted_0246 0	480628 5	480672 8	-1	P2 phage tail completion protein R (GpR)
480332 0	4810555	7235	Predicted by at least one method		Predicted_0246 1	480669 7	480725 7	-1	hypothetical protein
480332 0	4810555	7235	Predicted by at least one method		Predicted_0246 2	480729 1	480773 4	-1	hypothetical protein
480332 0	4810555	7235	Predicted by at least one method		Predicted_0246 3	480772 7	480803 2	-1	hypothetical protein
480332 0	4810555	7235	Predicted by at least one method		Predicted_0246 4	480803 6	480823 9	-1	Phage Tail Protein X
480332 0	4810555	7235	Predicted by at least one method		Predicted_0246 5	480823 6	480871 2	-1	Phage head completion protein (GPL)
480332 0	4810555	7235	Predicted by at least one method		Predicted_0246 6	480880 1	480945 7	-1	Phage small terminase subunit
480332 0	4810555	7235	Predicted by at least one method		Predicted_0246 7	480946 1	481055 5	-1	Phage major capsid protein, P2 family
495119 7	5040496	89299	Predicted by at least one method		Predicted_0426 3	495119 7	495198 5	-1	hypothetical protein

495119 7	5040496	89299	Predicted by at least one method		Predicted_0399 4	495345 7	495369 9	1	hypothetical protein
495119 7	5040496	89299	Predicted by at least one method		Predicted_0399 5	495426 0	495525 2	-1	hypothetical protein
495119 7	5040496	89299	Predicted by at least one method		Predicted_0399 6	495538 5	495638 3	-1	Reverse transcriptase (RNA- dependent DNA polymerase)
495119 7	5040496	89299	Predicted by at least one method		Predicted_0399 7	495648 7	495745 8	-1	hypothetical protein
495119 7	5040496	89299	Predicted by at least one method		Predicted_0399 8	495752 7	495945 2	-1	chromosome segregation protein
495119 7	5040496	89299	Predicted by at least one method	mdaB_1	Predicted_0439 9	496104 8	496119 7	1	Modulator of drug activity B
495119 7	5040496	89299	Predicted by at least one method	mdaB_2	Predicted_0440 0	496119 4	496136 1	1	Modulator of drug activity B
495119 7	5040496	89299	Predicted by at least one method		Predicted_0439 1	496277 5	496323 9	-1	Replication protein
495119 7	5040496	89299	Predicted by at least one method		Predicted_0440 3	496483 9	496503 0	1	hypothetical protein
495119 7	5040496	89299	Predicted by at least one method	wbbD	Predicted_0434 0	496801 1	496884 4	-1	UDP-Gal:alpha-D-GlcNAc- diphosphoundecaprenol beta- 1,3-galactosyltransferase
495119 7	5040496	89299	Predicted by at least one method		Predicted_0434 1	496896 5	496983 1	-1	Polysaccharide pyruvyl transferase
495119 7	5040496	89299	Predicted by at least one method		Predicted_0017 6	497449 1	497489 2	1	hypothetical protein
495119 7	5040496	89299	Predicted by at least one method		Predicted_0017 7	497518 3	497582 7	1	Pentapeptide repeats (8 copies)
495119 7	5040496	89299	Predicted by at least one method		Predicted_0017 8	497646 5	497690 5	1	hypothetical protein
495119 7	5040496	89299	Predicted by at least one method		Predicted_0420 3	498032 1	498082 7	-1	hypothetical protein
495119 7	5040496	89299	Predicted by at least one method		Predicted_0420 4	498083 8	498359 7	-1	hypothetical protein
495119 7	5040496	89299	Predicted by at least one method		Predicted_0440 1	498711 3	498737 9	1	hypothetical protein
495119 7	5040496	89299	Predicted by at least one method		Predicted_0440 8	499536 3	499555 4	-1	hypothetical protein
495119	5040496	89299	Predicted by at least		Predicted_0439	499679	499754	-1	TcdA/TcdB pore forming

7			one method		2	4	6		domain
495119	5040496	89299	Predicted by at least		Predicted_0429	500199	500211	-1	hypothetical protein
7			one method		8	4	3		
495119	5040496	89299	Predicted by at least	epsE	Predicted_0439	500577	500626	1	Putative glycosyltransferase
7			one method		0	9	4		EpsE
495119	5040496	89299	Predicted by at least		Predicted_0435	501171	501180	1	hypothetical protein
7			one method		4	4	6		
495119	5040496	89299	Predicted by at least		Predicted_0426	501409	501432	1	Prophage CP4-57 regulatory
7			one method		7	2	2		protein (AlpA)
495119	5040496	89299	Predicted by at least	papC_6	Predicted_0385	501892	502129	1	Outer membrane usher
7			one method		7	7	0		protein PapC precursor
495119	5040496	89299	Predicted by at least	papD_6	Predicted_0385	502129	502202	1	Chaperone protein PapD
7			one method		8	7	2		precursor
495119	5040496	89299	Predicted by at least		Predicted_0385	502202	502300	1	hypothetical protein
7			one method		9	6	3		
495119	5040496	89299	Predicted by at least	rscB_2	Predicted_0386	502307	502372	1	Transcriptional regulatory
7			one method		0	3	6		protein RcsB
495119	5040496	89299	Predicted by at least	rscC_2	Predicted_0386	502370	502691	-1	Sensor histidine kinase RcsC
7			one method		1	4	0		
495119	5040496	89299	Predicted by at least		Predicted_0439	502834	502910	1	hypothetical protein
7			one method		4	7	8		
495119	5040496	89299	Predicted by at least	epsJ_2	Predicted_0435	503015	503100	-1	putative glycosyltransferase
7			one method		6	0	4		EpsJ
495119	5040496	89299	Predicted by at least	gtf1	Predicted_0435	503117	503160	-1	Glycosyltransferase Gtf1
7			one method		7	8	6		
495119	5040496	89299	Predicted by at least		Predicted_0431	503327	503401	-1	hypothetical protein
7			one method		9	9	0		
495119	5040496	89299	Predicted by at least		Predicted_0432	503402	503437	-1	hypothetical protein
7			one method		0	9	9		
495119	5040496	89299	Predicted by at least		Predicted_0432	503452	503468	-1	hypothetical protein
7			one method		1	8	3		
495119	5040496	89299	Predicted by at least		Predicted_0377	503970	504015	1	hypothetical protein
7			one method		9	8	4		
495119	5040496	89299	Predicted by at least	hrsA	Predicted_0378	504017	504049	1	Heat-responsive suppressor
7			one method		0	6	6		HrsA
495119	5040496	89299	Predicted by at least	fruA_2	Predicted_0378	504048	504160	1	PTS system fructose-specific
7			one method		1	3	1		EIIBC component
496119	4968844	7650	Predicted by at least	mdaB_1	Predicted_0439	496104	496119	1	Modulator of drug activity B

4			one method		9	8	7		
496119	4968844	7650	Predicted by at least one method	mdaB_2	Predicted_0440	496119	496136	1	Modulator of drug activity B
4			one method		0	4	1		
496119	4968844	7650	Predicted by at least one method		Predicted_0439	496277	496323	-1	Replication protein
4			one method		1	5	9		
496119	4968844	7650	Predicted by at least one method		Predicted_0440	496483	496503	1	hypothetical protein
4			one method		3	9	0		
496119	4968844	7650	Predicted by at least one method	wbbD	Predicted_0434	496801	496884	-1	UDP-Gal:alpha-D-GlcNAc-diphosphoundecaprenol beta-1,3-galactosyltransferase
4			one method		0	1	4		
499536	5006264	10901	Predicted by at least one method		Predicted_0440	499536	499555	-1	hypothetical protein
3			one method		8	3	4		
499536	5006264	10901	Predicted by at least one method		Predicted_0439	499679	499754	-1	TcdA/TcdB pore forming domain
3			one method		2	4	6		
499536	5006264	10901	Predicted by at least one method		Predicted_0429	500199	500211	-1	hypothetical protein
3			one method		8	4	3		
499536	5006264	10901	Predicted by at least one method	epsE	Predicted_0439	500577	500626	1	Putative glycosyltransferase EpsE
3			one method		0	9	4		
507312	5079853	6731	Predicted by at least one method	aacA4_2	Predicted_0421	507312	507372	-1	Aminoglycoside N(6')-acetyltransferase type 1
2			one method		8	2	1		
507312	5079853	6731	Predicted by at least one method		Predicted_0425	507585	507671	-1	hypothetical protein
2			one method		6	1	4		
507312	5079853	6731	Predicted by at least one method	ant1_4	Predicted_0425	507704	507798	-1	Streptomycin 3"-adenylyltransferase
2			one method		7	3	7		
507312	5079853	6731	Predicted by at least one method		Predicted_0432	507951	507985	1	hypothetical protein
2			one method		7	5	3		
507994	5085829	5883	Predicted by at least one method	smfA_7	Predicted_0432	507994	508047	1	Fimbria A protein precursor
6			one method		8	6	9		
507994	5085829	5883	Predicted by at least one method	papH_6	Predicted_0432	508052	508108	1	PAP fimbrial minor pilin protein precursor
6			one method		9	6	3		
507994	5085829	5883	Predicted by at least one method	xerD_6	Predicted_0402	508302	508362	-1	Tyrosine recombinase XerD
6			one method		1	7	9		
507994	5085829	5883	Predicted by at least one method		Predicted_0402	508386	508432	1	hypothetical protein
6			one method		2	8	9		
507994	5085829	5883	Predicted by at least one method		Predicted_0402	508533	508582	-1	hypothetical protein
6			one method		3	5	9		

Localização das Ilhas genômicas PR02

Island start	Island end	Length	Method	Gene ID	Locus	Gene start	Gene end	Strand	Product
121511	130328	8817	Predicted by at least one method		Predicted_04029	121511	121912	1	hypothetical protein
121511	130328	8817	Predicted by at least one method	mrpA_4	Predicted_04030	121905	122438	1	Major MR/P fimbria protein precursor
121511	130328	8817	Predicted by at least one method		Predicted_04330	123815	124828	1	hypothetical protein
121511	130328	8817	Predicted by at least one method		Predicted_04358	126856	127881	-1	hypothetical protein
121511	130328	8817	Predicted by at least one method		Predicted_04359	127897	128157	-1	hypothetical protein
121511	130328	8817	Predicted by at least one method	papD_11	Predicted_04180	129618	130328	-1	Chaperone protein PapD precursor
1629087	1715575	86488	Predicted by at least one method	hcpC_6	Predicted_04197	1629087	1630115	-1	Putative beta-lactamase HcpC precursor
1629087	1715575	86488	Predicted by at least one method	rhsC	Predicted_00766	1631840	1634869	-1	Putative deoxyribonuclease RhsC
1629087	1715575	86488	Predicted by at least one method	yfbR_1	Predicted_00755	1641839	1642444	-1	5'-deoxynucleotidase YfbR
1629087	1715575	86488	Predicted by at least one method	hhaIM	Predicted_00749	1647201	1648826	-1	Modification methylase HhaI
1629087	1715575	86488	Predicted by at least one method		Predicted_00742	1651213	1651602	-1	hypothetical protein
1629087	1715575	86488	Predicted by at least one method	ascD_2	Predicted_00741	1651664	1651888	-1	CDP-6-deoxy-L-threo-D-glycero-4-hexulose-3-dehydrase reductase
1629087	1715575	86488	Predicted by at least one method	ssb_2	Predicted_00732	1660219	1660746	-1	Single-stranded DNA-binding protein
1629087	1715575	86488	Predicted by at least one method	cobS_3	Predicted_00730	1661729	1662697	-1	Aerobic cobaltochelate subunit CobS
1629087	1715575	86488	Predicted by at least one method	yfgF	Predicted_00723	1669601	1670272	-1	Cyclic di-GMP phosphodiesterase YfgF
1629087	1715575	86488	Predicted by at least one method	dsbC_3	Predicted_00718	1674800	1675507	-1	Thiol:disulfide interchange protein DsbC precursor
1629087	1715575	86488	Predicted by at least one method	topB_1	Predicted_00701	1694849	1697041	1	DNA topoisomerase 3

1629087	1715575	86488	Predicted by at least one method	ydiO	Predicted_00693	1701337	1702764	-1	putative BsuMI modification methylase subunit YdiO
1629087	1715575	86488	Predicted by at least one method	korB	Predicted_00684	1706956	1708137	-1	Transcriptional repressor protein KorB
1629087	1715575	86488	Predicted by at least one method	soj	Predicted_00683	1708141	1708926	-1	Sporulation initiation inhibitor protein Soj
1629087	1715575	86488	Predicted by at least one method	bin3_2	Predicted_00680	1709731	1710468	-1	Putative transposon Tn552 DNA-invertase bin3
1629087	1715575	86488	Predicted by at least one method	dhfrI	Predicted_00200	1712632	1713105	1	Dihydrofolate reductase type 1
1629087	1715575	86488	Predicted by at least one method	ant1_2	Predicted_00202	1713782	1714570	1	Streptomycin 3"-adenylyltransferase
1690871	1705023	14152	Predicted by at least one method	topB_1	Predicted_00701	1694849	1697041	1	DNA topoisomerase 3
1690871	1705023	14152	Predicted by at least one method	ydiO	Predicted_00693	1701337	1702764	-1	putative BsuMI modification methylase subunit YdiO
1708141	1717256	9115	Predicted by at least one method	soj	Predicted_00683	1708141	1708926	-1	Sporulation initiation inhibitor protein Soj
1708141	1717256	9115	Predicted by at least one method		Predicted_00682	1709100	1709411	-1	hypothetical protein
1708141	1717256	9115	Predicted by at least one method		Predicted_00681	1709510	1709734	-1	hypothetical protein
1708141	1717256	9115	Predicted by at least one method	bin3_2	Predicted_00680	1709731	1710468	-1	Putative transposon Tn552 DNA-invertase bin3
1708141	1717256	9115	Predicted by at least one method		Predicted_00679	1710575	1711066	1	hypothetical protein
1708141	1717256	9115	Predicted by at least one method	dhfrI	Predicted_00200	1712632	1713105	1	Dihydrofolate reductase type 1
1708141	1717256	9115	Predicted by at least one method		Predicted_00201	1713200	1713724	1	TDP-fucosamine acetyltransferase
1708141	1717256	9115	Predicted by at least one method	ant1_2	Predicted_00202	1713782	1714570	1	Streptomycin 3"-adenylyltransferase
1730364	1799760	69396	Predicted by at least one method	hpaIIM	Predicted_00234	1739414	1740319	-1	Modification methylase HpaI
1730364	1799760	69396	Predicted by at least one method		Predicted_00235	1740574	1740762	-1	hypothetical protein
1730364	1799760	69396	Predicted by at least one method		Predicted_00236	1740774	1741178	-1	hypothetical protein

1730364	1799760	69396	Predicted by at least one method	nrdH_1	Predicted_00237	1741192	1741521	-1	Glutaredoxin-like protein NrdH
1730364	1799760	69396	Predicted by at least one method		Predicted_00238	1741699	1742010	-1	hypothetical protein
1730364	1799760	69396	Predicted by at least one method		Predicted_00239	1742068	1742442	-1	hypothetical protein
1730364	1799760	69396	Predicted by at least one method		Predicted_00240	1742521	1743201	-1	hypothetical protein
1730364	1799760	69396	Predicted by at least one method	dinG_1	Predicted_00241	1743242	1745302	-1	putative ATP-dependent helicase DinG
1730364	1799760	69396	Predicted by at least one method		Predicted_00242	1745409	1745717	-1	hypothetical protein
1730364	1799760	69396	Predicted by at least one method		Predicted_00243	1745735	1746013	-1	hypothetical protein
1730364	1799760	69396	Predicted by at least one method	virB	Predicted_00244	1746290	1747309	-1	Virulence regulon transcriptional activator VirB
1730364	1799760	69396	Predicted by at least one method	parA_1	Predicted_00245	1747312	1748559	-1	Plasmid partition protein A
1730364	1799760	69396	Predicted by at least one method	ascD_1	Predicted_00261	1754946	1755203	-1	CDP-6-deoxy-L-threo-D-glycero-4-hexulose-3-dehydrase reductase
1730364	1799760	69396	Predicted by at least one method	flhC_1	Predicted_00272	1762176	1762691	1	Flagellar transcriptional regulator FlhC
1730364	1799760	69396	Predicted by at least one method		Predicted_00275	1764054	1766768	-1	conjugal transfer mating pair stabilization protein TraN
1730364	1799760	69396	Predicted by at least one method		Predicted_00277	1767190	1770348	-1	TraG-like protein, N-terminal region
1730364	1799760	69396	Predicted by at least one method		Predicted_00278	1770363	1771799	-1	Conjugative relaxosome accessory transposon protein
1730364	1799760	69396	Predicted by at least one method		Predicted_00289	1776409	1778187	-1	von Willebrand factor type A domain
1730364	1799760	69396	Predicted by at least one method	cobS_1	Predicted_00292	1779938	1780999	-1	Aerobic cobaltochelatase subunit CobS
1730364	1799760	69396	Predicted by at least one method		Predicted_00293	1781066	1781440	-1	hypothetical protein
1730364	1799760	69396	Predicted by at least one method		Predicted_00294	1781808	1782923	-1	YqaJ-like viral recombinase domain
1730364	1799760	69396	Predicted by at least one method		Predicted_00295	1782983	1784527	-1	recombination and repair

			one method						protein RecT
1730364	1799760	69396	Predicted by at least one method		Predicted_00296	1784613	1785302	-1	Single-strand binding protein family
1730364	1799760	69396	Predicted by at least one method		Predicted_00297	1785770	1786207	-1	hypothetical protein
1730364	1799760	69396	Predicted by at least one method		Predicted_00298	1786286	1786894	-1	hypothetical protein
1730364	1799760	69396	Predicted by at least one method		Predicted_00299	1787404	1788153	1	hypothetical protein
1730364	1799760	69396	Predicted by at least one method		Predicted_00300	1788297	1788827	1	Sel1 repeat
1730364	1799760	69396	Predicted by at least one method		Predicted_00301	1788878	1789963	-1	TraU protein
1730364	1799760	69396	Predicted by at least one method		Predicted_00302	1789960	1791219	-1	Type-F conjugative transfer system pilin assembly protein
1730364	1799760	69396	Predicted by at least one method		Predicted_00303	1791213	1791737	-1	signal peptidase I
1730364	1799760	69396	Predicted by at least one method		Predicted_00304	1791755	1792030	-1	hypothetical protein
1730364	1799760	69396	Predicted by at least one method		Predicted_00305	1792027	1794516	-1	AAA-like domain
1730364	1799760	69396	Predicted by at least one method	dsbC_1	Predicted_00306	1794540	1795283	-1	putative thiol:disulfide interchange protein DsbC precursor
1730364	1799760	69396	Predicted by at least one method		Predicted_00307	1795691	1796224	-1	Type IV conjugative transfer system lipoprotein (TraV)
1730364	1799760	69396	Predicted by at least one method		Predicted_00308	1796227	1797615	-1	Bacterial conjugation TrbI-like protein
1730364	1799760	69396	Predicted by at least one method		Predicted_00309	1797623	1798717	-1	TraK protein
1730364	1799760	69396	Predicted by at least one method		Predicted_00310	1798886	1799476	-1	TraE protein
1730364	1799760	69396	Predicted by at least one method		Predicted_00311	1799473	1799760	-1	TraL protein
1845550	1871329	25779	Predicted by at least one method	rdgC_1	Predicted_00371	1848184	1849422	-1	Recombination-associated protein RdgC
1845550	1871329	25779	Predicted by at least one method	dam_1	Predicted_00372	1849419	1850240	-1	DNA adenine methylase

1845550	1871329	25779	Predicted by at least one method		Predicted_00373	1850295	1850936	-1	hypothetical protein
1845550	1871329	25779	Predicted by at least one method	dnaN_1	Predicted_00374	1851163	1852431	-1	DNA polymerase III subunit beta
1845550	1871329	25779	Predicted by at least one method		Predicted_00383	1856238	1856588	-1	hypothetical protein
1845550	1871329	25779	Predicted by at least one method	ssb_1	Predicted_00384	1856656	1857177	-1	Single-stranded DNA-binding protein
1845550	1871329	25779	Predicted by at least one method		Predicted_00385	1857426	1857857	-1	hypothetical protein
1845550	1871329	25779	Predicted by at least one method		Predicted_00386	1857866	1858150	-1	hypothetical protein
1845550	1871329	25779	Predicted by at least one method	hflK_1	Predicted_00387	1858261	1859181	-1	Modulator of FtsH protease HflK
1845550	1871329	25779	Predicted by at least one method		Predicted_00388	1859205	1859636	-1	NfeD-like C-terminal, partner-binding
1845550	1871329	25779	Predicted by at least one method		Predicted_00396	1865879	1867270	-1	hypothetical protein
1845550	1871329	25779	Predicted by at least one method	tnpR_1	Predicted_00397	1867347	1867928	-1	Transposon gamma-delta resolvase
1845550	1871329	25779	Predicted by at least one method		Predicted_00398	1868113	1868463	1	hypothetical protein
1845550	1871329	25779	Predicted by at least one method	higA_1	Predicted_00399	1868465	1868764	1	Antitoxin HigA
1845550	1871329	25779	Predicted by at least one method		Predicted_00400	1868905	1869921	1	hypothetical protein
1845550	1871329	25779	Predicted by at least one method		Predicted_00401	1869972	1870373	-1	hypothetical protein
1845550	1871329	25779	Predicted by at least one method		Predicted_00402	1870805	1871329	-1	LexA repressor
1880359	1892698	12339	Predicted by at least one method	comR_1	Predicted_00417	1880359	1880940	-1	HTH-type transcriptional repressor ComR
1880359	1892698	12339	Predicted by at least one method		Predicted_00418	1881089	1881661	-1	hypothetical protein
1880359	1892698	12339	Predicted by at least one method		Predicted_00419	1881797	1882489	-1	hypothetical protein
1880359	1892698	12339	Predicted by at least one method	tus_1	Predicted_00420	1882545	1883426	-1	DNA replication terminus site-binding protein

1880359	1892698	12339	Predicted by at least one method		Predicted_00180	1889232	1889960	1	Tn3 transposase DDE domain
1880359	1892698	12339	Predicted by at least one method	aacA4_1	Predicted_00527	1891353	1891958	-1	Aminoglycoside N(6')-acetyltransferase type 1
1880359	1892698	12339	Predicted by at least one method	tnpR_2	Predicted_00526	1892141	1892698	-1	Transposon Tn3 resolvase
1889232	1896303	7071	Predicted by at least one method		Predicted_00180	1889232	1889960	1	Tn3 transposase DDE domain
1889232	1896303	7071	Predicted by at least one method	aacA4_1	Predicted_00527	1891353	1891958	-1	Aminoglycoside N(6')-acetyltransferase type 1
1889232	1896303	7071	Predicted by at least one method	tnpR_2	Predicted_00526	1892141	1892698	-1	Transposon Tn3 resolvase
1889232	1896303	7071	Predicted by at least one method		Predicted_00525	1892860	1895865	1	Tn3 transposase DDE domain
1889232	1896303	7071	Predicted by at least one method		Predicted_00524	1896214	1896303	1	hypothetical protein
1889232	1896303	7071	Predicted by at least one method	dmlR_1	Predicted_00523	1896281	1897150	-1	HTH-type transcriptional regulator DmlR
1903585	1915816	12231	Predicted by at least one method		Predicted_00516	1903585	1903809	1	hypothetical protein
1903585	1915816	12231	Predicted by at least one method		Predicted_00515	1904357	1904641	1	Transposase
1903585	1915816	12231	Predicted by at least one method		Predicted_00514	1904671	1905495	1	Integrase core domain
1903585	1915816	12231	Predicted by at least one method		Predicted_00513	1905885	1906526	-1	DNA replication terminus site-binding protein
1903585	1915816	12231	Predicted by at least one method	pir	Predicted_00510	1907016	1907852	-1	PI protein
1903585	1915816	12231	Predicted by at least one method		Predicted_00509	1909188	1909679	-1	hypothetical protein
1903585	1915816	12231	Predicted by at least one method		Predicted_00508	1909690	1910070	-1	Plasmid stability protein
1903585	1915816	12231	Predicted by at least one method	parM	Predicted_00507	1910075	1911034	-1	Plasmid segregation protein ParM
1903585	1915816	12231	Predicted by at least one method		Predicted_00506	1911331	1912143	-1	hypothetical protein
1903585	1915816	12231	Predicted by at least one method		Predicted_00505	1912467	1912886	1	hypothetical protein

1903585	1915816	12231	Predicted by at least one method		Predicted_00504	1913222	1913680	-1	hypothetical protein
1903585	1915816	12231	Predicted by at least one method		Predicted_00503	1914167	1914715	1	Transglycosylase SLT domain
1903585	1915816	12231	Predicted by at least one method		Predicted_00502	1914738	1915259	1	hypothetical protein
1903585	1915816	12231	Predicted by at least one method		Predicted_00501	1915256	1915816	1	transcriptional activator FlhC
1904357	1910070	5713	Predicted by at least one method		Predicted_00515	1904357	1904641	1	Transposase
1904357	1910070	5713	Predicted by at least one method		Predicted_00514	1904671	1905495	1	Integrase core domain
1904357	1910070	5713	Predicted by at least one method		Predicted_00513	1905885	1906526	-1	DNA replication terminus site-binding protein
1904357	1910070	5713	Predicted by at least one method		Predicted_00512	1906615	1906710	-1	hypothetical protein
1904357	1910070	5713	Predicted by at least one method		Predicted_00511	1906786	1907013	-1	hypothetical protein
1904357	1910070	5713	Predicted by at least one method	pir	Predicted_00510	1907016	1907852	-1	PI protein
1904357	1910070	5713	Predicted by at least one method		Predicted_00509	1909188	1909679	-1	hypothetical protein
1904357	1910070	5713	Predicted by at least one method		Predicted_00508	1909690	1910070	-1	Plasmid stability protein
1923186	1932903	9717	Predicted by at least one method		Predicted_00496	1923186	1924595	-1	Conjugative relaxosome accessory transposon protein
1923186	1932903	9717	Predicted by at least one method		Predicted_00495	1924611	1925642	-1	hypothetical protein
1923186	1932903	9717	Predicted by at least one method		Predicted_00494	1925878	1927653	-1	von Willebrand factor type A domain
1923186	1932903	9717	Predicted by at least one method		Predicted_00493	1927718	1928905	-1	hypothetical protein
1923186	1932903	9717	Predicted by at least one method	cobS_2	Predicted_00492	1929004	1930011	-1	Aerobic cobaltochelate subunit CobS
1923186	1932903	9717	Predicted by at least one method		Predicted_00491	1930065	1930433	-1	hypothetical protein
1923186	1932903	9717	Predicted by at least one method		Predicted_00490	1930471	1931565	-1	YqaJ-like viral recombinase domain

1923186	1932903	9717	Predicted by at least one method		Predicted_00489	1931614	1932903	-1	recombination and repair protein RecT
1961683	2089481	127798	Predicted by at least one method		Predicted_00458	1961683	1964370	1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method		Predicted_00457	1964402	1966432	1	Helicase conserved C-terminal domain
1961683	2089481	127798	Predicted by at least one method		Predicted_00456	1966436	1967695	1	T5orf172 domain
1961683	2089481	127798	Predicted by at least one method	mrr_1	Predicted_00455	1967708	1968706	1	Mrr restriction system protein
1961683	2089481	127798	Predicted by at least one method		Predicted_00454	1968877	1971429	-1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method		Predicted_00453	1971578	1972045	1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method		Predicted_00452	1972042	1972491	1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method		Predicted_00451	1972530	1972961	1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method		Predicted_00450	1973020	1973229	-1	Prophage CP4-57 regulatory protein (AlpA)
1961683	2089481	127798	Predicted by at least one method	engB_1	Predicted_00449	1973307	1974296	1	GTP-binding protein EngB
1961683	2089481	127798	Predicted by at least one method		Predicted_00448	1974406	1974852	1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method		Predicted_00447	1974894	1975538	-1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method		Predicted_00446	1975985	1976917	1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method		Predicted_00445	1977142	1977471	-1	Resolvase, N terminal domain
1961683	2089481	127798	Predicted by at least one method		Predicted_00444	1977639	1980626	1	Tn3 transposase DDE domain
1961683	2089481	127798	Predicted by at least one method		Predicted_00443	1980898	1982433	1	DNA-dependent helicase II
1961683	2089481	127798	Predicted by at least one method		Predicted_00442	1982529	1983605	-1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method		Predicted_00441	1983659	1984213	-1	hypothetical protein

1961683	2089481	127798	Predicted by at least one method		Predicted_00440	1984272	1985234	-1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method		Predicted_00439	1985324	1986259	-1	Restriction endonuclease
1961683	2089481	127798	Predicted by at least one method		Predicted_00438	1986277	1986846	-1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method		Predicted_00437	1986833	1987183	-1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method	sipT	Predicted_00436	1987170	1987919	-1	Signal peptidase I T
1961683	2089481	127798	Predicted by at least one method		Predicted_00435	1988000	1988422	-1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method		Predicted_00434	1988412	1989422	-1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method		Predicted_00433	1989443	1990261	-1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method	xerC_1	Predicted_00432	1990275	1991228	-1	Tyrosine recombinase XerC
1961683	2089481	127798	Predicted by at least one method		Predicted_00431	1991523	1991792	1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method		Predicted_00430	1991831	1992085	1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method		Predicted_00429	1992182	1992640	1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method		Predicted_00428	1992961	1993104	1	small toxic polypeptide
1961683	2089481	127798	Predicted by at least one method		Predicted_00427	1993191	1993571	-1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method	mazF_1	Predicted_00426	1993602	1993934	-1	mRNA interferase MazF
1961683	2089481	127798	Predicted by at least one method	mazE	Predicted_00425	1993934	1994179	-1	Antitoxin MazE
1961683	2089481	127798	Predicted by at least one method	parA_2	Predicted_00424	1994489	1995112	1	Chromosome partitioning protein ParA
1961683	2089481	127798	Predicted by at least one method		Predicted_00423	1995237	1995464	1	ParG
1961683	2089481	127798	Predicted by at least one method		Predicted_04342	1997580	1997819	1	hypothetical protein

1961683	2089481	127798	Predicted by at least one method	xerC_6	Predicted_04343	1998448	1998984	-1	Tyrosine recombinase XerC
1961683	2089481	127798	Predicted by at least one method		Predicted_01194	2000983	2001480	-1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method		Predicted_01193	2001483	2001971	-1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method		Predicted_01192	2002068	2002403	1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method		Predicted_01191	2002418	2002888	-1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method		Predicted_01190	2002881	2003252	-1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method		Predicted_01189	2003263	2003457	-1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method		Predicted_01188	2003457	2003708	-1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method		Predicted_01187	2003798	2004346	-1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method		Predicted_01186	2004509	2004859	1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method	higA_4	Predicted_01185	2004864	2005166	1	Antitoxin HigA
1961683	2089481	127798	Predicted by at least one method		Predicted_01184	2005193	2005486	-1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method	hupB_1	Predicted_01183	2005574	2005846	-1	DNA-binding protein HU-beta
1961683	2089481	127798	Predicted by at least one method	yncB	Predicted_01182	2005904	2006431	-1	Endonuclease YncB precursor
1961683	2089481	127798	Predicted by at least one method		Predicted_01181	2006448	2006636	-1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method		Predicted_01180	2006662	2007519	-1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method		Predicted_01179	2007506	2007733	-1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method		Predicted_01178	2007736	2008254	-1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method		Predicted_01177	2008251	2008697	-1	hypothetical protein

1961683	2089481	127798	Predicted by at least one method		Predicted_01176	2008697	2009056	-1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method		Predicted_01175	2009113	2009541	-1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method	bdbD_2	Predicted_01174	2009575	2010435	-1	Disulfide bond formation protein D precursor
1961683	2089481	127798	Predicted by at least one method	sppA_1	Predicted_01173	2010451	2011410	-1	Putative signal peptide peptidase SppA
1961683	2089481	127798	Predicted by at least one method		Predicted_01159	2021089	2021382	1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method		Predicted_01158	2021384	2021803	1	H-NS histone family
1961683	2089481	127798	Predicted by at least one method		Predicted_01157	2021863	2022414	-1	transcriptional activator FlhC
1961683	2089481	127798	Predicted by at least one method		Predicted_01156	2022411	2023076	-1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method		Predicted_01155	2023030	2023563	-1	Transglycosylase SLT domain
1961683	2089481	127798	Predicted by at least one method		Predicted_01154	2023563	2023835	-1	DNA-binding transcriptional regulator Nlp
1961683	2089481	127798	Predicted by at least one method		Predicted_01153	2024551	2024913	1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method		Predicted_01152	2024951	2028565	-1	TraG-like protein, N-terminal region
1961683	2089481	127798	Predicted by at least one method		Predicted_01151	2028578	2030011	-1	Conjugative relaxosome accessory transposon protein
1961683	2089481	127798	Predicted by at least one method		Predicted_01150	2030013	2031041	-1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method		Predicted_01149	2031164	2032675	-1	DNA helicase IV
1961683	2089481	127798	Predicted by at least one method		Predicted_01148	2032684	2032962	-1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method	smc_2	Predicted_01147	2032962	2034002	-1	Chromosome partition protein Smc
1961683	2089481	127798	Predicted by at least one method	tus_3	Predicted_01146	2034155	2035030	-1	DNA replication terminus site-binding protein
1961683	2089481	127798	Predicted by at least one method		Predicted_01145	2035357	2035848	-1	hypothetical protein

1961683	2089481	127798	Predicted by at least one method	dnaQ_1	Predicted_01144	2035845	2036714	-1	DNA polymerase III subunit epsilon
1961683	2089481	127798	Predicted by at least one method	xerD_2	Predicted_01143	2036719	2037552	-1	Tyrosine recombinase XerD
1961683	2089481	127798	Predicted by at least one method		Predicted_01135	2041517	2042449	-1	Transposase DDE domain
1961683	2089481	127798	Predicted by at least one method		Predicted_01134	2042538	2042864	1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method	dinB_3	Predicted_01133	2042900	2044177	-1	DNA polymerase IV
1961683	2089481	127798	Predicted by at least one method	lexA_3	Predicted_01132	2044183	2044611	-1	LexA repressor
1961683	2089481	127798	Predicted by at least one method		Predicted_01131	2044707	2044979	-1	Helix-turn-helix domain
1961683	2089481	127798	Predicted by at least one method		Predicted_01130	2045104	2045454	1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method		Predicted_01129	2045475	2045864	1	Transposase DDE domain
1961683	2089481	127798	Predicted by at least one method		Predicted_01128	2045932	2046483	1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method		Predicted_01127	2046705	2046944	-1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method		Predicted_01126	2047007	2047837	-1	Integrase core domain
1961683	2089481	127798	Predicted by at least one method		Predicted_01125	2047834	2048145	-1	Transposase
1961683	2089481	127798	Predicted by at least one method	tmrB	Predicted_01124	2048198	2048491	-1	Tunicamycin resistance protein
1961683	2089481	127798	Predicted by at least one method	yokD	Predicted_01123	2048504	2049364	-1	SPBc2 prophage-derived aminoglycoside N(3')-acetyltransferase-like protein YokD
1961683	2089481	127798	Predicted by at least one method	bla_1	Predicted_01122	2049506	2050366	-1	Beta-lactamase TEM precursor
1961683	2089481	127798	Predicted by at least one method	tnpR_4	Predicted_01121	2050549	2051106	-1	Transposon Tn3 resolvase
1961683	2089481	127798	Predicted by at least one method		Predicted_01120	2051270	2054278	1	Tn3 transposase DDE domain

1961683	2089481	127798	Predicted by at least one method		Predicted_00097	2056678	2058219	1	Putative transposase
1961683	2089481	127798	Predicted by at least one method	trpF	Predicted_00098	2058775	2059245	-1	N-(5'-phosphoribosyl)anthranilate isomerase
1961683	2089481	127798	Predicted by at least one method		Predicted_00099	2059355	2059906	1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method	blaNDM-1	Predicted_00100	2059787	2060599	-1	Beta-lactamase NDM-1 precursor
1961683	2089481	127798	Predicted by at least one method		Predicted_00101	2060700	2061710	-1	Integrase core domain
1961683	2089481	127798	Predicted by at least one method	aphA_1	Predicted_00102	2061877	2062656	-1	Aminoglycoside 3'-phosphotransferase
1961683	2089481	127798	Predicted by at least one method		Predicted_00103	2062762	2062992	-1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method		Predicted_00104	2062982	2063425	-1	Transposase
1961683	2089481	127798	Predicted by at least one method		Predicted_00105	2063716	2065977	-1	Tn3 transposase DDE domain
1961683	2089481	127798	Predicted by at least one method	hin	Predicted_00106	2065980	2066612	-1	DNA-invertase hin
1961683	2089481	127798	Predicted by at least one method	ttgC	Predicted_00107	2066668	2068155	-1	Toluene efflux pump outer membrane protein TtgC precursor
1961683	2089481	127798	Predicted by at least one method		Predicted_00108	2068142	2068615	-1	Cytochrome C'
1961683	2089481	127798	Predicted by at least one method	lolD_1	Predicted_00109	2068644	2069354	-1	Lipoprotein-releasing system ATP-binding protein LolD
1961683	2089481	127798	Predicted by at least one method		Predicted_00110	2069359	2070561	-1	FtsX-like permease family
1961683	2089481	127798	Predicted by at least one method	mdtA_1	Predicted_00111	2070558	2071709	-1	Multidrug resistance protein MdtA precursor
1961683	2089481	127798	Predicted by at least one method	slmA_1	Predicted_00112	2071706	2072314	-1	Nucleoid occlusion factor SlmA
1961683	2089481	127798	Predicted by at least one method		Predicted_00113	2072395	2073069	-1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method		Predicted_00114	2073624	2073971	1	hypothetical protein

1961683	2089481	127798	Predicted by at least one method		Predicted_00115	2074016	2077042	-1	Tn3 transposase DDE domain
1961683	2089481	127798	Predicted by at least one method	bin3_1	Predicted_00116	2077204	2077848	1	Putative transposon Tn552 DNA-invertase bin3
1961683	2089481	127798	Predicted by at least one method	betT_1	Predicted_00117	2078032	2078361	1	High-affinity choline transport protein
1961683	2089481	127798	Predicted by at least one method	merR	Predicted_00118	2078351	2078806	-1	Mercuric resistance operon regulatory protein
1961683	2089481	127798	Predicted by at least one method		Predicted_00119	2078878	2079243	1	MerT mercuric transport protein
1961683	2089481	127798	Predicted by at least one method	merP	Predicted_00120	2079259	2079534	1	Mercuric transport protein periplasmic component precursor
1961683	2089481	127798	Predicted by at least one method	merC	Predicted_00121	2079562	2079987	1	Mercuric resistance protein MerC
1961683	2089481	127798	Predicted by at least one method	merA	Predicted_00122	2080026	2081711	1	Mercuric reductase
1961683	2089481	127798	Predicted by at least one method		Predicted_00123	2081729	2082094	1	zinc-responsive transcriptional regulator
1961683	2089481	127798	Predicted by at least one method		Predicted_00124	2082091	2082327	1	MerE protein
1961683	2089481	127798	Predicted by at least one method		Predicted_00125	2082393	2082605	1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method		Predicted_00126	2082861	2083682	1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method		Predicted_00127	2084142	2084636	1	hypothetical protein
1961683	2089481	127798	Predicted by at least one method	folP_1	Predicted_00128	2084742	2085581	-1	Dihydropteroate synthase
1961683	2089481	127798	Predicted by at least one method	emrE_1	Predicted_00129	2085575	2085922	-1	Multidrug transporter EmrE
1961683	2089481	127798	Predicted by at least one method	ant1_1	Predicted_00130	2086086	2086865	-1	Streptomycin 3"-adenylyltransferase
1961683	2089481	127798	Predicted by at least one method	dfrA	Predicted_00131	2087285	2087782	-1	Dihydrofolate reductase
1961683	2089481	127798	Predicted by at least one method	xerD_1	Predicted_00132	2087927	2088940	1	Tyrosine recombinase XerD
1961683	2089481	127798	Predicted by at least one method		Predicted_00133	2088927	2089481	-1	hypothetical protein

			one method						
1973020	1980626	7606	Predicted by at least one method		Predicted_00450	1973020	1973229	-1	Prophage CP4-57 regulatory protein (AlpA)
1973020	1980626	7606	Predicted by at least one method	engB_1	Predicted_00449	1973307	1974296	1	GTP-binding protein EngB
1973020	1980626	7606	Predicted by at least one method		Predicted_00448	1974406	1974852	1	hypothetical protein
1973020	1980626	7606	Predicted by at least one method		Predicted_00447	1974894	1975538	-1	hypothetical protein
1973020	1980626	7606	Predicted by at least one method		Predicted_00446	1975985	1976917	1	hypothetical protein
1973020	1980626	7606	Predicted by at least one method		Predicted_00445	1977142	1977471	-1	Resolvase, N terminal domain
1973020	1980626	7606	Predicted by at least one method		Predicted_00444	1977639	1980626	1	Tn3 transposase DDE domain
1985324	1990261	4937	Predicted by at least one method		Predicted_00439	1985324	1986259	-1	Restriction endonuclease
1985324	1990261	4937	Predicted by at least one method		Predicted_00438	1986277	1986846	-1	hypothetical protein
1985324	1990261	4937	Predicted by at least one method		Predicted_00437	1986833	1987183	-1	hypothetical protein
1985324	1990261	4937	Predicted by at least one method	sipT	Predicted_00436	1987170	1987919	-1	Signal peptidase I T
1985324	1990261	4937	Predicted by at least one method		Predicted_00435	1988000	1988422	-1	hypothetical protein
1985324	1990261	4937	Predicted by at least one method		Predicted_00434	1988412	1989422	-1	hypothetical protein
1985324	1990261	4937	Predicted by at least one method		Predicted_00433	1989443	1990261	-1	hypothetical protein
1991523	2001971	10448	Predicted by at least one method		Predicted_00431	1991523	1991792	1	hypothetical protein
1991523	2001971	10448	Predicted by at least one method		Predicted_00430	1991831	1992085	1	hypothetical protein
1991523	2001971	10448	Predicted by at least one method		Predicted_00429	1992182	1992640	1	hypothetical protein
1991523	2001971	10448	Predicted by at least one method		Predicted_00428	1992961	1993104	1	small toxic polypeptide
1991523	2001971	10448	Predicted by at least		Predicted_00427	1993191	1993571	-1	hypothetical protein

			one method						
1991523	2001971	10448	Predicted by at least one method	mazF_1	Predicted_00426	1993602	1993934	-1	mRNA interferase MazF
1991523	2001971	10448	Predicted by at least one method	mazE	Predicted_00425	1993934	1994179	-1	Antitoxin MazE
1991523	2001971	10448	Predicted by at least one method	parA_2	Predicted_00424	1994489	1995112	1	Chromosome partitioning protein ParA
1991523	2001971	10448	Predicted by at least one method		Predicted_00423	1995237	1995464	1	ParG
1991523	2001971	10448	Predicted by at least one method		Predicted_04342	1997580	1997819	1	hypothetical protein
1991523	2001971	10448	Predicted by at least one method	xerC_6	Predicted_04343	1998448	1998984	-1	Tyrosine recombinase XerC
1991523	2001971	10448	Predicted by at least one method		Predicted_01194	2000983	2001480	-1	hypothetical protein
1991523	2001971	10448	Predicted by at least one method		Predicted_01193	2001483	2001971	-1	hypothetical protein
2002881	2009541	6660	Predicted by at least one method		Predicted_01191	2002418	2002888	-1	hypothetical protein
2002881	2009541	6660	Predicted by at least one method		Predicted_01190	2002881	2003252	-1	hypothetical protein
2002881	2009541	6660	Predicted by at least one method		Predicted_01189	2003263	2003457	-1	hypothetical protein
2002881	2009541	6660	Predicted by at least one method		Predicted_01188	2003457	2003708	-1	hypothetical protein
2002881	2009541	6660	Predicted by at least one method		Predicted_01187	2003798	2004346	-1	hypothetical protein
2002881	2009541	6660	Predicted by at least one method		Predicted_01186	2004509	2004859	1	hypothetical protein
2002881	2009541	6660	Predicted by at least one method	higA_4	Predicted_01185	2004864	2005166	1	Antitoxin HigA
2002881	2009541	6660	Predicted by at least one method		Predicted_01184	2005193	2005486	-1	hypothetical protein
2002881	2009541	6660	Predicted by at least one method	hupB_1	Predicted_01183	2005574	2005846	-1	DNA-binding protein HU-beta
2002881	2009541	6660	Predicted by at least one method	yncB	Predicted_01182	2005904	2006431	-1	Endonuclease YncB precursor
2002881	2009541	6660	Predicted by at least one method		Predicted_01181	2006448	2006636	-1	hypothetical protein

			one method						
2002881	2009541	6660	Predicted by at least one method		Predicted_01180	2006662	2007519	-1	hypothetical protein
2002881	2009541	6660	Predicted by at least one method		Predicted_01179	2007506	2007733	-1	hypothetical protein
2002881	2009541	6660	Predicted by at least one method		Predicted_01178	2007736	2008254	-1	hypothetical protein
2002881	2009541	6660	Predicted by at least one method		Predicted_01177	2008251	2008697	-1	hypothetical protein
2002881	2009541	6660	Predicted by at least one method		Predicted_01176	2008697	2009056	-1	hypothetical protein
2002881	2009541	6660	Predicted by at least one method		Predicted_01175	2009113	2009541	-1	hypothetical protein
2010451	2024913	14462	Predicted by at least one method	sppA_1	Predicted_01173	2010451	2011410	-1	Putative signal peptide
2010451	2024913	14462	Predicted by at least one method		Predicted_01172	2011410	2012264	-1	peptidase SppA
2010451	2024913	14462	Predicted by at least one method		Predicted_01171	2012269	2012562	-1	hypothetical protein
2010451	2024913	14462	Predicted by at least one method		Predicted_01170	2012573	2013046	-1	hypothetical protein
2010451	2024913	14462	Predicted by at least one method		Predicted_01169	2013143	2013664	-1	hypothetical protein
2010451	2024913	14462	Predicted by at least one method		Predicted_01168	2013667	2014188	-1	hypothetical protein
2010451	2024913	14462	Predicted by at least one method		Predicted_01167	2014193	2014801	-1	hypothetical protein
2010451	2024913	14462	Predicted by at least one method		Predicted_01166	2015070	2016185	1	hypothetical protein
2010451	2024913	14462	Predicted by at least one method		Predicted_01165	2016203	2016637	1	hypothetical protein
2010451	2024913	14462	Predicted by at least one method		Predicted_01164	2017047	2017328	-1	hypothetical protein
2010451	2024913	14462	Predicted by at least one method		Predicted_01163	2017674	2018738	-1	hypothetical protein
2010451	2024913	14462	Predicted by at least one method		Predicted_01162	2018759	2019046	-1	hypothetical protein
2010451	2024913	14462	Predicted by at least one method		Predicted_01161	2019051	2019590	-1	hypothetical protein

			one method						
2010451	2024913	14462	Predicted by at least one method		Predicted_01160	2020089	2021072	1	hypothetical protein
2010451	2024913	14462	Predicted by at least one method		Predicted_01159	2021089	2021382	1	hypothetical protein
2010451	2024913	14462	Predicted by at least one method		Predicted_01158	2021384	2021803	1	H-NS histone family
2010451	2024913	14462	Predicted by at least one method		Predicted_01157	2021863	2022414	-1	transcriptional activator FlhC
2010451	2024913	14462	Predicted by at least one method		Predicted_01156	2022411	2023076	-1	hypothetical protein
2010451	2024913	14462	Predicted by at least one method		Predicted_01155	2023030	2023563	-1	Transglycosylase SLT domain
2010451	2024913	14462	Predicted by at least one method		Predicted_01154	2023563	2023835	-1	DNA-binding transcriptional regulator Nlp
2010451	2024913	14462	Predicted by at least one method		Predicted_01153	2024551	2024913	1	hypothetical protein
2030013	2060599	30586	Predicted by at least one method		Predicted_01150	2030013	2031041	-1	hypothetical protein
2030013	2060599	30586	Predicted by at least one method		Predicted_01149	2031164	2032675	-1	DNA helicase IV
2030013	2060599	30586	Predicted by at least one method		Predicted_01148	2032684	2032962	-1	hypothetical protein
2030013	2060599	30586	Predicted by at least one method	smc_2	Predicted_01147	2032962	2034002	-1	Chromosome partition protein Smc
2030013	2060599	30586	Predicted by at least one method	tus_3	Predicted_01146	2034155	2035030	-1	DNA replication terminus site-binding protein
2030013	2060599	30586	Predicted by at least one method		Predicted_01145	2035357	2035848	-1	hypothetical protein
2030013	2060599	30586	Predicted by at least one method	dnaQ_1	Predicted_01144	2035845	2036714	-1	DNA polymerase III subunit epsilon
2030013	2060599	30586	Predicted by at least one method	xerD_2	Predicted_01143	2036719	2037552	-1	Tyrosine recombinase XerD
2030013	2060599	30586	Predicted by at least one method		Predicted_01135	2041517	2042449	-1	Transposase DDE domain
2030013	2060599	30586	Predicted by at least one method		Predicted_01134	2042538	2042864	1	hypothetical protein
2030013	2060599	30586	Predicted by at least one method	dinB_3	Predicted_01133	2042900	2044177	-1	DNA polymerase IV

			one method						
2030013	2060599	30586	Predicted by at least one method	lexA_3	Predicted_01132	2044183	2044611	-1	LexA repressor
2030013	2060599	30586	Predicted by at least one method		Predicted_01131	2044707	2044979	-1	Helix-turn-helix domain
2030013	2060599	30586	Predicted by at least one method		Predicted_01130	2045104	2045454	1	hypothetical protein
2030013	2060599	30586	Predicted by at least one method		Predicted_01129	2045475	2045864	1	Transposase DDE domain
2030013	2060599	30586	Predicted by at least one method		Predicted_01128	2045932	2046483	1	hypothetical protein
2030013	2060599	30586	Predicted by at least one method		Predicted_01127	2046705	2046944	-1	hypothetical protein
2030013	2060599	30586	Predicted by at least one method		Predicted_01126	2047007	2047837	-1	Integrase core domain
2030013	2060599	30586	Predicted by at least one method		Predicted_01125	2047834	2048145	-1	Transposase
2030013	2060599	30586	Predicted by at least one method	tmrB	Predicted_01124	2048198	2048491	-1	Tunicamycin resistance protein
2030013	2060599	30586	Predicted by at least one method	yokD	Predicted_01123	2048504	2049364	-1	SPBc2 prophage-derived aminoglycoside N(3')-acetyltransferase-like protein YokD
2030013	2060599	30586	Predicted by at least one method	bla_1	Predicted_01122	2049506	2050366	-1	Beta-lactamase TEM precursor
2030013	2060599	30586	Predicted by at least one method	tnpR_4	Predicted_01121	2050549	2051106	-1	Transposon Tn3 resolvase
2030013	2060599	30586	Predicted by at least one method		Predicted_01120	2051270	2054278	1	Tn3 transposase DDE domain
2030013	2060599	30586	Predicted by at least one method		Predicted_00097	2056678	2058219	1	Putative transposase
2030013	2060599	30586	Predicted by at least one method	trpF	Predicted_00098	2058775	2059245	-1	N-(5'-phosphoribosyl)anthranilate isomerase
2030013	2060599	30586	Predicted by at least one method		Predicted_00099	2059355	2059906	1	hypothetical protein
2030013	2060599	30586	Predicted by at least one method	blaNDM-1	Predicted_00100	2059787	2060599	-1	Beta-lactamase NDM-1 precursor

2082861	2089481	6620	Predicted by at least one method		Predicted_00126	2082861	2083682	1	hypothetical protein
2082861	2089481	6620	Predicted by at least one method		Predicted_00127	2084142	2084636	1	hypothetical protein
2082861	2089481	6620	Predicted by at least one method	folP_1	Predicted_00128	2084742	2085581	-1	Dihydropteroate synthase
2082861	2089481	6620	Predicted by at least one method	emrE_1	Predicted_00129	2085575	2085922	-1	Multidrug transporter EmrE
2082861	2089481	6620	Predicted by at least one method	ant1_1	Predicted_00130	2086086	2086865	-1	Streptomycin 3"-adenylyltransferase
2082861	2089481	6620	Predicted by at least one method	dfrA	Predicted_00131	2087285	2087782	-1	Dihydrofolate reductase
2082861	2089481	6620	Predicted by at least one method	xerD_1	Predicted_00132	2087927	2088940	1	Tyrosine recombinase XerD
2082861	2089481	6620	Predicted by at least one method		Predicted_00133	2088927	2089481	-1	hypothetical protein
2591142	2599539	8397	Predicted by at least one method		Predicted_00637	2591142	2594150	-1	Tn3 transposase DDE domain
2591142	2599539	8397	Predicted by at least one method	tnpR_3	Predicted_00638	2594314	2594886	1	Transposon Tn3 resolvase
2591142	2599539	8397	Predicted by at least one method	fdtC	Predicted_00639	2595006	2595458	1	dTDP-3-amino-3,6-dideoxy-alpha-D- galactopyranose 3-N-acetyltransferase
2591142	2599539	8397	Predicted by at least one method	hapE	Predicted_00640	2595490	2597022	-1	4-hydroxyacetophenone monooxygenase
2591142	2599539	8397	Predicted by at least one method		Predicted_00641	2597019	2597288	-1	PQ loop repeat
2591142	2599539	8397	Predicted by at least one method	ntdC	Predicted_00642	2597290	2598306	-1	Glucose-6-phosphate 3-dehydrogenase
2591142	2599539	8397	Predicted by at least one method	ntdB	Predicted_00643	2598303	2599136	-1	Kanosamine-6-phosphate phosphatase
2591142	2599539	8397	Predicted by at least one method	ntdA_1	Predicted_00644	2599120	2599539	-1	3-oxo-glucose-6-phosphate:glutamate aminotransferase
2594314	2599539	5225	Predicted by at least one method	tnpR_3	Predicted_00638	2594314	2594886	1	Transposon Tn3 resolvase
2594314	2599539	5225	Predicted by at least one method	fdtC	Predicted_00639	2595006	2595458	1	dTDP-3-amino-3,6-dideoxy-alpha-D- galactopyranose 3-N-acetyltransferase

2594314	2599539	5225	Predicted by at least one method	hapE	Predicted_00640	2595490	2597022	-1	4-hydroxyacetophenone monooxygenase
2594314	2599539	5225	Predicted by at least one method		Predicted_00641	2597019	2597288	-1	PQ loop repeat
2594314	2599539	5225	Predicted by at least one method	ntdC	Predicted_00642	2597290	2598306	-1	Glucose-6-phosphate 3-dehydrogenase
2594314	2599539	5225	Predicted by at least one method	ntdB	Predicted_00643	2598303	2599136	-1	Kanosamine-6-phosphate phosphatase
2594314	2599539	5225	Predicted by at least one method	ntdA_1	Predicted_00644	2599120	2599539	-1	3-oxo-glucose-6-phosphate:glutamate aminotransferase
3267726	3284238	16512	Predicted by at least one method	ppnK_2	Predicted_02302	3267726	3268625	1	putative inorganic polyphosphate/ATP-NAD kinase
3267726	3284238	16512	Predicted by at least one method	recN	Predicted_02303	3268712	3270373	1	DNA repair protein RecN
3267726	3284238	16512	Predicted by at least one method	bamE	Predicted_02304	3270486	3270878	1	Outer membrane protein assembly factor BamE precursor
3267726	3284238	16512	Predicted by at least one method	pasI	Predicted_02305	3271052	3271357	-1	Persistence and stress-resistance antitoxin PasI
3267726	3284238	16512	Predicted by at least one method	pasT	Predicted_02306	3271341	3271775	-1	Prophage CP4-57 integrase
3267726	3284238	16512	Predicted by at least one method	smpB	Predicted_02307	3271929	3272411	1	SsrA-binding protein
3267726	3284238	16512	Predicted by at least one method	intA_4	Predicted_02309	3272989	3274197	1	Prophage CP4-57 integrase
3267726	3284238	16512	Predicted by at least one method		Predicted_02310	3274243	3276297	1	hypothetical protein
3267726	3284238	16512	Predicted by at least one method		Predicted_02311	3276399	3277670	-1	hypothetical protein
3267726	3284238	16512	Predicted by at least one method	era_2	Predicted_03129	3279073	3279945	1	GTPase Era
3267726	3284238	16512	Predicted by at least one method	kIcA_3	Predicted_03128	3280037	3280462	1	Antirestriction protein KlcA
3267726	3284238	16512	Predicted by at least one method		Predicted_03127	3280476	3280907	1	hypothetical protein
3267726	3284238	16512	Predicted by at least one method	xerD_4	Predicted_03126	3280993	3282003	-1	Tyrosine recombinase XerD

			one method						
3267726	3284238	16512	Predicted by at least one method	xerC_4	Predicted_03125	3281996	3282961	-1	Tyrosine recombinase XerC
3267726	3284238	16512	Predicted by at least one method	xerC_3	Predicted_03124	3282961	3284238	-1	Tyrosine recombinase XerC
3296833	3302010	5177	Predicted by at least one method		Predicted_04334	3296833	3297108	-1	InsA C-terminal domain
3296833	3302010	5177	Predicted by at least one method		Predicted_04335	3297222	3297341	1	hypothetical protein
3296833	3302010	5177	Predicted by at least one method		Predicted_02210	3299136	3299243	-1	hypothetical protein
3296833	3302010	5177	Predicted by at least one method		Predicted_02209	3299279	3299545	-1	hypothetical protein
3296833	3302010	5177	Predicted by at least one method		Predicted_02208	3299535	3300593	-1	hypothetical protein
3296833	3302010	5177	Predicted by at least one method	hcpA_5	Predicted_02207	3300611	3301090	-1	Major exported protein
3296833	3302010	5177	Predicted by at least one method	intA_3	Predicted_02206	3301576	3302010	1	Prophage CP4-57 integrase
3623549	3629520	5971	Predicted by at least one method		Predicted_00860	3623549	3623920	-1	hypothetical protein
3623549	3629520	5971	Predicted by at least one method		Predicted_00861	3623979	3624461	-1	hypothetical protein
3623549	3629520	5971	Predicted by at least one method		Predicted_00862	3624647	3625042	-1	hypothetical protein
3623549	3629520	5971	Predicted by at least one method		Predicted_00863	3625141	3625626	-1	hypothetical protein
3623549	3629520	5971	Predicted by at least one method	repA	Predicted_00864	3626145	3627017	-1	Replication protein RepA
3623549	3629520	5971	Predicted by at least one method		Predicted_00865	3627321	3627626	-1	hypothetical protein
3623549	3629520	5971	Predicted by at least one method		Predicted_00866	3627623	3627940	-1	hypothetical protein
3623549	3629520	5971	Predicted by at least one method		Predicted_00867	3628060	3628626	-1	hypothetical protein
3623549	3629520	5971	Predicted by at least one method	higA_2	Predicted_00868	3629104	3629520	1	Antitoxin HigA
4097099	4102094	4995	Predicted by at least		Predicted_01114	4097099	4097785	-1	hypothetical protein

			one method						
4097099	4102094	4995	Predicted by at least one method		Predicted_01113	4097789	4101121	-1	ATP-dependent helicase HepA
4097099	4102094	4995	Predicted by at least one method		Predicted_01112	4101159	4102094	-1	hypothetical protein
4468672	4477529	8857	Predicted by at least one method	arsD_2	Predicted_03796	4468672	4469037	-1	Arsenical resistance operon trans-acting repressor ArsD
4468672	4477529	8857	Predicted by at least one method	arsR_3	Predicted_03797	4469085	4469420	-1	Arsenical resistance operon repressor
4468672	4477529	8857	Predicted by at least one method		Predicted_03798	4469903	4470646	-1	hypothetical protein
4468672	4477529	8857	Predicted by at least one method		Predicted_03799	4470648	4471184	-1	hypothetical protein
4468672	4477529	8857	Predicted by at least one method		Predicted_03421	4472613	4472939	1	hypothetical protein
4468672	4477529	8857	Predicted by at least one method		Predicted_03420	4472936	4474015	1	hypothetical protein
4468672	4477529	8857	Predicted by at least one method		Predicted_03419	4474019	4474963	-1	hypothetical protein
4468672	4477529	8857	Predicted by at least one method	thyA_2	Predicted_03418	4474967	4475944	-1	Thymidylate synthase
4468672	4477529	8857	Predicted by at least one method	intA_5	Predicted_03417	4476255	4477529	-1	Prophage CP4-57 integrase
4469085	4475944	6859	Predicted by at least one method	arsR_3	Predicted_03797	4469085	4469420	-1	Arsenical resistance operon repressor
4469085	4475944	6859	Predicted by at least one method		Predicted_03798	4469903	4470646	-1	hypothetical protein
4469085	4475944	6859	Predicted by at least one method		Predicted_03799	4470648	4471184	-1	hypothetical protein
4469085	4475944	6859	Predicted by at least one method		Predicted_03421	4472613	4472939	1	hypothetical protein
4469085	4475944	6859	Predicted by at least one method		Predicted_03420	4472936	4474015	1	hypothetical protein
4469085	4475944	6859	Predicted by at least one method		Predicted_03419	4474019	4474963	-1	hypothetical protein
4469085	4475944	6859	Predicted by at least one method	thyA_2	Predicted_03418	4474967	4475944	-1	Thymidylate synthase
4803320	4810555	7235	Predicted by at least		Predicted_02454	4802716	4803327	-1	Phage tail protein (Tail_P2_I)

			one method						
4803320	4810555	7235	Predicted by at least one method	Predicted_02455	4803320	4804231	-1	Baseplate J-like protein	
4803320	4810555	7235	Predicted by at least one method	Predicted_02456	4804234	4804575	-1	Gene 25-like lysozyme	
4803320	4810555	7235	Predicted by at least one method	Predicted_02457	4804572	4805195	-1	Phage-related baseplate assembly protein	
4803320	4810555	7235	Predicted by at least one method	Predicted_02458	4805202	4805582	-1	hypothetical protein	
4803320	4810555	7235	Predicted by at least one method	Predicted_02459	4805671	4806288	-1	Phage virion morphogenesis family	
4803320	4810555	7235	Predicted by at least one method	Predicted_02460	4806285	4806728	-1	P2 phage tail completion protein R (GpR)	
4803320	4810555	7235	Predicted by at least one method	Predicted_02461	4806697	4807257	-1	hypothetical protein	
4803320	4810555	7235	Predicted by at least one method	Predicted_02462	4807291	4807734	-1	hypothetical protein	
4803320	4810555	7235	Predicted by at least one method	Predicted_02463	4807727	4808032	-1	hypothetical protein	
4803320	4810555	7235	Predicted by at least one method	Predicted_02464	4808036	4808239	-1	Phage Tail Protein X	
4803320	4810555	7235	Predicted by at least one method	Predicted_02465	4808236	4808712	-1	Phage head completion protein (GPL)	
4803320	4810555	7235	Predicted by at least one method	Predicted_02466	4808801	4809457	-1	Phage small terminase subunit	
4803320	4810555	7235	Predicted by at least one method	Predicted_02467	4809461	4810555	-1	Phage major capsid protein, P2 family	
4951197	5040496	89299	Predicted by at least one method	Predicted_04263	4951197	4951985	-1	hypothetical protein	
4951197	5040496	89299	Predicted by at least one method	Predicted_03994	4953457	4953699	1	hypothetical protein	
4951197	5040496	89299	Predicted by at least one method	Predicted_03995	4954260	4955252	-1	hypothetical protein	
4951197	5040496	89299	Predicted by at least one method	Predicted_03996	4955385	4956383	-1	Reverse transcriptase (RNA-dependent DNA polymerase)	
4951197	5040496	89299	Predicted by at least one method	Predicted_03997	4956487	4957458	-1	hypothetical protein	
4951197	5040496	89299	Predicted by at least one method	Predicted_03998	4957527	4959452	-1	chromosome segregation	

			one method						protein
4951197	5040496	89299	Predicted by at least one method	mdaB_1	Predicted_04399	4961048	4961197	1	Modulator of drug activity B
4951197	5040496	89299	Predicted by at least one method	mdaB_2	Predicted_04400	4961194	4961361	1	Modulator of drug activity B
4951197	5040496	89299	Predicted by at least one method		Predicted_04391	4962775	4963239	-1	Replication protein
4951197	5040496	89299	Predicted by at least one method		Predicted_04403	4964839	4965030	1	hypothetical protein
4951197	5040496	89299	Predicted by at least one method	wbbD	Predicted_04340	4968011	4968844	-1	UDP-Gal:alpha-D-GlcNAc-diphosphoundecaprenol beta-1,3-galactosyltransferase
4951197	5040496	89299	Predicted by at least one method		Predicted_04341	4968965	4969831	-1	Polysaccharide pyruvyl transferase
4951197	5040496	89299	Predicted by at least one method		Predicted_00176	4974491	4974892	1	hypothetical protein
4951197	5040496	89299	Predicted by at least one method		Predicted_00177	4975183	4975827	1	Pentapeptide repeats (8 copies)
4951197	5040496	89299	Predicted by at least one method		Predicted_00178	4976465	4976905	1	hypothetical protein
4951197	5040496	89299	Predicted by at least one method		Predicted_04203	4980321	4980827	-1	hypothetical protein
4951197	5040496	89299	Predicted by at least one method		Predicted_04204	4980838	4983597	-1	hypothetical protein
4951197	5040496	89299	Predicted by at least one method		Predicted_04401	4987113	4987379	1	hypothetical protein
4951197	5040496	89299	Predicted by at least one method		Predicted_04408	4995363	4995554	-1	hypothetical protein
4951197	5040496	89299	Predicted by at least one method		Predicted_04392	4996794	4997546	-1	TcdA/TcdB pore forming domain
4951197	5040496	89299	Predicted by at least one method		Predicted_04298	5001994	5002113	-1	hypothetical protein
4951197	5040496	89299	Predicted by at least one method	epsE	Predicted_04390	5005779	5006264	1	Putative glycosyltransferase EpsE
4951197	5040496	89299	Predicted by at least one method		Predicted_04354	5011714	5011806	1	hypothetical protein
4951197	5040496	89299	Predicted by at least one method		Predicted_04267	5014092	5014322	1	Prophage CP4-57 regulatory protein (AlpA)

4951197	5040496	89299	Predicted by at least one method	papC_6	Predicted_03857	5018927	5021290	1	Outer membrane usher protein PapC precursor
4951197	5040496	89299	Predicted by at least one method	papD_6	Predicted_03858	5021297	5022022	1	Chaperone protein PapD precursor
4951197	5040496	89299	Predicted by at least one method		Predicted_03859	5022026	5023003	1	hypothetical protein
4951197	5040496	89299	Predicted by at least one method	rcsB_2	Predicted_03860	5023073	5023726	1	Transcriptional regulatory protein RcsB
4951197	5040496	89299	Predicted by at least one method	rcsC_2	Predicted_03861	5023704	5026910	-1	Sensor histidine kinase RcsC
4951197	5040496	89299	Predicted by at least one method		Predicted_04394	5028347	5029108	1	hypothetical protein
4951197	5040496	89299	Predicted by at least one method	epsJ_2	Predicted_04356	5030150	5031004	-1	putative glycosyltransferase EpsJ
4951197	5040496	89299	Predicted by at least one method	gtf1	Predicted_04357	5031178	5031606	-1	Glycosyltransferase Gtf1
4951197	5040496	89299	Predicted by at least one method		Predicted_04319	5033279	5034010	-1	hypothetical protein
4951197	5040496	89299	Predicted by at least one method		Predicted_04320	5034029	5034379	-1	hypothetical protein
4951197	5040496	89299	Predicted by at least one method		Predicted_04321	5034528	5034683	-1	hypothetical protein
4951197	5040496	89299	Predicted by at least one method		Predicted_03779	5039708	5040154	1	hypothetical protein
4951197	5040496	89299	Predicted by at least one method	hrsA	Predicted_03780	5040176	5040496	1	Heat-responsive suppressor HrsA
4951197	5040496	89299	Predicted by at least one method	fruA_2	Predicted_03781	5040483	5041601	1	PTS system fructose-specific EIIBC component
4961194	4968844	7650	Predicted by at least one method	mdaB_1	Predicted_04399	4961048	4961197	1	Modulator of drug activity B
4961194	4968844	7650	Predicted by at least one method	mdaB_2	Predicted_04400	4961194	4961361	1	Modulator of drug activity B
4961194	4968844	7650	Predicted by at least one method		Predicted_04391	4962775	4963239	-1	Replication protein
4961194	4968844	7650	Predicted by at least one method		Predicted_04403	4964839	4965030	1	hypothetical protein

4961194	4968844	7650	Predicted by at least one method	wbbD	Predicted_04340	4968011	4968844	-1	UDP-Gal:alpha-D-GlcNAc-diphosphoundecaprenol beta-1,3-galactosyltransferase
4995363	5006264	10901	Predicted by at least one method		Predicted_04408	4995363	4995554	-1	hypothetical protein
4995363	5006264	10901	Predicted by at least one method		Predicted_04392	4996794	4997546	-1	TcdA/TcdB pore forming domain
4995363	5006264	10901	Predicted by at least one method		Predicted_04298	5001994	5002113	-1	hypothetical protein
4995363	5006264	10901	Predicted by at least one method	epsE	Predicted_04390	5005779	5006264	1	Putative glycosyltransferase EpsE
5073122	5079853	6731	Predicted by at least one method	aacA4_2	Predicted_04218	5073122	5073721	-1	Aminoglycoside N(6')-acetyltransferase type 1
5073122	5079853	6731	Predicted by at least one method		Predicted_04256	5075851	5076714	-1	hypothetical protein
5073122	5079853	6731	Predicted by at least one method	ant1_4	Predicted_04257	5077043	5077987	-1	Streptomycin 3"-adenylyltransferase
5073122	5079853	6731	Predicted by at least one method		Predicted_04327	5079515	5079853	1	hypothetical protein
5079946	5085829	5883	Predicted by at least one method	smfA_7	Predicted_04328	5079946	5080479	1	Fimbria A protein precursor
5079946	5085829	5883	Predicted by at least one method	papH_6	Predicted_04329	5080526	5081083	1	PAP fimbrial minor pilin protein precursor
5079946	5085829	5883	Predicted by at least one method	xerD_6	Predicted_04021	5083027	5083629	-1	Tyrosine recombinase XerD
5079946	5085829	5883	Predicted by at least one method		Predicted_04022	5083868	5084329	1	hypothetical protein
5079946	5085829	5883	Predicted by at least one method		Predicted_04023	5085335	5085829	-1	hypothetical protein

Molecular characterization of *Providencia rettgeri* clinical strains carrying the *bla*_{NDM} and *bla*_{TEM} genes

Abstract

Providencia rettgeri is a Gram-negative bacillus widely distributed in the environment and is currently considered an emerging pathogen, mainly associated with nosocomial infections. The aim this study was to analyze the profile genomic of the two *P. rettgeri* strains isolated from clinical samples. The presence of the *bla*_{NDM} and *bla*_{TEM} genes was determined by *Multiplex-PCR*. The genomes of the PR01 and PR02 strains, characterized as carrier of the *bla*_{NDM-1} and *bla*_{TEM} genes were sequenced using the Illumina-MiSeq platform. After sequencing, the pre-assembled genomic DNA sequences were annotated using the Prokka software. A complementary genome analysis was performed using the Fast Annotation Subsystem (RAST) technology. A partial genomic characterization for the *P. rettgeri* PR01 and *P. rettgeri* PR02 strains using RAST indicated a wide range of genes related to efflux pumps of antimicrobial drugs (genetic systems were characterized, such as multiple drug extrusion (MATE), and the large family of the facilitator (MFS), in which a total of 91 genes were identified that were related to the virulence, disease and defense subsystem. In addition, several genetic systems were identified for the production and release of siderophores, which are associated with iron absorption, and other genes as receptors and transporters for heme ring and hemin. About 43 genes were identified and expressed proteins for uptake and metabolism. Analysis of the genome of the *P. rettgeri* PR01 and *P. rettgeri* PR02 strains showed a high concentration of insertion sequences, mobile and *prophages* elements, suggesting the existence of a genome. It was also verified the presence of several genomic islands, associated with persistence markers in relation to chemical and nutritional stress, such as the toxin-antitoxin system, containing the *PasI* and *PasT* genes. The diversity of genetic elements in *P. rettgeri* related to mechanisms of antimicrobial resistance can lead in the short term to an inefficiency of antimicrobial therapies in the fight against infections caused by this microorganism and its rapid access as pathogen is a result of great genetic plasticity.

Introduction

The *Providencia* genus is part of the Enterobacteriaceae family that contain Gram-negative bacilli With mobility producers of urease. Currently, the genus *Providencia* genus is composed of nine species named *P. alcalifaciens*, *P. stuartii*, *P. rettgeri*, *P. rustigianii*, *P. heimbachae*, *P. vermicola*, *P. sneebia*, *P. burhodogranariae*, and *P. thailandensis* (Shima et al., 2016). Some bacterial species of the genus contribute to a series of human infections, highlighting, among them the *P. stuartii* and *P. rettgeri* species (Washington and Barnhill, 2015).

The species *P. rettgeri* can be found in many environments such as water, soil and still inhabiting plant tissues, also part of the normal flora of the human gastrointestinal tract (Clifford et al.; 2012). *P. rettgeri* has been implicated in infections of the urinary tract, and in some cases these infections are associated with gastroenteritis and bacteremia frame (Yoh et al., 2005; Tada et al., 2014). *P. rettgeri* It is considered an emerging pathogenic bacteria and can exhibit high antimicrobial resistance rates commonly used in medical practice (Tshisevhe et al., 2017). *P. rettgeri* has intrinsic resistance to antibiotics polymyxin B and E (colistin), which are antimicrobial drugs of last choice for the treatment of infections caused by bacteria resistant to β -lactam antibiotics such as carbapenems (Olaitan et al., 2016).

P. rettgeri is a bacterial species that have been described in recent years as a microorganism key to spread the gene New Delhi Metallo- β -lactamase (Commonly called *bla_{NDM}*) in South America (Carvalho-Assef et al, 2013; Marquez-Ortiz, 2017). Enzymes classified as NDM, are characterized as carbapenemases, currently featuring 16 variants detectable in Gram-negative bacteria. These enzymes are capable of inactivating some of the class of β -lactam antibiotics (Khan; Maryam; Zarrilli, 2017). The NDM enzyme variants are widely distributed among Gram-negative bacteria and are responsible for triggering an increased level of bacterial resistance to antimicrobials agents (Khan; Maryam; Zarrilli, 2017).

The NDM-1 enzyme, which is encoded by *bla_{NDM-1}* gene was first detected in

of the *Escherichia coli* and *Klebsiella pneumoniae* strains isolated from clinical samples from a Swedish patient, whose medical history showed his hospitalization in India in the year of 2009 (Pillal; McGeer, Low, 2011). The spread of *bla*_{NDM-1} gene and its variants presents a challenge for health professionals on how to deal with infected patients (Olaitan et al., 2016).

The high mobility *bla*_{NDM} gene is related to the mechanisms of horizontal transfer of genes (THG), which occurs mainly via plasmid (conjugation events), coupled to the bacteria against the environment (Rolain; Parola; Cornaglia, 2010). These resistance genes are readily mobilized due to their location on mobile genetic elements (MGEs), as transposon Tn125 containing 10092 pb also found in *Acinetobacter baumannii* specie (Poirel et al, 2012). Germinally these meetings greatly contribute to bacterial survival (Jackson et al., 2011). The *bla*_{NDM-1} gene has been detected in plasmid pMR0211 in *P. stuartii* (Gann et al, 2012), and the plasmid pPrY2001 carried by *P. rettgeri* M15628 (Mataseje, 2014). However, *bla*_{NDM-1} gene in some bacterial strains may be integrated into the bacterial chromosome, as evidenced in *P. rettgeri* (Gefen-Halevi et al., 2013).

After his description in 2009, *bla*_{NDM-1} gene It was identified in bacteria isolated from clinical samples of UK hospital patients, India and other regions of the world, especially in isolated *E. coli* and *K. pneumoniae* (Pillal; McGeer, Low, 2011). In Brazil in 2013 bacterial strains harboring the *bla*_{NDM-1} gene have been detected in clinical samples of two patients in a hospital of Rio Grande do Sul State, strain CCBH11880 with Genbank accession number: GCA000805715.1 (Carvalho-Assef et al., 2013). The second detection *bla*_{NDM-1} gene in bacteria circulating in patients was reported in Brazil in the São Paulo. In this description a patient was colonized and infected by another *P. rettgeri* (Carmo-Junior et al., 2015).

The study main objective characterization of the genes and molecular mechanisms involved in the spread of resistance to carbapenems and other antimicrobial drugs in clinical isolates of *P. rettgeri*. Further, characterizing the genes and proteins related to virulence in two strains of *P. rettgeri* PR01 and PR02 strains from partial sequencing of the genome.

99 **Material and methods**

100 **Collection and identification of bacterial isolates**

101 This study included eight (8) strains of *Providencia rettgeri* obtained
102 from clinical specimens from blood culture and urinary sample from patients
103 treated in Maranhão state hospitals. These micro-organisms were provided the
104 routine service of Analysis Laboratory Clinica Cedar, located in São Luis-MA.

105 The identification of the bacterial species was obtained by MALDI-TOF
106 MS system (Matrix-Assisted Laser Desorption Ionization-Time of Flight Mass
107 Spectrometry) using Biotyper system (Bruker, Billerica, MA). Bacterial cells were
108 grown on TSB agar plates (Himedia, India) for 24 h at 37 ° C. For the test, a
109 sample of colony was transferred with the aid of a bacteriological loop metered 1
110 uL to a metal blade so as to present a thin layer over the detection plate, and then
111 were added to sample 1 µL of a solution of matrix (α -cyano-4-hydroxycinnamic
112 acid). The samples were subjected to the incidence of a laser beam for extracting
113 energized ribosomal peptides. Finally, the mass spectra acquired for each
114 bacterial strain were compared with the known mass spectra contained in the
115 classification software (Version 3.1, Library 1.0).

116 **Susceptibility tests to antimicrobial agentes**

117 The susceptibility profile to antimicrobial drugs of bacterial strains was
118 determined on the AST No. 105 cards for automated VITEK 2 (CLSI, 2016)
119 (BioMerieux SA, Marcy-l'Etoile, France). Using antibiotics, amikacin, ampicillin,
120 ampicillin / sulbactam, cefepime, ceftazidime, ceftriaxone, cefuroxime,
121 cefuroxime axetil, ciprofloxacin, ertapenem, gentamicin, imipenem, meropenem,
122 piperacillin / tazobactam. The sensitivity analyzes were performed according to
123 the manufacturer's recommendations. Bacteria were stored in BHI plus 20%
124 glycerol and stored in cryovials at -80 ° C.

125 **Molecular identification of the *bla*_{KPC}, *bla*_{NDM-1} *bla*_{TEM}, and *bla*_{SHV} *bla*_{AmpC}** 126 **genes**

127 **DNA Extraction**

Research on the presence of genes related to β -lactamase enzymes was conducted by multiplex-PCR reactions. For this purpose, bacterial cultures obtained from bacterial growth were used at 37 °C for up to 18 hours in medium liquid BHI (Difco, Detroit, MI, USA). After incubation, 500 μ L of culture was centrifuged at 10,000 rpm for 20 minutes to obtain a pellet for using DNA extraction kit "Wizard® Genomic DNA Purification" (Promega Corporation, Madison, WI USA), following the original protocol manufacturer. Genomic DNA was quantified in Nanodrop™ 1000-260 and 280 nm and used for amplification reactions related gene fragments resistance to β -lactams.

Multiplex-PCR

Multiplex-PCR reactions were performed using a specific pair of primers (primers) in a final volume of 25 μ L reaction containing 12.5 μ L of a solution of GoTaq® Green Master Mix (Promega, Madison, WI, USA). The sequences of the specific primers that were used in this study are listed in Table 1

Amplification reactions of the *multiplex-PCR* assay were performed according to standard methods by the following authors: Cunningham et al. (2013), Liu et al., (2012), Khalilzadegan et al., (2016) and Dallenne et al. (2010). PCR products were separated on agarose gel at a concentration of 1.5% and visualized under ultraviolet light exposure. For *bla_{NDM}* gene sequencing, the fragments were subjected to sequencing reactions using the DYEnamic ET Terminator Cycle Sequencing kit (GE Healthcare Life Sciences, Buckinghamshire, UK) according to the manufacturer's instructions. The fragments were analyzed on the automatic sequencing system ABI PRISM® 3100 Genetic Analyzer (Applied Biosystems, USA). The quality of the sequences obtained electropherograms during the sequencing process was analyzed with the software ChromasPro (<http://www.technelysium.com.au/chromas.html>). For alignment, at least three consensus sequences of each sequenced fragment NDM type were selected from the database for alignments using MEGA 6.0 (Tamura et al., 2013). Similarities between the obtained nucleotide sequences were verified using available BLASTn <https://blast.ncbi.nlm.nih.gov/Blast.cgi>.

To identify the gene variants *bla*_{NDM}, all sequences were translated into amino acids using the software ExPASy [translation tool (<http://web.expasy.org/translate/>)]. The correct translation was chosen based on data available in GenBank. The amino acid sequences were compared with the sequences obtained from the GenBank protein NDM using BLASTx. The values of similarity to the amino acid sequences ranged 99 to 100%, indicating a highly conserved region.

2.4 Characterization of the incompatibility groups of plasmids

The determination of incompatibility groups of plasmids was performed according to the method described by Carattoli et al. (2005). Through several PCR reactions (5 multiplexes and 3 simplex) are tested the most representative types of plasmids of the Enterobacteriaceae family, such as FIA, FIB, FIC HI1, HI2, I1-Iy L/M, N, P, W, T, A/C, K, B/O, X, Y, F and FIIA. PCR reactions were performed in a final volume of 25 µL reaction containing 12.5 µL of a solution of GoTaq® Green Master Mix (Promega, Madison, WI, USA) and 100 ng of template DNA per reaction. The thermocycler program, except for that used for simplex-F reaction (whose difference is the annealing temperature of 52 ° C), consisted of an initial step of 5 minutes at 94 ° C followed by 30 cycles of 1 minute at 94 ° C 30 seconds at 60 ° C and 1 min.

Polymerase chain reaction for the detection of multiplex-integrans class 1, 2 and 3

The characterization of integrans it was performed having as the target sequences intl1, intl2 and intl3, these were amplified by PCR reaction method, using multiplex primers described by Goldstein et al (2001). PCR reactions were performed using a specific pair of primers (primers) in a final volume of 25 µL reaction containing 12.5 µL of a solution of GoTaq® Green Master Mix (Promega, Madison, WI, USA), 25 µmol of each primer and 1 µL DNA template (100 ng DNA/reaction). The PCR conditions were used as described by Khoramrooz et al. (2016). The amplified products 280 bp to integron class 1, class 233 bp to 600 bp integron 2 and the class 3 integron were separated by electrophoresis in 1% agarose gel containing 0.5 µg/mL ethidium bromide.

Sequencing of the genome and notes

The genome of lines PR01 and PR02 blaNDM carrying the gene were subjected to data obtained by the assembly process MiSeq as A5 pipeline aid (<http://www.ncbi.nlm.nih.gov/pubmed/23028432>). This uses a new sequencing approach for prokaryote genomes A5 (Andrew And Aaron's Awesome Assembly pipeline). The pipeline and associated programs are open source distributed under GPLv3 license, runs under Linux and is installed on the local server for bioinformatics analysis Neopropecta (Setting: 03 Linux servers, Core i7, 64GB RAM, 2TB HD).

The genes were sequenced as predicted using the inference constant BLAST protein (<http://www.ncbi.nlm.nih.gov/>). From this analysis, it was identified related sequences in other bacterial species of interest listed in the database RAST (Rapid Annotation using Subsystem Technology), which is an important tool bioinformatics to predict putative genes (available on <http://rast.nmpdr.org>).

Construction of the phylogenetic tree

The sequences of *P. rettgeri* PR01 and *P. rettgeri* PR02 strains were subjected to analysis to this RNAmmer software: <http://www.cbs.dtu.dk/services/RNAmmer> (Lagesen et al., 2007) for identification of the 16S ribosomal gene. The analyzes were performed with the *BLAST algorithm*, using at least one sequence of the 16S rDNA gene. Some sequences of 16S rDNA of the genus *Providencia* were used for the preparation of phylogenetic relationships. To determine the root of the tree (outgroup) was used partial sequence of the 16S rRNA (ribosomal RNA) *Serratia plymuthica* K-7 (NR_037111). The sequences obtained from bacterial species were aligned so the software MEGA 7.0 (Kumar; Stecher; Tamura, 2016). For construction of the phylogenetic tree was used the maximum likelihood method (*Maximum Likelihood*) MUSCLE applying the algorithm (Multiple Sequence Comparison by Log-Expectation) (Edgar; Drive; Valley, 2004). The values of *Bootstrap* lower 70% were hidden in forming the phylogenetic tree of Fig.

Analysis of genomic islands

The genomes of the *P. rettgeri* PR01 and *P. rettgeri* strains PR02 were analyzed for the presence of genomic islands, pathogenicity islands (PAIs) and antibiotic resistance (RIs). For these analyzes, the genome draft were worked in house using scripts to the concatenation of contigs / scaffolds. For these analyzes it was first a search for bacteria of the same genus genomes or as close as possible to the genomes of strains *P. rettgeri* PR01 and *P. rettgeri* PR02. In this quest came to *Providencia alcalifaciens* DSM 30120. The analyzes were conducted with *P. rettgeri* PR01 (query) and a reference genome, in this case *P. alcalifaciens* (subject) for prediction of genomic islands. After the islands predicted, a figure in BRIG was generated using *P. rettgeri* PR01 reference. In the same figure, They were added to the coordinates of the islands previously predicted. Additionally, most were plotted two rings contigs corresponding to the genome of *P. rettgeri* PR01, where the pairs contigs were marked in blue and the odd in green. A manual curation was made on the data generated by the Gipsy software. Thus, PAIs were identified, gels and mixed Islands (MSIs), these regions are predicted as much as PAIs, as IRs by Gipsy software. Parallel genomic islands *P. rettgeri* PR01 and *P. rettgeri* PR02. They were predicted using software IslandViewer 4 (Bertelli et al., 2017) associated with the programs IslandPick, SIGI-HMM, and IslandPath-DIMOB.

Results

Determination of the susceptibility profile to antimicrobial agents

In this study it was observed that eight (08) strains of *P. rettgeri* designated as PR01, PR02, PR12, PR13, PR18, PR20, PR21 and RP27 showed resistance to multiple antimicrobial drugs of different classes (Table 2), amikacin, and gentamicin (aminoglycosides) resistance with an MIC $\geq$ 16 μ g/mL, cephalosporins MIC $\geq$ 8 μ g/mL and with carbapenem MIC $\geq$ 4 μ g/mL. Where PR27 was the only isolate that showed a pattern sensitivity to all tested antibiotics, including β -lactam antibiotics.

Detection of genes related enzymes β -lactamases

In this work, the *bla*_{TEM} genes have been identified, and *bla*_{AmpC} *bla*_{NDM} in all strains of the *P. rettgeri*. However for *bla*_{SHV} *bla*_{KPC} genes and the results

were negative in all strains of *P. rettgeri*, as shown in Table 3.

Identification and characterization of the incompatibility groups and integrons

All strains of *P. rettgeri* presented integrons class 1 and 2 as well as the plasmid incompatibility group FIIA as seen in Table 3.

The characterization of plasmid incompatibility groups, it was observed that all isolates of *P. rettgeri* replicon were FIIA carriers with integrons class 1 and 2 (Table 3).

Phylogenetic analysis of *P. rettgeri* PR01 and *P. rettgeri* PR02 strains

The evolutionary relationships among isolates *P. rettgeri* PR01 and *P. rettgeri* PR02 were determined by maximum likelihood method, the results indicated that the closer to *P. rettgeri* were isolated strains DSM4542 *P. rettgeri* NCTC11801 (89% bootstrap) and having a genetic similarity to strain RB151 as seen in figure 2.

Genomic characterization of *P. rettgeri* PR01 and *P. rettgeri* PR02 strains

The general data annotation of partial genomes of the *P. rettgeri* PR01 and *P. rettgeri* PR02 strains are shown in Table 04. The identification of genomic or coding sequences (CDS, coding sequence) was present in the lines 4425 and 4477 for PR01 and PR02 strains, respectively. In RAST analysis performed by several genes were observed associated with various categories of subsystems (Figures 3,4). However, this study highlight the proteins associated with efflux pumps to antibiotics and biocides, iron shot to proteins, proteins associated with motility and elements associated with the gene transfer mechanisms. These elements were compared by BLASTn and finally their identities were then given in Supplement I. In addition, We verified the presence of genes associated with mobility genome such as those related to protein mobilities and bacteriophages. Finally, the analysis of genes and related proteins were also directed to the related sub-system metabolism and capture iron.

Resistance profile characterization and virulence of the *P. rettgeri* PR01 and

***P. rettgeri* PR02 strains**

Drug efflux pumps and biocides

For a more specific analysis of the genetic features of the *P. rettgeri* PR01 and *P. rettgeri* PR02 strains, these were characterized at the genomic level after the next generation sequencing using Illumina- Miseq system. In these analyzes were identified sixteen (16) types of efflux pumps antimicrobial agents and biocides (Figures 5 and 6).

Among the sixteen (16) efflux pumps present in *P. rettgeri* PR01 and *P. rettgeri* PR02 strains were identified apple (Macrolide-specific efflux protein apple), MACB (Macrolide export ATP-binding/permease protein Mac TolC (RND efflux system, outer membrane lipoprotein CMEC), AcrB (RND multidrug efflux transporter, Acriflavin resistance protein), MATE_Family_MDR_Pump (Multi antimicrobial extrusion protein (Na (+) / drug antiporter, MATE family of MDR efflux pumps).

Characterization of iron uptake system

The *P. rettgeri* PR01 and *P. rettgeri* PR02 presented six categories of genes related to iron uptake and metabolism, and in two major groups represented caputra categories heme proteins and related Hemia with 22 followed by 14 Hemin transport proteins. As seen in Figure 7, having the iutA genes associated proteins, and FhuC with the location in BLASTn gi |490378242| WP_004257840.1 with 99% identity to iutA, and two genes for FhuC with their locations gi | 490382157 |WP_004261670.1 and gi | 757595221 | WP_042843418.1 both with 100% identity. Introducing a wide range of systems siderophoros the type and aerobactin iron transport (TonB) shown in Supplement I.

The PR01 and PR02 strains, have the iutA genes associated proteins, and FhuC with the location in BLASTn gi | 490378242 | WP_004257840.1 with 99% identity to iutA, and two genes for FhuC with their locations gi | 490382157 | WP_004261670.1 and gi | 757595221 | WP_042843418.1 both with 100% identity.

Characterization Genomic Islands

In the analysis of genomic islands carried by IslandViewer program 4 was observed 22 genomic islands in the genomes of PR01 and PR02 lines as seen in Figures (11, 12) with a great diversity of genes related to bacterial virulence, drug resistance antimicrobial and mercury, toxin-antitoxin system, generic control, among others in the supplement II. The major identified genomic island in strain showed PR01 114 821 bp (> 10 kb) and harboring the gene for resistance to aminoglycosides and β -lactam. This genomic island are the *bla*_{TEM} and *bla*_{NDM} gene, and a gene for the transposon Tn3 resolvase associated. Parallel to the analysis program BRIG 6 pathogenicity islands compared with strains 6, 2 and 4 *P. rettgeri* *P. stuartii* with an identity of 100% as shown in Figure 10.

Analysis of bacterial mobiloma

Insertion Sequences

All analyzes the genome of the *P. rettgeri* PR01 strain using Issaga software (Sequence Insertion semi-automatic annotation genome were detected twelve (12) Putative types of insertion, with a total of 54 elements, with the largest percentage for inserts were to IS3 with 19.64% and 19.23% for PR01 to PR02 iS5 followed by 12.5% to 13.46 to *P. rettgeri* PR01 and *P. rettgeri* PR02. About 8.93% of insertion sequences and PR01 11.54% in PR02 were not identified by the database, being named ISNCY (English, IS not yet classified, unclassified IS).

All analyzes were also identified that about 10.71% and 13.46% of insertion elements PR01 and PR02, respectively, belonging to the transposon Tn3. The Tn3 transposon was characterized with an average variation 2306-3282bp presenting an identity of 99% (2892/2896) housed in sequence with the plasmid carried by pNDM15-1091 *P. rettgeri* N15-01091 strain (Genbank code: CP012903.1). This nucleotide sequence, in turn mainly relates to a gene for a 1845 bp transposase.

DNA sequences associated with phage

In the analysis of CDs for bacteriophages sequences in genome

performed by PHAST program were observed 6 regions related to these elements. Of the sequences found five (05) presenting was incomplete 2.4 Kb to 30 Kb in size, and related sequences found in bacteria of the family Enterobacteriaceae (such as *Salmonella* spp.), and as Gram-positive *Staphylococcus* spp. and mycobacteria. The only intact bacteriophage sequence found was related to phage (PHAGE_Salmon_RE_2010_NC_019488), a sequence of 33.7 kb (at genome position: 4233321-4267110). This sequence shows 49 CDs and the majority of these sequences are derived from a bacteriophage of *Salmonella* spp. (Code in GenBank NC019488).

Transposable elements

In analyzes performed by RAST, it was observed several genes associated with transposable elements or transposons. These elements had their identities compared by BLASTn algorithm. *P. rettgeri* PR01 and *P. rettgeri* PR02 harboring two types of transposable elements, these elements are responsible for implementing proteins, namely Tn7 and Tn21.

Still, it was possible to verify that sequences associated with Tn7 are present in both *P. rettgeri* PR01 and *P. rettgeri* PR02 strains being related to fragments of 1026 and 882 bp which are located integrons Tn7. These genes are responsible for the implementation of tnsA protein, and protein tnsB, respectively. Yet PR02 strains contains only one TN21 associated sequence indicated by the sequence 372 bp in PR01 already has three sequences related to TN21 fragments of 372, 252 and 240 bp with their respective locations and the% identity of BLASTn [gi | 519733601 | EPP24804.1](#) and 98% ID, [gi | 1119200185 | WP_072209121.1](#) 100% ID [gi | 1119200185 | WP_072209121.1](#) ID 99%.

Discussion

Profile susceptibility to antimicrobial agents

In this study, 8 isolates of *P. rettgeri* PR01, PR02, PR12, PR13, PR18, PR20 and PR21 were analyzed for antimicrobial agents susceptibility profile. Six

bacterial isolates were resistant carbapenem antibiotics, namely ertapenem, imipenem and meropenem MIC was with in ≥ 16 $\mu\text{g/mL}$ (CLSI, 2016). A particular feature of these bacterial isolates is that they originate from two hospitals located 300 km away. Focusing on an inter-hospital transmission, occurring by professionals or patients transferred between the units. Since carbapenems are among the best options for treatment of infections by Gram-negative microorganisms resistant to multiple antimicrobial drugs (Johnson, Woodford, 2017; Haciseyitoglu et al., 2017; Leylabadl et al., 2015), it is evident therefore an alarming situation public health, where these microorganisms are a common source of hospital-acquired infections (Rolain; Parola; Cornaglia, 2010). The profile arising MDR genes related efflux pumps, or more specific genes such as those coding for β -lactamase enzymes, such as *bla*_{NDM} and *bla*_{KPC} genes. These genes are commonly found in species of bacteria of the Enterobacteriaceae family (Wu et al., 2015).

β lactamases Genes related enzyme

The *bla*_{TEM} and *bla*_{AmpC} genes are responsible for the expression of a cephalosporinase ESBL type and a β -lactamase AmpC type, respectively. Resistance to antibiotics amikacin, gentamicin, ampicillin, sulbactam, cefepime, ceftazidime, ceftriaxone, cefuroxime, ciprofloxacin, piperacillin, and tazobactam is given the correlation of various defense mechanisms that have MDR's as efflux pumps, and specific enzymes (Li; Plésiat, Nikaido 2015).

The gene *bla*_{TEM} expresses a cephalosporinase ESBL the type often detected in *E. coli* and *Klebsiella* spp. (Oduro-Mensah et al., 2016). The presence of the *bla*_{TEM} in *P. rettgeri* indicates a high probability that this occurrence is being made possible by the transfer of movable horizontally transposable elements. A feature of the resistance genes and their association with a Cassette region of integrons region which is located between two recombination sites (ETTA and attC), initial recombination sites and termination, respectively, in this way the resistance genes may be integral DNA molecule (Deng et al., 2015).

The gene *bla*_{NDM} has shown a clinical relevance in recent years, since this gene is responsible for the production of carbapenemase enzyme, a metallo-

β-lactamase zinc-dependent, acting in the hydrolysis of the carbapenem antibiotics (Galdiero et al., 2012). The carbapenemases are usually able to hydrolyse not only carbapenems, but also all β-lactam antibiotics such as cephalosporins, monobactams and penicillin (Bernabeu, et al., 2012). **Integrans and incompatibility groups**

Integrans class I and II were identified in strains of *P. rettgeri* all clinical samples obtained from the aspect of resistance presented possibly the resistance markers βlactam are present recombination between the sites of integrans, which are commonly drug resistance or pathogenicity genes (Rajpara et al., 2015). Such genes may be allocated in the chromosome since the presence of integrans class I together with Tn3 family of transposons enable such insertion gene (Deng et al, 2015; Rajpara et al, 2015).

The incompatibility group FIIA detected in strains *P. rettgeri strains* gives a unique pattern with respect to transconjugate barriers between these strains. The FIIA replicon (replicon typing) is called conjugative plasmid or *incf incf* is the reference for the design of primers (primer) (Carattoli et al., 2005). This incompatibilidade group was first identified in *Salmonella enterica* (Typhimurium). The replicon FIIA has also been identified in a plasmid recovered from a strain of *Salmonella enterica* serovar Kentucky, isolated from chicken fecal samples (Fricke et al., 2009). The FIIA incompatibility group was related to virulence factors and resistance to streptomycin and tetracycline these *Salmonella enterica* strains (Fricke et al., 2009). However, there are few reports on the presence of the FIIA incompatibility group in other species of bacteria in the Enterobacteriaceae family.

Moreover, it is believed that the presence of *repliconFIIA* is rare in other species of Gram-negative bacteria, yet its acquisition by other species such as *P. rettgeri* may be possible, especially if these species relate in environments such as the gastrointestinal tract of animals such as birds or human hosts. Recently, FIIA replicon was identified in strains of *Enterobacter* spp. clinical origin (Logan et al., 2016). These bacterial strains were carriers of *blaACT* / *MIR* genes that confer resistance to oxyimino-β-lactams and other β-lactam antibiotics (Logan et al., 2016). It appears then, that there is a possibility of replicon FIIA not

be restricted to *Salmonella* spp. as had previously been proposed, can circulate in other genera of bacteria, however with a lower transmission frequency.

Phylogenetic analysis of *P. rettgeri* PR01 and *P. rettgeri* PR02 strains

The *P. rettgeri* PR01 and *P. rettgeri* PR02 strains showed a large genetic similarity to *P. rettgeri* strain RB151 that is a pathogen carrier *bla*_{NDM-1} gene was also related to the same group (89% bootstrap) and which is lineages, and PR01 PR02, the pathogen is one of the few strains of the species is described for carrying metallo- β -lactamase (Cardozo Castro et al., 2017).

The *P. rettgeri* RB151 strain was isolated in 2013, from a urine sample from a female patient of 58 years old. This patient was diagnosed with urinary tract infection and was being treated at the emergency department of a hospital in the Bucaramanga city, Colombia (Saavedra-Rojas et al., 2015). Interestingly, the *bla*_{NDM-1} gene present in strain RB151, It is housed in a so-called plasmid pRB151-NDM (Marquez-Ortiz et al., 2017), which may not occur with the lines PR01 and PR02. It was observed in a first analysis after partial sequencing, the plasmid PR01 (pPR01 9.8 kb) showed no *bla*_{NDM-1} the gene harbored only *bla*_{TEM} (supplement II). This plasmid showed 99% identity with the pCR14_2 plasmid carried by *K. pneumoniae* strain CR14 (ID in the Genbank: CP015394.1). Possibly *bla*_{NDM-1} gene may be integrated into the chromosome in *P. rettgeri* PR01 strain as has been previously reported for other species, as combination assays failed to demonstrate clinical transfer between the horizontal strains *E. coli* (positive NDM) and the permissive receptor strain *E. coli* J53 (Shaheen et al., 2013).

Analyzes of the subsystems of the *P. rettgeri* PR01 and *P. rettgeri* PR02 strains

In *P. rettgeri* PR01 and *P. rettgeri* PR02 strains the systems MATE, TolC, MacAB, AcrB RND are connected in a direct way resistance to antimicrobial drugs such, as fluoroquinolones (ciprofloxacin, norfloxacin, norfloxacin), rifampicin and macrolide antibiotics (ritromicina, azithromycin and clarithromycin). In addition, of bacterial resistance to cobalt and zinc, this evidence was correlated with the pattern of resistance in the MIC test.

In genomes of the PR01 and PR02 strains it was also possible to detect the presence of 04 (four) genes for β -lactamases, which are related to the resistance to penicillins ID BLASTn [gi | 926465406 | ALD19783.1](#) 100% identity, 1st and 2nd generation cephalosporins, BLASTn ID [gi | 490383288 | WP_004262799.1](#) with 99% identity and [gi | 446804399 | WP_000881655.1](#) 100% identity, 3rd and 4th generation BLASTn ID [gi | 1002048020 | AMM70781.1](#) 100% identity.

The iron-binding proteins in PR01 and PR02 isolates were iutA (aerobactin receptor), FhuB (ferrichrome permease) and FhuC (ATP binding protein), which are characterized as iron uptake receptors (Cabrera et al. 2001). Also, they are elements responsible for the breakdown of the microbial cell wall, consequently iron chelators such as enterobactin are released into the extracellular space (West et al., 1987).

The FhuC gene was previously identified in some species of Enterobacteriaceae such as *Edwardsiella ictaluri* (Abdelhamed et al., 2016) and *Yersinia enterocolitica* biovar 1 (Kanauija; Bajaj; Viridi, 2015). *E. ictaluri* the FhuC protein contributes to the acquisition of ferric hidroxamate (Abdelhamed et al., 2016). The detection of a great diversity of genes in *P. rettgeri* that are responsible for production of many elements such as siderophores and transport and membrane proteins that contribute to the uptake and transport of iron, indicate that these products may be important in virulence and rapid multiplication of this bacterial species.

Analysis of genomic islands and islands of pathogenicity

The great diversity of genes acquired by horizontal transfer in *P. rettgeri* PR01 and *P. rettgeri* PR02, indicate a high resiliency with respect to the species

to survive adverse environments and stressful conditions, such as in the presence of oxidizing agents, biocides, and environments with nutrient limitation.

The toxin-antitoxin system represented by different genetic sequences in the supplement II. The *P. rettgeri* PR01 and *P. rettgeri* PR02 found in genomic islands can contribute to biofilm formation on abiotic surfaces and colonization of host tissues, such as urinary tract, as well as generating persistent cells in inhospitable conditions, as observed for cells that survive in the presence antimicrobial drugs (Page; Peti, 2016; Wang; Wood, 2011). These assumptions were raised, taking into account that the toxin-antitoxin system is an important factor in *E. coli* uropathogenic for niche-specific colonization (urinary tract), resistance to stress caused by ciprofloxacin in addition to the nitrite oxidation resistance sodium and other reactive nitrogen intermediates (Norton; Mulvey, 2012).

Analysis of insertion and transposon sequences in the bacterial mobiloma

The transposase of Tn7 is heteromeric, which distinguishes itself by the complexity of its target site insertion transposition att Tn7, it requires four proteins, Tn7, TnsABC + D and two substrate DNA (Young et al., 2013). The Tn7 transposase protein has two, TnsA and TnsB. TnsA performs cleavage of the donor at the 5' transposition and transposition TnsB acting directly on the ends 3', causing splitting and joining the target DNA (Holder; Craig, 2010; Young et al, 2013).

The presence of DNA sequences in *P. rettgeri* PR01 and *P. rettgeri* PR02, Tn7 transposon Related indicates a high versatility of these strains in the spread of antibiotic resistance genes for this micro-organism. The transposon Tn7 is a particularly sophisticated mobile element, and developed in some Gram microorganisms negative, such as *E. coli* and *Acidithiobacillus ferrooxidans*, which have completely different lifestyles (Oppon et al., 1998; Peters, Craig, 2001). The Tn7 transposon has alternative mechanisms to promote their spread among bacterial populations. The movable element can move to low frequency insertion sites, such as transposable other furniture elements, in that implements in many places without the existence of a specific sequence in the DNA recipient.

Moreover, this element can preferably move to an insertion sequence for Tn7 in some replicons, preferably in conjugative plasmids. The latter contributes significantly to the spread of the bacterial population Tn7, thus resulting in the acquisition of genes associated with antibiotic resistance in many bacterial species (Peters, Craig, 2001).

The Tn21, is a transposon belonging to the Tn3 family, is an important determinant for resistance to antibiotics, for example aminoglycosides, mainly by the gene adduce for 2"-aminoglycoside nucleotidyl transferase ANT (2 " ') (Schmidt; Nucken; Henschke 1988). More recently it has been linked to cAMP Tn21 gene, which confers resistance to penicillins and cephalosporins in the 1st and K. *pneumoniae* strain LCT-KP214 and KP289-LCT (Guo et al., 2014). A particular feature of Tn21, is its high exchange capacity and resistance locus accumulation in plasmids and transposons other (Liebert, Hall; Summers, 1999). The transposition occurs by the Tn21 TnpA (transposase A), with the insertion sequence, a 5 bp duplication of target DNA, usually rich in AT (Liebert, Hall; Summers, 1999). As the horizontal and vertical transfer of genes related to antibiotic resistance in Enterobacteriaceae genus are derived from plasmids and transposons, end up covering a high variety of bacterial hosts.

The Tn3 transposon has been related to the presence *bla*_{TEM-1D} gene (a beta-lactamase class A) in some bacterial species of the family Enterobacteriaceae (Sandner-Miranda et al., 2016). The *P. rettgeri* PR01 and *P. rettgeri* PR02, characterized in this study *bla*_{TEM} carrying the gene, and there is probably a good correlation being the presence of the transposon Tn3 and the spread of this gene.

The contribution of insertion sequences for the genetic plasticity in *Providencia* spp. It has not been thoroughly studied, but can provide that its contribution can be very similar to those observed for other species of bacteria of the family Enterobacteriaceae. In *E. coli* O157 (a burial-hemorrhagic serotype), for example, it has been suggested that IS elements may have a role in the inactivation and immobilization of phage and plasmids received by bacterial strains (Ooka et al., 2009).

The presence of bacteriophage sequences in bacterial genomes, especially in enterobacteria may be the result of the acquisition by horizontal transfer genetic material into more specific niches such as the gastrointestinal tract of the host animals (Huddleston, 2014). This assumption is based on the evidence of the existence of a greater availability of viral genetic material into the intestinal lumen after its release during the lytic cycle of the bacteria in this niche (Sunagawa; Koonin; Bork, 2014). Experimental evidence using quantitative PCR for DNA bacteriophages present in human faeces indicates that these elements. They carry a large amount of resistance genes such as genes for β -lactamases and *bla*_{TEM} and *bla*_{CTX-M-1} (Modi et al., 2013). These observations can corroborate to measure the impact of bacteriophage genomes in movement and placement of antibiotic resistance genes for the bacterial population (Modi et al., 2013). Finally, it is believed that the bacteriophages associated with transduction events can increase the contribution mechanisms in resistoma the bacterial population in the gastrointestinal tract especially among Enterobacteriaceae, so have a great impact on the development of emerging pathogens such as *P. rettger*

This work showed various molecular characteristics analyzed after partial sequencing of the genome of two isolates of *P. rettgeri* (PR01 and PR02). These characteristics are related to the ability of these strains to endure the presence of various antibiotics, especially beta-lactam clinical use as cephalosporins and carbapenems. The presence of a wide variety of efflux pumps, in addition to inactivating enzymes aminoglycosides, beta-lactamases are expressed from the *bla*_{NDM} gene and are *bla*_{TEM} contributing to the rapid dissemination and accession of this species in Brazilian nosocomial environment, here in the first instance detected in isolated Maranhenses hospitals. It was also found that strains of *P. rettgeri*. They can adapt to various environments characteristics, especially with respect to iron capitation and self-regulation systems such, as system toxin-antitoxin. In addition, partial genome analysis also indicated that the species has a highly dynamic genome consists of a mobiloma wherein several gene segments related to mobility and bacteriophage protein genes. Finally, the dissemination of genetic elements can contribute to transmission of genes associated with virulence and resistance to biocides and bacterial antibiotic drugs and *other species of the Enterobacteriaceae family*.

Conflict of Interest Statement

The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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