Highly pathogenic avian influenza A(H5N1) virus has circulated in poultry in Indonesia since 2003 (1,2). The phylogeny of A(H5N1) viruses detected during 2003–2011 indicated all genes descended from 1 ancestral virus with a clade 2.1 hemagglutinin (HA) introduced into Indonesia before 2003 (3). These viruses became enzootic and evolved into second-, third-, and fourth-order HA clades, leading to the recent dominance of clade 2.1.3.2 viruses (4). Outbreaks in poultry typically caused high mortality rates among gallinaceous birds, especially layer, broiler, and native chickens. The virus seemed less pathogenic in rates among gallinaceous birds, especially layer, broiler, and native chickens. The virus seemed less pathogenic in aquatic birds (5). However, reports of duck deaths and a higher than usual mortality rate (100% in some outbreaks) in backyard farms in Central Java, Jogjakarta, and East Java Provinces, Indonesia, in September 2012 triggered a joint outbreak investigation by animal and public health authorities (6). We describe the genetic characteristics of viruses isolated from A(H5N1) infection outbreaks in these 3 provinces on Java Island, where a previously unrecognized clade was detected.

The Study

We investigated 9 small-holding duck farms that reported bird deaths during September 12–November 5, 2012 (6). Cloacal swab samples were collected from sick birds, placed in 1,000 µL of viral transport medium, and sent for testing at laboratories of the regional Ministry of Agriculture Disease Investigation Center, Jogjakarta. Seventeen A(H5N1)–positive samples were forwarded to the National Animal Health Laboratory, Indonesian Research Center for Veterinary Science (IRCVS), for virus isolation and genome sequencing.

In addition, IRCVS collected 122 cloacal swab samples from birds and 58 environmental swab samples (from defeathering machines) at 5 live-bird markets (LBMs) in East Java Province during November 5–8, 2012. RNA extracted from farm and LBM specimens was tested for influenza A matrix gene to identify presumptive A(H5N1)–positive samples (7). Select positive samples were inoculated in 9–11-day-old embryonated, specific pathogen–free eggs. Allantoic fluid was harvested 36 h postinfection and tested for HA with chicken erythrocytes to confirm virus isolation (8).

Samples showing suspected A(H5N1) infection were propagated in a Biosafety Level-3 laboratory at IRCVS in compliance with biosafety regulations. Ten virus isolates (7 from duck farms, 3 from LBMs) were chosen for full-length HA gene sequencing (GenBank accession nos. KC417271–KC417277, KC757643); 4 were selected for genome sequencing. Results of reverse transcription PCR and sequencing primers are available on request. Sequencing and consensus sequence generation were conducted as described (9). Phylogenetic trees were generated by using MEGA4 (10) (Figure; online Technical Appendix 1, wwwnc.cdc.gov/EID/article/20/4/13-0517-Techapp1.pdf).

Phylogenetic analysis revealed that A(H5N1) isolates from samples collected from duck farm outbreaks and an LBM were not related to isolates in long-established Indonesian clade 2.1; rather, the HA genes closely resembled those of clade 2.3.2.1 viruses recently found in Vietnam, China, and Hong Kong (Figure). Full-length HA genes showed 97%–98%-nt identity with recent viruses from Vietnam and clustered in a larger group containing viruses from many Asian regions during 2009–2012. The environmental sample from an East Java LBM shared >99% nt similarity with viruses from samples at duck farms, indicating spread of this A(H5N1) clade into the marketing chain. A poultry sample from the same district as the 2.3.2.1 virus was identified as clade 2.1.3.2 (Figure), indicating likely cocirculation.

The 8 clade 2.3.2.1 HA genes analyzed possessed a multibasic amino acid cleavage site (Table 1). The cleavage site sequence of the clade 2.3.2.1 viruses from Indonesian (PQREdelRRRKR↓G) differed from recent clade 2.1.3.2...
viruses (PQRSRRKKG) by a Ser deletion at position 325 and a K328R substitution. Like other serotype H5N1 HA proteins, all isolates possessed a conserved glutamine at position 222 (equivalent to H3 position 226) and glycine at position 224 (H3 position 228), indicating no substantial changes in avian receptor-binding specificity (Table 1). The clade 2.3.2.1 viruses from Indonesia possessed 6 or 7 potential N-linked glycosylation sites (7 in clade 2.1.3.2 viruses), but unlike 2.1.3.2 viruses, all 2.3.2.1 viruses lacked the potential glycosylation site at position 154. Up to 29 conserved amino acid changes occurred in the mature HA1 protein between clade 2.3.2.1 and clade 2.1.3.2 viruses found recently in Indonesia, indicating these A(H5N1) virus subgroups probably diverged substantially.

**Table 1. Genetic characteristics of influenza A(H5N1) clade 2.3.2.1 viruses found in Indonesia, 2012**

<table>
<thead>
<tr>
<th>Strain name</th>
<th>PB2 aa</th>
<th>PB1-F2 truncation</th>
<th>HA1 aa</th>
<th>Cleavage site</th>
<th>NA aa</th>
<th>M2 aa</th>
<th>NS sequence</th>
</tr>
</thead>
<tbody>
<tr>
<td>A/Hubei/1/2010</td>
<td>627 D</td>
<td>701</td>
<td>Q G</td>
<td>PQERRRKRKG</td>
<td>I I S</td>
<td>Yes</td>
<td>ESEV</td>
</tr>
<tr>
<td>A/Hong Kong/6841/2010</td>
<td>E D</td>
<td>90 aa N</td>
<td>Q G</td>
<td>PQERRRKRKG</td>
<td>I I S</td>
<td>Yes</td>
<td>ESEV</td>
</tr>
<tr>
<td>A/env/East Java/LBM-LM13/2012</td>
<td>E D</td>
<td>57 aa N</td>
<td>Q G</td>
<td>PQERRRKRKG</td>
<td>V I S</td>
<td>Yes</td>
<td>ESEV</td>
</tr>
<tr>
<td>A/duck/Sukoharjo/BBVW-1428-9/2012</td>
<td>E D</td>
<td>57 aa N</td>
<td>Q G</td>
<td>PQERRRKRKG</td>
<td>I I S</td>
<td>Yes</td>
<td>ESEV</td>
</tr>
<tr>
<td>A/duck/Bantul/BBVW-1443-9/2012</td>
<td>E D</td>
<td>57 aa N</td>
<td>Q G</td>
<td>PQERRRKRKG</td>
<td>I I S</td>
<td>Yes</td>
<td>ESEV</td>
</tr>
<tr>
<td>A/duck/Slemang/BBVW-1463-10/2012</td>
<td>E D</td>
<td>57 aa N</td>
<td>Q G</td>
<td>PQERRRKRKG</td>
<td>ND I S</td>
<td>Yes</td>
<td>ESEV</td>
</tr>
<tr>
<td>A/moncorvo/BBVW-1732-11/2012</td>
<td>ND ND</td>
<td>ND ND</td>
<td>Q G</td>
<td>PQERRRKRKG</td>
<td>ND ND</td>
<td>ND ND</td>
<td>ND</td>
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<tr>
<td>A/dk/Bilgar/BBVW-1731-11/2012</td>
<td>ND ND</td>
<td>ND ND</td>
<td>Q G</td>
<td>PQERRRKRKG</td>
<td>ND ND</td>
<td>ND ND</td>
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<tr>
<td>A/dk/Tegal/BBVW-1727-11/2012</td>
<td>ND ND</td>
<td>ND ND</td>
<td>Q G</td>
<td>PQERRRKRKG</td>
<td>ND ND</td>
<td>ND ND</td>
<td>ND</td>
</tr>
<tr>
<td>A/dk/Wonogiri/BBVW-1730-11/2012</td>
<td>ND ND</td>
<td>ND ND</td>
<td>Q G</td>
<td>PQERRRKRKG</td>
<td>ND ND</td>
<td>ND ND</td>
<td>ND</td>
</tr>
</tbody>
</table>

*HA, hemagglutinin; NA, neuraminidase; M2, matrix 2; NS, nonstructural; ND, not determined.
†Numbering of the first and last nucleotide position of the gene that was sequenced is as follows: PB2, 1618–2192; PB1, 130–632; PA, 34–429; HA, 1–1710; NP, 268–755; NA, 55–1314; M, 55–919; NS, 25–690.
‡Glycosylation motif at 154 was absent for all strains.
in antigenicity. In contrast, the HA1 of the new viruses collected in Indonesia differed by 8–10 aa from A/Hubei/1/2010, the most closely related clade 2.3.2.1 A(H5N1) candidate vaccine virus recommended by the World Health Organization (online Technical Appendix 2, wwwnc.cdc.gov/EID/article/20/4/13-0517-Techapp2.pdf) (7).

To test the antigenic relationship of the clade 2.3.2.1 virus to the endemic clade 2.1.3.2 virus, we conducted a hemagglutination-inhibition test with ferret antiserum raised against viruses from these and other H5N1 clades (Table 2) (8). As the HA1 protein sequence differences suggest, clade 2.1.3.2 antiserum did not inhibit hemagglutination by a representative clade 2.3.2.1 virus from Indonesia, A/environment/East Java/LBM-LM13/2012. In contrast, this virus cross-reacted with antiserum to clade 2.3.2.1 viruses and the previously identified clade 2.1.3.2 genotype virus in Indonesia.

Conclusions
Detection of a novel clade of A(H5N1) virus in Indonesia marks a potential turning point in the molecular epidemiology of this virus. Indonesia has the highest number of human A(H5N1) infections because of ongoing outbreaks in poultry (14,15). Whether this new virus will become entrenched, as did clade 2.1.3 viruses over the past decade, remains to be seen, as do its effects on the incidence of human infection. Potential cocirculation of subtypes of 2 different clades warrants review of diagnostic methods and vaccination strategy to maximize effectiveness of disease control interventions. The lack of antigenic relatedness between the clade 2.3.2.1 and 2.1.3.2 viruses must be considered when evaluating A(H5N1) serologic diagnostic reagents used in Indonesia. This change also may have implications in selecting prepanemic candidate vaccine virus for the region. Furthermore,
poultry vaccines may need to be matched antigenically to circulating virus if clade 2.3.2.1 virus continues to circulate in Indonesia. Introduction of this virus is a stark reminder of the value of control measures to reduce the spread of subtype H5N1 and the need for enhanced surveillance of humans and poultry to monitor changes in its genetic and immunologic features.

Dr Dharmayanti is a researcher in the Virology Department at the Indonesian Research Center for Veterinary Science, Ministry of Agriculture. Her primary research interest is avian influenza.

References


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Genetic Characterization of Clade 2.3.2.1 Avian Influenza A(H5N1) Viruses, Indonesia, 2012

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Technical Appendix 1.
Technical Appendix 1, A–G. Phylogenetic tree of PB2, PB1, PA, NP, NA, M, NS (A = PB2; B = PB1; C = PA; D = NP; E = NA; F = M; G = NS). The phylogenetic tree was generated in MEGA version 4, using neighbor-joining analysis with 1000 bootstrap replicates using the Kimura-2 parameter model. Viruses characterized in this study are indicated with a bar showing samples collected in Indonesia from September to November 2012.