





## Complete Genome Sequence of *Streptococcus pneumoniae* Serotype 19A, a Blood Clinical Isolate from Northeast Mexico

® Antonio Ali Perez-Maya,ª Rosa Maria Hinojosa-Robles,<sup>b</sup> Jose Ramon Barcenas-Walls,<sup>c</sup> Armando Vignau-Cantu,ª Hugo A. Barrera-Saldaña,ª Rocio Ortiz-Lopez<sup>a,c</sup>

Departamento de Bioquímica y Medicina Molecular, Facultad de Medicina, Universidad Autónoma de Nuevo León, Monterrey, Nuevo León, Mexico<sup>a</sup>; Hospital Universitario Jose Eleuterio Gonzalez, Universidad Autónoma de Nuevo León, Monterrey, Nuevo León, Mexico<sup>b</sup>; Unidad de Genómica; Universidad Autónoma de Nuevo León, Centro de Investigación y Desarrollo en Ciencias de la Salud, Monterrey, Nuevo León, Mexico<sup>c</sup>

We report here the draft genome sequence of a *Streptococcus pneumoniae* strain isolated in Monterrey, Mexico, MTY1662SN214, from a man with purpura fulminans. The strain belongs to the invasive and multidrug-resistant serogroup 19A, sequence type 320 (ST320). The draft genome sequence consists of 60 large contigs, a total of 2,069,474 bp, and has a G+C content of 39.7%.

Received 12 February 2016 Accepted 18 February 2016 Published 31 March 2016

Citation Perez-Maya AA, Hinojosa-Robles RM, Barcenas-Walls JR, Vignau-Cantu A, Barrera-Saldaña HA, Ortiz-Lopez R. 2016. Complete genome sequence of *Streptococcus pneumoniae* serotype 19A, a blood clinical isolate from Northeast Mexico. Genome Announc 4(2):e00195-16. doi:10.1128/genomeA.00195-16.

Copyright © 2016 Perez-Maya et al. This is an open-access article distributed under the terms of the Creative Commons Attribution 4.0 International license.

Address correspondence to Rocio Ortiz-Lopez, rortizlopez@gmail.com

A fter the introduction of the 7-valent pneumococcal conjugate vaccine (PCV7) in the United States and other western countries, there was a significant reduction in the incidence of invasive pneumococcal disease (IPD) (1, 2). However, the number of cases of IPD, caused by *Streptococcus pneumoniae* serotypes not included in PCV7 (2, 3), increased. Particularly, the serotype 19A-sequence type 320 (ST320) has emerged worldwide as an important pathogen associated with IPD and multidrug resistance (MDR), but the causal factors remain unclear (4–6).

S. pneumoniae 19A ST320 has been implicated in both IPD and noninvasive pneumococcal disease (6). ST320 19A is a double-locus variant of the originally described Taiwan19F-14 (ST236) clone with high antibiotic resistance, which was isolated from a Taiwanese hospital in 1997. This strain underwent capsular modification, likely through recombination that altered the wzy polymerase gene and gave rise the serotype 19A-ST320 (7, 8).

S. pneumoniae strain MTY1662SN214 was isolated from a blood sample in 2014 at the Dr. José Eleuterio González University Hospital, Universidad Autónoma de Nuevo León (UANL), in Northeast Mexico. The patient was a 43-year-old male with purpura fulminans. The strain belonged to serotype 19A and ST320. Strain MTY1662SN214 was multidrug resistant, with resistance to penicillin, erythromycin, trimethoprim-sulfamethoxazole, tetracycline, quinolone, clindamycin, and cefotaxime (9).

The genome sequence was determined using Illumina MiSeq platform, which generated 487,316 total reads (483,492 reads after trimming) with an average length of  $250 \times 2$  nucleotides (nt), providing about 25-fold genome coverage. *De novo* assemblies were performed by the SPAdes Genome Assembler version 3.5.0 software (10). The unclosed draft genome of MTY1662SN214 was assembled into 60 contigs ( $\geq$ 500 bp), with a total length of 2,069,474 bp and a G+C content of 39.7%.

In order to obtain a functional annotation of the genome, all the assembled contigs were annotated by NCBI Prokaryotic Genome Annotation Pipeline (11). The genome possesses a total of 2,221 genes, including 2,073 coding sequences (CDSs), 5 rRNAs (three copies of 5S, one copy of 16S, and one copy of 23S rRNA genes), 47 tRNAs, and 95 pseudogenes.

**Nucleotide sequence accession numbers.** This whole-genome shotgun project has been deposited at DDBJ/EMBL/GenBank under the accession no. LKAA000000000. The version described in this paper is version LKAA01000000.

## **FUNDING INFORMATION**

This research received no specific grant from any funding agency in the public, commercial, or not-for-profit sectors.

## **REFERENCES**

- Whitney CG, Farley MM, Hadler J, Harrison LH, Bennett NM, Lynfield R, Reingold A, Cieslak PR, Pilishvili T, Jackson D, Facklam RR, Jorgensen JH, Schuchat A, Active Bacterial Core Surveillance of the Emerging Infections Program Network. 2003. Decline in invasive pneumococcal disease after the introduction of protein-polysaccharide conjugate vaccine. N Engl J Med 348:1737–1746. http://dx.doi.org/10.1056/ NEIMoa022823.
- Del Grosso M, Camilli R, D'Ambrosio F, Petrucci G, Melchiorre S, Moschioni M, Gherardi G, Pantosti A. 2013. Increase of pneumococcal serotype 19A in Italy is due to expansion of the piliated clone ST416/ CC199. J Med Microbiol 62:1220–1225. http://dx.doi.org/10.1099/ imm.0.061242-0.
- Hsu HE, Shutt KA, Moore MR, Beall BW, Bennett NM, Craig AS, Farley MM, Jorgensen JH, Lexau CA, Petit S, Reingold A, Schaffner W, Thomas A, Whitney CG, Harrison LH. 2009. Effect of pneumococcal conjugate vaccine on pneumococcal meningitis. N Engl J Med 360: 244–256. http://dx.doi.org/10.1056/NEJMoa0800836.
- Hanage WP, Bishop CJ, Lee GM, Lipsitch M, Stevenson A, Rifas-Shiman SL, Pelton SI, Huang SS, Finkelstein JA. 2011. Clonal replacement among 19A Streptococcus pneumoniae in Massachusetts, prior to 13 valent conjugate vaccination. Vaccine 29:8877–8881. http://dx.doi.org/ 10.1016/j.vaccine.2011.09.075.
- Pillai DR, Shahinas D, Buzina A, Pollock RA, Lau R, Khairnar K, Wong A, Farrell DJ, Green K, McGeer A, Low DE. 2009. Genome-wide dissection of globally emergent multi-drug resistant serotype 19A Streptococcus pneumoniae. BMC Genomics 10:642. http://dx.doi.org/10.1186/1471 -2164-10-642.

- Ramos V, Parra EL, Duarte C, Moreno J. 2014. Characterization of Streptococcus pneumoniae invasive serotype 19A isolates recovered in Colombia. Vaccine 32:755-758. http://dx.doi.org/10.1016/ i.vaccine.2013.12.024.
- Hsieh YC, Lin TL, Chang KY, Huang YC, Chen CJ, Lin TY, Wang JT. 2013. Expansion and evolution of *Streptococcus pneumoniae* serotype 19A ST320 clone as compared to its ancestral clone, Taiwan19F-14 (ST236). J Infect Dis 208:203–210. http://dx.doi.org/10.1093/infdis/jit145.
- Ip M, Ang I, Liyanapathirana V, Ma H, Lai R. 2015. Genetic analyses of penicillin binding protein determinants in multidrug-resistant *Streptococ-cus pneumoniae* serogroup 19 CC320/271 clone with high-level resistance to third-generation cephalosporins. Antimicrob Agents Chemother 59: 4040–4045. http://dx.doi.org/10.1128/AAC.00094-15.
- 9. CLSI. 2012. Performance standards for antimicrobial susceptibility

- testing: 22nd informational supplement. CLSI document M100-S22. Clinical and Laboratory Standards Institute, Wayne, Pennsylvania, USA.
- Bankevich A, Nurk S, Antipov D, Gurevich AA, Dvorkin M, Kulikov AS, Lesin VM, Nikolenko SI, Pham S, Prjibelski AD, Pyshkin AV, Sirotkin AV, Vyahhi N, Tesler G, Alekseyev MA, Pevzner PA. 2012. SPAdes: a new genome assembly algorithm and its applications to single-cell sequencing. J Comput Biol 19:455–477. http://dx.doi.org/10.1089/cmb.2012.0021.
- Angiuoli SV, Gussman A, Klimke W, Cochrane G, Field D, Garrity GM, Kodira CD, Kyrpides N, Madupu R, Markowitz V, Tatusova T, Thomson N, White O. 2008. Toward an online repository of Standard Operating Procedures (SOPs) for (meta)genomic annotation. OMICS 12: 137–141. http://dx.doi.org/10.1089/omi.2008.0017.