

Elsevier has created a Monkeypox Information Center in response to the declared public health emergency of international concern, with free information in English on the monkeypox virus. The Monkeypox Information Center is hosted on Elsevier Connect, the company's public news and information website.

Elsevier hereby grants permission to make all its monkeypox related research that is available on the Monkeypox Information Center - including this research content - immediately available in publicly funded repositories, with rights for unrestricted research re-use and analyses in any form or by any means with acknowledgement of the original source.

These permissions are granted for free by Elsevier for as long as the Monkeypox Information Center remains active.



Genomic characterisation of human monkeypox virus in Nigeria

Published Online
January 16, 2018
http://dx.doi.org/10.1016/
\$1473-3099(18)30043-4

This online publication has been corrected. The corrected version first appeared at thelancet. com/infection on February 21, 2018

See Online for appendix

Monkeypox virus (MPXV) is a large, double-stranded DNA virus belonging to the Orthopox genus in the family Poxviridae. First identified in 1958, MPXV has caused sporadic human outbreaks in central and west Africa, with a mortality rate between 1% and 10%. Viral genomes from west Africa and the Congo Basin separate into two clades, the latter being more virulent. Recently, MPXV outbreaks have occurred in Sudan (2005), the Republic of the Congo and Democratic Republic of the Congo (2009), and the Central African Republic (2016).

A suspected outbreak of human MPXV was reported to WHO on Sept 26, 2017, by the Nigeria Centre for Disease Control (NCDC) after a cluster of suspected cases had occurred in Yenagoa Local Government Area, Bayelsa State, Nigeria.⁴ Since the onset of the outbreak, 155 cases have been reported by the NCDC, of which 56 were confirmed.⁴ A subset of these samples was sent to the WHO Collaborating Center at the Institut Pasteur de Dakar (IPD) in Senegal for confirmation by PCR.

This was not the first report of MPXV cases in Nigeria. Between 1971 and 1978, ten human cases were reported, with three confirmed and two sequenced. Since then, no other cases have been reported in the area. Because of the lapse in MPXV cases in the region and recent cases in the Congo Basin, the origin of the Nigerian outbreak needed to be identified, as did whether the outbreak was a result of a local zoonotic spillover event or importation from another endemic country.

To help support advanced characterisation of pathogens in the region, 29 samples positive according to PCR were prepared for sequencing at IPD and enriched

using a pan-viral probe set designed by Illumina (San Diego, CA, USA) and the US Army Medical Research Institute of Infectious Diseases Center for Genomic Sciences (Frederick, MD, USA), and synthesised by Twist Bioscience (San Francisco, CA, USA). Analysis of the sequencing data found reads aligning to MPXV in 22 of the 29 samples. Three of the samples (297 957, 298 464, and 298481) were determined to have reads covering 99.8%, 94.2%, and 95.0% of the closest related MPXV genome sequence on GenBank (a Nigeria strain, KJ642617) with average depths of 512, 56, and 25 x, respectively. Reads aligned to the repetitive terminal ends but could not be used to definitely resolve the termini using the short Illumina reads.

Phylogenetic analysis (appendix) indicates that the closest relative of the three outbreak isolates were the two Nigerian strains available on GenBank (KJ642615 and KJ642617), within the west African clade. The isolates grouped most closely to KJ642617, a genome isolated from a human MPXV case in Ihie, Abia State, Nigeria, in 1971, which is relatively close to the epicentre of the current outbreak.

These findings support the hypothesis that the index case of the current outbreak in Nigeria was not imported, but probably originated from a spillover event or events involving reservoir hosts. These results emphasise the value of local surveillance for the early detection of viral spillovers and the need for advanced characterisation to help determine the origins of outbreaks.

MRW reports grants from the Defense Threat Reduction Agency and Joint Program Executive Office for Chemical and Biological Defense. All other authors declare no competing interests. The content of this publication does not necessarily reflect the views or policies of the US Army, the US Department of Defense, the US Department of Health and Human Services, or the institutions or companies affiliated with the authors. The research described herein was supported by the Institute Pasteur in Dakar, Senegal, and the US Defense Threat Reduction Agency (CB10246) and supported with equipment provided by the Targeted Acquisition of Reference Materials Augmenting

Capabilities (TARMAC) initiative and the Defense Biological Product Assurance Office (DBPAO) through a task order award to the National Strategic Research Institute, FA4600-12-D-9000. All the outbreak control teams were made up of staff from the Bayelsa State Ministry of Health, Niger Delta University Teaching Hospital, Nigeria Field and Laboratory Training Programme, and the Nigeria Centre for Disease Control. OA, CBP, and MF contributed equally to this research; GP, AAS, and CI are joint senior authors.

Ousmane Faye, Catherine B Pratt,
Martin Faye, Gamou Fall,
Joseph A Chitty, Moussa M Diagne,
Michael R Wiley,
Adesola F Yinka-Ogunleye, Sola Aruna,
Ebitimitula N Etebu, Neni Aworabhi,
Dimie Ogoina, Wari Numbere,
Nwando Mba, *Gustavo Palacios,
Amadou A Sall, Chikwe Ihekweazu
qustavo.f.palacios.ctr@mail.mil

Institute Pasteur, Dakar, Senegal (OF, MF, GF, MMD, AAS); Center for Genome Sciences, US Army Medical Research Institute of Infectious Disease, Frederick, MD 21702, USA (CBP, JAC, MRW, GP); College of Public Health, University of Nebraska Medical Center, Omaha, NE, USA (CBP, MRW); Nigeria Centre for Disease Control, Abuja, Nigeria (AFY-O, SA, NM, CI); Measure Evaluation, Abuja, Nigeria (SA); Bayelsa State Ministry of Health, Yenagoa, Nigeria (ENE, NA); Niger Delta University Teaching Hospital/Niger Delta University, Yenagoa, Nigeria (DO); and Nigeria Field Epidemiology and Laboratory Training Program, Ibadan, Nigeria (WN)

- Damon IK. Status of human monkeypox: clinical disease, epidemiology and research. Vaccine 2011; 29 (suppl): D54–59.
- 2 Likos AM, Sammons SA, Olson VA, et al. A tale of two clades: monkeypox viruses. J Gen Virol 2005; 86: 2661–72.
- 3 WHO. Media centre: monkeypox. Fact sheet, November, 2016. http://www.who.int/ mediacentre/factsheets/fs161/en/ (accessed Dec 7, 2017).
- 4 Nigeria Centre for Disease Control. Monkeypox outbreak in Nigeria: situation report (no. 009). Nov 23, 2017. https://reliefweb.int/sites/reliefweb.int/files/resources/An%20Update%20of%20 Monkeypox%20Outbreak%20in%20 Nigeria_231117_48.pdf (accessed Dec 7, 2017).

Beyond one virus: vaccination against hepatitis B after hepatitis C treatment

New treatments for hepatitis C virus (HCV) infection with direct-acting antivirals provide an extraordinary cure rate. A recent Article by