

Complete Genome Sequence of Human Adenovirus Type 7 Associated with Fatal Infant Pneumonia

Liuying Tang, a Junjing An, b Pengbo Yu, c Wenbo Xua

National Institute for Viral Disease Control and Prevention, Chinese Centre for Disease Control and Prevention, Beijing, People's Republic of China^a; Fengtai District, Beijing Center for Disease Control and Prevention, Beijing, People's Republic of China^b; Shaanxi Center for Disease Control and Prevention, Xi'an, People's Republic of China^c

The Chinese human adenovirus 7 (HAdV7) 0901HZ/ShX/CHN/2009 was isolated from the hydrothorax fluid of an infant with fatal pneumonia in Shaanxi, China, in 2009. Comparison of the entire genome with the genomes of the other 10 strains of HAdV7 from GenBank revealed homologies of 89.9 to 99.9%, with geographic polymorphism among HAdV-7 field strains circulating in mainland China.

Received 6 December 2012 Accepted 18 December 2012 Published 14 February 2013

Citation Tang L, An J, Yu P, Xu W. 2013. Complete genome sequence of human adenovirus type 7 associated with fatal infant pneumonia. Genome Announc. 1(1):e00182-12. doi:10.1128/genomeA.00182-12.

Copyright © 2013 Tang et al. This is an open-access article distributed under the terms of the Creative Commons Attribution 3.0 Unported license. Address correspondence to Liuying Tang, tangly@ivdc.chinacdc.cn.

uman adenoviruses (HAdVs) can cause a broad range of clinical manifestations in humans, with billions of people infected worldwide (1). These adenoviruses are grouped into seven species (A to G), and to date fifty-six types (HAdV-1 to HAdV-56) have been identified on the basis of nucleotide and deduced amino acid sequences (2, 3). The increasing application of genomics and bioinformatics approaches to the investigation of HAdV genomes will lead to an understanding of the emergence of novel HAdV pathogens, the identification and performance of surveillance for these pathogens, and perhaps the control of epidemic outbreaks with effective and appropriate vaccines (4).

An outbreak of infant severe pneumonia was reported in Shaanxi, China, in 2009 (5), and two adenovirus 7 strains were isolated from the hydrothorax fluid of the patients. For one strain, named 0901HZ/ShX/CHN/2009 (HAdV7-0901HZ), sequencing and analysis of the whole genome were conducted. For genome sequencing, a standard PCR was conducted by using a previously described procedure (6). Raw sequence data were assembled using Sequencher software (version 4.0.5).

Pipmaker (http://bio.cse.psu.edu/pipmaker) (7) was used to perform nucleotide dot blot identity-based analyses of the whole HAdV genomes now available from GenBank. A total of 10 complete genome sequences of HAdV7 (not including the one in this study) are available in the GenBank database, including the prototype type, vaccine strains, and field strains, of which three were isolated from the south of China, Guangzhou, and Chongqing. The strain 0901HZ was isolated from the northwest of China, in the Shaanxi Province.

The strain HAdV7-0901HZ has a genome of 35,239 bp in length and a GC content of 51.08%. A total of 48 coding sequences were identified in this genome sequence, including seven predicted genes.

The genome sequence alignment of HAdV7-0901HZ with other HAdV7 strains revealed high homologies of 99.7 to 99.9% (accession numbers JX625134, JN860677, and GQ478341). An exception was HAdV7-GZ07, which has a large fragment deletion at position 28334 to 31162 of the E3 region (accession number

HQ659699) and homology of 89.9%. For the vaccine and Gomen strains (accession nos. AY594255 and AY594256, respectively) the nucleotide sequence similarities were 97.7% and 99.4%, respectively

Adenovirus infections in the immunocompetent host are usually limited by the immune system. But HAdV7, a member of the B1 subspecies, has been identified in epidemics to be highly virulent and is associated with clinical manifestations of considerable severity, including residual lung damage and fatal outcomes (8–10). This pathogen has been found in outbreaks in the United States and Asia and has become a global concern (11–13). It also poses a special threat for infants and young children as their immune systems are not completely established.

Comparative genomic analysis of 11 strains of human adenovirus type 7 isolated from this study and the references from GenBank showed that HAdV-7 circulating in mainland China has multiple transmission chains, suggesting the need for surveillance in high-risk populations.

Nucleotide sequence accession number. The nucleotide sequence of the complete genome of HAdV7 strain 0901HZ/ShX/CHN/2009 has been submitted to GenBank under accession number JF800905.

ACKNOWLEDGMENT

This study was supported by the "China Mega-Project for Infectious Disease (2011ZX10004-001)" from The People's Republic of China.

REFERENCES

- Smith JG, Wiethoff CM, Stewart PL, Nemerow GR. 2010. Adenovirus. Curr. Top. Microbiol. Immunol. 343:195–224.
- 2. Robinson CM, Singh G, Henquell C, Walsh MP, Peigue-Lafeuille H, Seto D, Jones MS, Dyer DW, Chodosh J. 2011. Computational analysis and identification of an emergent human adenovirus pathogen implicated in a respiratory fatality. Virology 409:141–147.
- 3. Walsh MP, Seto J, Jones MS, Chodosh J, Xu W, Seto D. 2010. Computational analysis identifies human adenovirus type 55 as a re-emergent acute respiratory disease pathogen. J. Clin. Microbiol. 48:991–993.
- 4. Mahadevan P, Seto J, Tibbetts C, Seto D. 2010. Natural variants of

- human adenovirus type 3 provide evidence for relative genome stability across time and geographic space. Virology 397:113–118.
- Tang L, Wang L, Tan X, Xu W. 2011. Adenovirus serotype 7 associated with a severe lower respiratory tract disease outbreak in infants in Shaanxi Province, China. Virol. J. 8:23.
- Yang Z, Zhu Z, Tang L, Wang L, Tan X, Yu P, Zhang Y, Tian X, Wang J, Zhang Y, Li D, Xu W. 2009. Genomic analyses of recombinant adenovirus type 11a in China. J. Clin. Microbiol. 47:3082–3090.
- Schwartz S, Zhang Z, Frazer KA, Smit A, Riemer C, Bouck J, Gibbs R, Hardison R, Miller W. 2000. PipMaker: a web server for aligning two genomic DNA sequences. Genome Res. 10:577–586.
- Kajon AE, Dickson LM, Fisher BT, Hodinka RL. 2011. Fatal disseminated adenovirus infection in a young adult with systemic lupus erythematosus. J. Clin. Virol. 50:80–83.
- Selvaraju SB, Kovac M, Dickson LM, Kajon AE, Selvarangan R. 2011. Molecular epidemiology and clinical presentation of human adenovirus infections in Kansas City children. J. Clin. Virol. 51:126–131.
- Stercz B, Nagy K, Ongrádi J. 2012. Adenovirus infections in immuncompromised patients. Orv. Hetil. 153:1896–1904.
- 11. Azar R, Varsano N, Mileguir F, Mendelson E. 1998. Molecular epidemiology of adenovirus type 7 in Israel: identification of two new genome types, Ad7k and Ad7d2. J. Med. Virol. 54:291–299.
- 12. Gerber SI, Erdman DD, Pur SL, Diaz PS, Segreti J, Kajon AE, Belkengren RP, Jones RC. 2001. Outbreak of adenovirus genome type 7d2 infection in a pediatric chronic-care facility and tertiary-care hospital. Clin. Infect. Dis. 32:694–700.
- 13. Ikeda Y, Yamaoka K, Noda M, Ogino T. 2003. Genome types of adenovirus type 7 isolated in Hiroshima city. J. Med. Virol. 69:215–219.