First detection of highly pathogenic H5N6 avian influenza virus on the African continent

Ismaila Shittu, Alice Bianco, Dorcas Gado, Nicodemus Mkpuma, Lanre Sulaiman, Agnes Laleye, Federica Gobbo, Alessio Bortolami, Francesco Bonfante, Columba Vakuru, Clement Meseko, Alice Fusaro, David Shamaki, Olaniran Alabi, Calogero Terregino and Tony Joannis

ABSTRACT
Since 2013, highly pathogenic avian influenza (HPAI) subtype H5N6 (clade 2.3.4.4) has been reported in wild birds and poultry in Asia as well as in other parts of the globe. In Africa, information on the presence of this virus subtype is lacking. This study reports the first detection of a HPAI (H5N6) virus (clade 2.3.4.4b) in a duck from a live bird market in Nigeria, whose genome is closely related to the European 2017–2018 H5N6 viruses, indiricting a recent virus introduction into the African continent.

ARTICLE HISTORY Received 6 February 2020; Revised 13 April 2020; Accepted 14 April 2020

KEYWORDS Highly pathogenic avian influenza virus; H5N6; live bird market; Nigeria; duck

In January 2006, the first outbreak of a highly pathogenic avian influenza (HPAI) subtype H5N1 (clade 2.2) was detected in Nigeria [1]. Within two years from its introduction, the disease had spread to 67.6% of the states in the country in circumstances suspected to be linked to wild bird migration and/or trade [2]. The HPAI outbreaks resulted in colossal economic losses and public health concerns. Following the first epidemic wave of HPAI (2006–2007), in 2008, a distinct H5N1 virus (clade 2.2.1) was detected in ducks from live bird market (LBM) surveillance in Northeast (Gombe) Nigeria [2]. In January 2015, another HPAI H5N1 incursion (clade 2.3.2.1.c) was reported from a LBM and from poultry farms in Lagos and Kano states, respectively [3]. A year later, in November 2016, a HPAI H5N8 virus of clade 2.3.4.4.b, responsible for one of the most devastating epizootic in poultry and wild birds in Europe [4], was detected in the northern state of Kano [5]. Since late-2017, a new reassortant HPAI clade 2.3.4.4b H5N6 has been reported in wild and domestic birds in Northern Europe. This study describes the first detection, in June 2019, of the same H5N6 subtype in a duck from LBM surveillance in Northwest (Sokoto) Nigeria (Figure 1).

During a recent surveillance activity for HPAI conducted from late-June to mid-August 2019 by the Department of Veterinary and Pest Control Services of the Federal Ministry of Agriculture and Rural Development, Abuja, an isolate of avian influenza was identified and characterized. In the course of the surveillance activity, 3131 tracheal and cloacal samples collected from 13 bird species were tested at the NVRI using real-time RT–PCR for avian influenza virus (AIV) targeting the matrix (M) gene [6]. AIV M-gene positive samples were subjected to specific protocols for subtyping avian influenza viruses [7,8]. In addition, the M-gene positive samples were inoculated in 9–11-day-old embryonated chicken eggs of specific antibody-negative origin according to the standard procedure [9].

A H5N6 virus, designated A/duck/Nigeria/SK28T_19VIR8424-2/2019, was isolated and sent to the Istituto Zooprofilattico Sperimentale delle Venezie (IZSVe), Padua, Italy, for subtype confirmation. The intravenous pathogenicity index (IVPI) was conducted to assess the pathogenicity of the virus (supplementary data). To trace the virus origin and evaluate its genetic properties, whole-genome sequencing was performed on the isolate using an Illumina MiSeq platform (Illumina, San Diego, CA, USA) (supplementary data). The sequences were deposited at the GenBank under the accession numbers: MN889503-MN889510.

The maximum likelihood phylogenetic tree of the haemagglutinin (HA) gene segment obtained by using IQTREE (Figure 1) shows that the HPAI H5N6 virus detected in Nigeria in summer 2019 falls within...
genetic clade 2.3.4.4b [10] and groups together with European H5N6 viruses identified in wild and domestic birds in 2017–2018 (similarity range between 98.9% and 99%) (Supplementary Table 1). The phylogenetic trees of all other gene segments reflect the same topology as the HA phylogeny (data not shown).

The amino acid sequences of the haemagglutinin (HA) gene show that the H5N6 virus possesses a multi-basic cleavage site (PLREKRRKR*GLF) typical of HPAI viruses. Similarly, the IVPI test result, 2.89, confirmed the highly pathogenic nature of the isolate [9]. In the neuraminidase (NA) gene segment of the virus, mutation N403H removes a potential N-glycosylation site at position 403, at the level of the sialic acid-binding domain of the protein [11]. This mutation could potentially affect both the antigenic and the receptor binding properties of this strain. Nevertheless, haemagglutination inhibition assays conducted with ferret antisera generated against HPAI viruses belonging to the 2.3.4.4 clade, H5N6 A/Sichuan/26221/2014 (SICH-26221), H5N8 A/Fujian-Sanyuan/21099/2017XPR8 (FUJIAN-21099) and the recent 2.3.4.4b H5N8 A/turkey/Italy/17VIR576-11/2017 (ITALY-576) strains, recorded titers that were either identical to the ones observed against the homologous antigens or within 1 log₂ difference. Genetic and antigenic data were generated and shared within the OFFLU network to contribute to the WHO biannual report of “Antigenic and genetic characteristics of zoonotic influenza viruses and development of candidate vaccine viruses for pandemic preparedness.”

Here, the first H5N6 clade 2.3.4.4.b introduction into Africa from Europe, most likely via wild birds is reported. HPAI H5N6 bearing the same gene constellation of the Nigerian strain, had already caused a total of 98 outbreaks in Europe between late-2017 and early 2019, of which 92% had occurred in wild birds [12]. This finding confirms our previous study, which recognized West Africa, and in particular Nigeria, as one of the most important hotspot for Gs/GD/96 HPAI H5Nx introduction into Africa [5]. Europe and West Africa are naturally connected by the Black Sea/Mediterranean flyway and, as suggested for previous virus spreads [5], wild migratory birds may have played an important role also in this new virus incursion. The virus may have circulated in the West African wild or domestic population for several months before its detection, highlighting the need to enhance surveillance, in particular in the areas close to wetlands. To prevent the spread of infection to other countries in the sub-region, coordinated control strategies have been applied, which include, but are not limited to, controlling the movement of poultry and poultry products with neighbouring countries. In addition, implementation of surveillance plans is mandatory to avoid possible gaps in monitoring the virus evolution and the epidemiological scenario.
Co-circulation of multiple subtypes (H5N8, H5N6 and H9N2) (T. Joannis pers. comm.) in the country may represent a diagnostic challenge, considering that new reassortant viruses could emerge, as recently reported in Egypt [13,14], and spread. Urgent actions to strengthen surveillance efforts, combined with eradication measures, adequate compensations and education of farmers are needed to contain and monitor virus spread and the emergence of novel viruses of animal and public health concern.

Acknowledgements

The authors appreciate the excellent technical assistance of Kayode Olawuyi, Chinonyerem Chinyere, Daspan Amos, Rachel Ogar, Lohbyen Amos, Judith Bakam, Seyi Oyetunde, Ruth Akintola, and Jesse Jonathan. The field officers from the Department of Veterinary and Pest Control Services of the Federal Ministry of Agriculture and Rural Development, Abuja Nigeria involved in sample collection are appreciated. We acknowledge the World Organisation for Animal Health (OIE) for supporting the Twinning Project “Improving NVRI laboratory capacity for a better control of the Avian Influenza virus at National and Regional level” between the Istituto Zooprofylattico Sperimentale delle Venezie (IZSVe) and the National Veterinary Research Institute (NVRI). We thank Francesca Ellero for editing the manuscript. We acknowledge the authors, originating and submitting laboratories for the sequences from GISAID’s EpiFlu™ Database on which this research is based.

Disclosure statement

No potential conflict of interest was reported by the author(s).

References

[1] Joannis T, Lombin LH, De Benedictis P, et al. Confirmation of H5N1 avian influenza in Africa. Vet Rec. 2006;158:309–310. doi:10.1136/vr.158.9.309-b.
[2] Fusaro A, Joannis T, Monnie I, et al. Introduction into Nigeria of a distinct genotype of avian influenza virus (H5N1). Emerg Infect Dis. 2009;15:445–447. doi:10.1007/s13337-012-0073-4.
[3] Monnie I, Meseko C, Joannis T, et al. Highly pathogenic avian influenza A(H5N1) virus in poultry, Nigeria, 2015. Emerg Infect Dis. 2015;21:1275–1277. doi:10.3201/eid2107.150421.
[4] Alarcon P, Brouwer A, Venkatesh D, et al. Comparison of 2016–17 and previous Epizootics of highly pathogenic avian influenza H5 Guangdong lineage in Europe. Emerg Infect Dis 2018; 24: 2270–2283. doi:10.3201/eid2412.171860
[5] Fusaro A, Zecchin B, Vranckx B, et al. Disentangling the role of Africa in the global spread of H5 highly pathogenic avian influenza. Nat Commun. 2019;10:531. doi:10.1038/s41467-019-13287-y.
[6] Spackman E, Senne DA, Myers TJ, et al. Development of a real-time reverse transcriptase PCR assay for type A influenza virus and the avian H5 and H7 hemagglutinin subtypes. J Clin Microbiol. 2002;40:3256–3260. doi:10.1128/jcm.40.9.3256-3260.2002.
[7] Slomka MJ, Pavlidis T, Banks J, et al. Validated H5 Eurasian real-time reverse transcriptase-polymerase chain reaction and its application in H5N1 outbreaks in 2005-2006. Avian Dis. 2007;51:373–377. doi:10.1637/77664-060906R1.1.
[8] Hoffmann B, Hoffmann D, Henritzi D, et al. Riems influenza a typing array (RITA): an RT-qPCR-based low density array for subtyping avian and mammalian influenza a viruses. Sci Rep. 2016;6:27211. doi:10.1038/srep27211.
[9] World Organisation for Animal Health. Avian influenza. In: Alexander DJ, editor. Manual of diagnostic tests and vaccines for terrestrial animals. OIE: Geneva, 2008. p. 465–481. Available from: http://www.oie.int/eng/normes/mmanual/A_summary.htm.
[10] Lee D-H, Bahl J, Torchetti MK, et al. Highly pathogenic avian influenza viruses and generation of novel Reassortants, United States, 2014–2015. Emerg Infect Dis. 2016;22:1283–1285. doi:10.3201/eid2207.160048.
[11] McAuley JL, Gilbertson BP, Trikovic S, et al. Influenza virus neuraminidase structure and functions. Front Microbiol. 2019;10:39. doi:10.3389/fmicb.2019.00039.
[12] Adlhoch C, Kuiken T, Monnie I, et al. Avian influenza overview November 2018 – February 2019. EFSA J; 2019;17. doi:10.2901/j.efsa.2019.5664.
[13] Hagag NM, Erfan AM, El-Husseiny M, et al. Isolation of a novel reassortant highly pathogenic avian influenza (H5N2) virus in Egypt. Viruses. 2019;11:565. doi:10.3390/v11030565.
[14] Hassan KE, King J, El-Kady M, et al. Novel reassortant highly pathogenic avian influenza A(H5N2) virus in Broiler Chickens, Egypt. Emerg Infect Dis. 2020;26:129–133. doi:10.3201/eid2601.190570.