



Draft Genome Sequence of *Escherichia albertii* Strain Mex-12/320a, Isolated from an Infant with Diarrhea and Harboring Virulence Genes Associated with Diarrheagenic Strains of Enteropathogenic *Escherichia coli*

Samantha Maldonado-Puga,^a Mario Meza-Segura,^a Adriana Becerra,^a Mussaret B. Zaidi,^{b,c} Teresa Estrada-Garcia^a

^aDepartment of Molecular Biomedicine, CINVESTAV-IPN, Mexico City, Mexico

^bInfectious Diseases Research Unit, Hospital General O'Horan, Merida, Mexico

^cDepartment of Epidemiology and Biostatistics, Michigan State University, Lansing, Michigan, USA

ABSTRACT *Escherichia albertii* is an emerging human enteropathogen. We report the draft genome sequence of *E. albertii* strain Mex-12/320a, isolated from an infant with diarrhea. The presence of the pathogenic island O122/IE6 and the *nleA* gene, previously found in diarrheagenic enteropathogenic *Escherichia coli* strains, suggests that *E. albertii* may cause acute diarrhea.

Escherichia albertii is an emerging human enteropathogen which belongs to the attaching and effacing (A/E) family of gastrointestinal pathogens that include well-known human pathogens such as enteropathogenic *Escherichia coli* (EPEC) and enterohemorrhagic *E. coli* (1). The genes responsible for the A/E phenotype are carried on a pathogenicity island (PAI) known as the locus of enterocyte effacement (LEE), which encodes several effector proteins that are all injected into the host cell by the type III secretion system (T3SS) (2). Other non-LEE-encoded (NLE) EPEC effectors are translocated by the T3SS (1). EPEC non-LEE genes, such as *nleA* and those harbored in the PAI O122/IE6 (*efa1* or *lifA*, *espL*, *nleB*, and *nleE*), are significantly more prevalent among EPEC strains isolated from children with diarrhea than in those without (3–5). Most of these genes are involved in bacterial evasion of the host innate immune response (6, 7).

We report the draft genome sequence of *E. albertii* strain Mex-12/320a, a lactose-negative *E. coli* strain that was isolated from the stool of a 2-year-old girl with acute diarrhea. She was admitted to a hospital in Yucatan, Mexico, in 2012 for dehydration after 4 days of illness with 4 stool movements per day and 12 episodes of vomiting. This strain was isolated and characterized as previously described (8), was originally classified as atypical EPEC, and was the sole pathogenic isolate identified in the fecal sample. It was subsequently reclassified as *E. albertii* by a multiplex *in silico* PCR that discriminates between different *Escherichia* species using primers for the DNA-binding transcriptional activator of a cysteine biosynthesis gene (393-bp product) that appears to be specific to this species (9).

One colony from MacConkey agar was selected and incubated in LB broth for 18 h at 37°C without agitation. Genomic DNA was extracted using a phenol-chloroform protocol, prepared as Illumina sequencing libraries using the Nextera XT kit, and sequenced by the GAllx system (Illumina, San Diego, CA, USA), generating 3,148,027 reads. Quality control, trimming, and filtering of raw sequencing data were performed with Trim Galore v0.4.1 (https://www.bioinformatics.babraham.ac.uk/projects/trim_galore/). The genome was assembled *de novo* using the SPAdes genome assembler v3.9.1 (<http://cab.spbu.ru/software/spades/>), and the accuracy of the genome assemblies was

Citation Maldonado-Puga S, Meza-Segura M, Becerra A, Zaidi MB, Estrada-Garcia T. 2019. Draft genome sequence of *Escherichia albertii* strain Mex-12/320a, isolated from an infant with diarrhea and harboring virulence genes associated with diarrheagenic strains of enteropathogenic *Escherichia coli*. Microbiol Resour Announc 8:e00208-19. <https://doi.org/10.1128/MRA.00208-19>.

Editor David Rasko, University of Maryland School of Medicine

Copyright © 2019 Maldonado-Puga et al. This is an open-access article distributed under the terms of the [Creative Commons Attribution 4.0 International license](https://creativecommons.org/licenses/by/4.0/).

Address correspondence to Teresa Estrada-Garcia, testrada@cinvestav.mx.

Received 5 March 2019

Accepted 4 June 2019

Published 3 July 2019

evaluated with REAPR 1.0.16 (10). Protein-coding sequences and RNA genes were predicted, and functional annotation was performed with Prokka v1.11. Preassembled contigs were scaffolded using SSPACE_Standard_v3.0.pl (11), and gaps were closed with GapFiller_v1-10 (12). Default parameters were used for all software during the assembly and annotation process.

The draft chromosome sequence of *E. albertii* strain Mex-12/320a consists of 4,941,694 bp in 268 scaffolds, with an N_{50} value 130,167 bp, a maximum scaffold size of 341 kbp, and 49.54% G+C content. The genome contains 5,090 predicted genes, of which 4,763 are coding sequences (CDSs), 5 are rRNA (*rrn*) operons, and 79 are tRNA genes. The genome sequence confirmed the presence of not only the LEE genes but also *nleA* and *cdt-II* (cytolethal distending toxin II) virulence genes previously found in other *E. albertii* strains (13, 14). To the best of our knowledge, this is the first report of the presence of the complete PAI O122/IE6 (*efa1* or *lifa*, *espL*, *nleB*, and *nleE* genes) in an *E. albertii* strain.

This work confirms that the *E. albertii* Mex-12/320a genome harbors virulence genes that have been previously reported in EPEC isolates from children with diarrhea. This finding adds to the growing body of evidence that *E. albertii* is an emerging pathogen of diarrheal disease (15) and that *nleA* and the PAI O122/IE6 are virulence markers of A/E *Escherichia* pathogens. This report, in conjunction with those for other genomes of *E. albertii* (13, 16, 17), will contribute to a greater understanding of the phylogeny and evolution of *Escherichia* species and to the identification of new virulence factors among A/E bacteria.

Data availability. The draft genome assembly of *E. albertii* strain Mex-12/320a was deposited in DDBJ/ENA/GenBank under the accession number [SIZV00000000](https://www.ncbi.nlm.nih.gov/nuclink/SIZV00000000) and BioProject number [PRJNA523447](https://www.ncbi.nlm.nih.gov/bioproject/PRJNA523447). Sequencing reads for *E. albertii* strain Mex-12/320a have been deposited in the NCBI Sequence Read Archive (SRA) under the accession number [SRR9024355](https://www.ncbi.nlm.nih.gov/sra/SRR9024355).

ACKNOWLEDGMENTS

This work was supported by CONACYT scholarships to S.M.-P. (award 261205), M.M.-S. (award 368026), and A.B. (award 483617) and a CONACYT grant (254994) to T.E.-G.

REFERENCES

- Slater SL, Sãgfors AM, Pollard DJ, Ruano-Gallego D, Frankel G. 2018. The type III secretion system of pathogenic *Escherichia coli*. *Curr Top Microbiol Immunol* 416:51–72. https://doi.org/10.1007/82_2018_116.
- Gaytán MO, Martínez-Santos VI, Soto E, González-Pedrajo B. 2016. Type three secretion system in attaching and effacing pathogens. *Front Cell Infect Microbiol* 6:129. <https://doi.org/10.3389/fcimb.2016.00129>.
- Afset J, Bruant G, Brousseau R, Harel JJ, Anderssen E, Bevanger L, Bergh KK. 2006. Identification of virulence genes linked with diarrhea due to atypical enteropathogenic *Escherichia coli* by DNA microarray analysis and PCR. *J Clin Microbiol* 44:3703–3711. <https://doi.org/10.1128/JCM.00429-06>.
- Mercado EH, Piscoche C, Contreras C, Durand D, Riveros M, Ruiz J, Ochoa TJ. 2016. Pathogenicity island O-122 in enteropathogenic *Escherichia coli* strains is associated with diarrhea severity in children from Lima Peru. *Int J Med Microbiol* 306:231–236. <https://doi.org/10.1016/j.ijmm.2016.05.005>.
- Salvador FA, Hernandez RT, Vieira MAM, Rockstroh AC, Gomes T. 2014. Distribution of non-LEE-encoded type 3 secretion system dependent effectors in enteropathogenic *Escherichia coli*. *Braz J Microbiol* 45: 851–855. <https://doi.org/10.1590/S1517-83822014000300014>.
- Santos AS, Finlay BB. 2015. Bringing down the host: enteropathogenic and enterohaemorrhagic *Escherichia coli* effector-mediated subversion of host innate immune pathways. *Cell Microbiol* 17:318–332. <https://doi.org/10.1111/cmi.12412>.
- Yen H, Sugimoto N, Tobe T. 2015. Enteropathogenic *Escherichia coli* uses NleA to inhibit NLRP3 inflammasome activation. *PLoS Pathog* 11: e1005121. <https://doi.org/10.1371/journal.ppat.1005121>.
- Patzi-Vargas S, Zaidi MB, Perez-Martinez I, León-Cen M, Michel-Ayala A, Chaussabel D, Estrada-Garcia T. 2015. Diarrheagenic *Escherichia coli* carrying supplementary virulence genes are an important cause of moderate to severe diarrhoeal disease in Mexico. *PLoS Negl Trop Dis* 9:e0003510. <https://doi.org/10.1371/journal.pntd.0003510>.
- Lindsey RL, Garcia-Toledo L, Fasulo D, Gladney LM, Strockbine N. 2017. Multiplex polymerase chain reaction for identification of *Escherichia coli*, *Escherichia albertii* and *Escherichia fergusonii*. *J Microbiol Methods* 140: 1–4. <https://doi.org/10.1016/j.mimet.2017.06.005>.
- Hunt M, Kikuchi T, Sanders M, Newbold C, Berriman M, Otto TD. 2013. REAPR: a universal tool for genome assembly evaluation. *Genome Biol* 14:R47. <https://doi.org/10.1186/gb-2013-14-5-r47>.
- Boetzer M, Pirovano W. 2012. Toward almost closed genomes with GapFiller. *Genome Biol* 13:R56. <https://doi.org/10.1186/gb-2012-13-6-r56>.
- Boetzer M, Henkel CV, Jansen HJ, Butler D, Pirovano W. 2011. Scaffolding pre-assembled contigs using SSPACE. *Bioinformatics* 27:578–579. <https://doi.org/10.1093/bioinformatics/btq683>.
- Ooka T, Ogura Y, Katsura K, Seto K, Kobayashi H, Kawano K, Tokuoka E, Furukawa M, Harada S, Yoshino S, Seto J, Ikeda T, Yamaguchi K, Murase K, Gotoh Y, Imuta M, Nishi J, Gomes TA, Beutin L, Hayashi T. 2015. Defining the genome features of *Escherichia albertii*, an emerging enteropathogen closely related to *Escherichia coli*. *Genome Biol Evol* 7:3170–3179. <https://doi.org/10.1093/gbe/ew211>.
- Hinenoya A, Yasuda N, Hibino T, Shima A, Nagita A, Tsukamoto T, Yamasaki S. 2017. Isolation and characterization of an *Escherichia albertii* strain pro-

- ducing three different toxins from a child with diarrhea. *Jpn J Infect Dis* 70:252–257. <https://doi.org/10.7883/yoken.JJID.2016.186>.
15. Huys G, Cnockaert M, Janda JM, Swings J. 2003. *Escherichia albertii* sp. nov., a diarrhoeagenic species isolated from stool specimens of Bangladeshi children. *Int J Syst Evol Microbiol* 53:807–810. <https://doi.org/10.1099/ijs.0.02475-0>.
 16. Murakami K, Etoh Y, Tanaka E, Ichihara S, Horikawa K, Kawano K, Ooka T, Kawamura Y, Ito K. 2014. Shiga toxin 2f-producing *Escherichia albertii* from a symptomatic human. *Jpn J Infect Dis* 67:204–208. <https://doi.org/10.7883/yoken.67.204>.
 17. Romão FT, Hernandes RT, Ooka T, Hayashi T, Sperandio V, Gomes TAT. 2018. Complete genome sequence of *Escherichia albertii* strain 1551-2, a potential extracellular and intracellular pathogen. *Genome Announc* 6:e00075–18. <https://doi.org/10.1128/genomeA.00075-18>.