Identification of H2N3 influenza A viruses from swine in the United States

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Although viruses of each of the 16 influenza A HA subtypes are potential human pathogens, only viruses of the H1, H2, and H3 subtype are known to have been successfully established in humans. H2 influenza viruses have been absent from human circulation since 1968, and as such they pose a substantial human pandemic risk. In this report, we isolate and characterize genetically similar avian/swine virus reassortant H2N3 influenza A viruses isolated from diseased swine from two farms in the United States. These viruses contained leucine at position 226 of the H2 protein, which has been associated with increased binding affinity to the mammalian α2,6Gal-linked sialic acid virus receptor. Correspondingly, the H2N3 viruses were able to cause disease in experimentally infected swine and mice without prior adaptation. In addition, the swine H2N3 virus was infectious and highly transmissible in swine and ferrets. Taken together, these findings suggest that the H2N3 virus has undergone some adaptation to the mammalian host and that their spread should be very closely monitored.

The generation of a pandemic influenza virus first requires interspecies transmission, and the virus must then genetically adapt to the new host species (1, 2) via either point mutations (antigenic drift) or reassortment (antigenic shift); the latter is the exchange of gene segments between two different influenza viruses. Molecular epidemiology suggests that the 1918 Spanish flu pandemic was caused by a wholly avian H1N1 influenza virus that was introduced into humans (3). The 1957 (H2N2) and 1968 (H3N2) pandemic viruses were generated through genetic reassortment of human and avian strains acquiring the neuraminidase (NA) and/or HA and the polymerase basic 1 (PB1) gene from an avian virus and other genes from the previously circulating human virus (4). The reassortment may have occurred either in infected humans or in an intermediate host, e.g., swine or quail (5, 6), before human infection. Swine are referred to as a “mixing vessel” because of their susceptibility to both human and avian influenza viruses (7, 8). Therefore, reassortment of avian and mammalian influenza viruses in this intermediate host may produce new viruses that are transmissible to humans.

H2N2 influenza virus has not circulated in the human population for the past 40 years and is currently detected only in avian species (9–11). There are two distinct lineages of avian H2 influenza viruses. The Eurasian lineage is genetically more similar to human H2 viruses (12) than the American lineage. Nevertheless, some H2 viruses isolated from North American shorebirds carry HA of the Eurasian lineage, suggesting interregional transmission of the H2 gene (13). H2 subtypes are presently circulating in birds, especially migratory birds. Here we describe the isolation and characterization of H2N3 influenza A viruses from pigs with respiratory disease from two farms in the United States, a subtype not previously reported in swine. These H2N3 reassortant viruses contain genes derived from avian and swine influenza viruses. We also investigated the pathogenicity and transmissibility of the H2N3 isolates in different mammalian hosts. The H2N3 virus was able to replicate in pigs, mice, and ferrets and was transmitted among pigs and ferrets. Serologic evidence suggests that the virus continues to circulate in the affected swine production systems.

Results

Analysis of Clinical Samples. In September 2006, the influenza virus A/Swine/Missouri/4296424/2006 (Sw/4296424) was isolated from several 5- to 6-week-old pigs with multifocal bronchopneumonia at a multisourced commercial swine nursery. Lung lesions included moderate, subacute to chronic, purulent bronchopneumonia and interstitial pneumonia with bronchioliths and perilbronchiitis. Lung tissue was negative for porcine reproductive and respiratory syndrome virus (PRRSV), porcine circovirus type 2 (PCV2), and Mycoplasma hyopneumoniae but was positive for Streptococcus suis. Because of the characteristic influenza-like lesions and clinical signs of pneumonia, lung tissue homogenate was inoculated on Madin–Darby canine kidney (MDCK) cells. Cytopathic effects were detected on day 3 postinoculation (p.i.). The influenza virus nucleoprotein (NP) gene was detected in the infected cells by RT-PCR. The virus did not react with reference swine anti-sera (A/Sw/IA/1973 H1N1, A/Sw/TX/1998 H3N2, A/Sw/NC/2001 H1N1) in hemagglutination inhibition (HI) assays, and multiplex RT-PCR detected no H1N1 or H3N2 genes (14). The virus was submitted to the National Animal Disease Center (NADC) in February 2007 for subtyping and sequencing.

After the September isolate had been subtyped and sequenced (described below), a search of case records revealed that another “untypable” influenza isolate had been submitted in April 2006, A/Swine/Missouri/2124514/2006 (Sw/2124514) had been isolated from a 12-week-old pig with respiratory disease at another commercial grower–finisher swine farm. Lung lesions were histopathologically characteristic of swine influenza (severe, subacute inflammation of alveoli and bronchi with bronchiolar epithelial cell necrosis and metaplasia). The lung was negative for PRRSV, PCV2, and M. hyopneumoniae but was positive for influenza A virus by RT-PCR (specific for the NP gene) and S. suis. The virus was submitted to the NADC in March 2007 for subtyping and sequencing.
Subtyping and Phylogenetic Analysis. To identify and characterize both influenza viruses, nucleic acid sequencing and molecular and phylogenetic analysis were conducted. Both viruses were directly sequenced from low-passage isolates by using MDCK cells, and the sequences were confirmed after plaque purification and resequencing. They were identified as H2N3 viruses by nucleotide sequence and a BLAST search of the Influenza Sequence Database (www.flu.lanl.gov). The HA gene segment of Sw/4296424 most closely matched those of H2 viruses isolated from mallards in North America [up to 97.8% nucleotide sequence identity; supporting information (SI Table 4)]. Its PA segment was closely related to that of an H4N3 avian influenza virus (AIV) isolated from blue-winged teal (98.3% identity). With the exception of the polymerase acidic (PA) gene, its internal genes were derived from contemporary triple-reassortant swine influenza viruses currently found in the United States. These viruses carry internal genes from human (PB1), avian (PB2, PA), and swine [NP, matrix (M), nonstructural (NS)] influenza virus origin (SI Table 4). Its PA segment was 99.2% identical to that of the H6N5 AIV isolated from mallard ducks (SI Table 4). The Sw/2124514 and Sw/4296424 viruses showed 99.3–99.9% total nucleotide sequence identity (SI Table 5). Both isolates were repeatedly plaque cloned, retested, and confirmed by sequencing to belong to the H2N3 subtype. The H2N3 subtype was serologically confirmed by hemagglutination inhibition and neuraminidase inhibition assays. Phylogenetic analysis based on the HA and NA genes showed that these two viruses belong to the American avian lineage that is distinct from the Eurasian avian strains and the H2N2 viruses isolated from humans after the 1957 influenza pandemic (Fig. 1).

Molecular Analysis of the HA and NA Surface Proteins. Influenza A viruses contain two surface proteins: the HA is the receptor-binding and membrane-fusion glycoprotein, and the NA is a receptor-destroying enzyme. The viral HA is a critical factor of host species specificity of influenza viruses (15). To characterize residues within HA that may be associated with adaptation of an avian virus to the mammalian host, we compared the amino acid sequences of swine HAs with those of the putative reference avian viruses. Molecular comparison of the HA molecules of the two swine H2N3 isolates revealed that they differ from the putative reference H2N3 virus isolated from mallards by six common amino acid substitutions (D36N, Q226L, T274I, V316I, L419L, and L506V) (SI Table 6). The substitution Q226L was found in both swine H2N3 isolates, whereas position 228 contained G, identical to the avian consensus sequence (Table 1) (16). In contrast, human HA molecules of H2 subtype contain 226L and 228S, whereas early human H2 isolates contain 226L and 228G (Table 1), similar to the swine isolates. Positions 36N, 274I, 316I, and 419L are unique to the two swine H2N3 isolates (SI Table 6), whereas the respective positions in human and avian isolates depicted in Fig. 1a are 36D, 274T, 316V, and 419L. For the influenza isolates depicted in Fig. 1a, position 506S is conserved among human, two swine H2N3 isolates, and the avian isolates, except for A/mallard/Alberta/2004 (H2N3) as shown in SI Table 6. Two common amino acid changes in the NA amino acid sequence of both swine isolates were found when compared with the reference H4N3 virus isolated from blue-winged teal: H47Y and H253Y (SI Table 7). The position 47Y in both swine H2N3 isolates is the same as the respective amino acid in Eurasian avian isolates depicted in Fig. 1b; conversely, the position in North American avian isolates is 47H. The position 253Y is unique to the swine H2N3 isolates, and the position 253H is conserved in Eurasian and North American avian isolates depicted in Fig. 1b. Interestingly, Sw/4296424 (H2N3), isolated 5 months later than Sw/2124514 (H2N3), had two additional substitutions (P162S and L321V) in the HA molecule, and had three additional substitutions (V301I, 49T, and A135T) in the NA molecule when compared with the HA and NA of Sw/2124514 (SI Tables 6 and 7). The position 30I (Sw/4296424) in the NA molecule is similar to Eurasian isolates, whereas the position 30V (Sw/2124514) is conserved in Northern American avian isolates.

Pathogenicity and Transmissibility of H2N3 Swine Influenza Viruses in Pigs. To investigate the extent of swine adaptation of the H2N3 virus, we investigated its pathogenicity in this host by inoculating 20 4-week-old pigs with 2 × 105 50% tissue culture infective dose (TCID50) of the Sw/4296424 virus. Only one H2N3 virus was chosen, because of the high identity between the two isolates. Twelve control pigs were mock-inoculated with noninfectious cell culture supernatant. We assessed transmissibility by cohousing 10 age-matched contact pigs with the inoculated pigs, starting on day 3 p.i. All pigs used for the study were seronegative at day 0 for antibodies against swine influenza H1N1, H1N2, H2N3, and H3N2 viruses by HI assay. Five inoculated pigs and three control pigs were euthanized for necropsy on days 3, 5, and 7 p.i. The 10 contact pigs and 5 virus-inoculated pigs were serologically tested by HI assay with H2N3 on day 24 after contact or day 27 p.i., respectively. No acute respiratory signs were observed. Necropsy revealed severe macroscopic lung lesions (plum-colored, consolidated areas) in pigs inoculated with the H2N3 virus.
(BALF) and isolated from nasal swab samples. Virus titers in the vessels (Fig. 2). Virus was titrated in bronchoalveolar lavage fluid with slight lymphocytic cuffing of bronchioles and inoculated pigs euthanized on day 3, 5, or 7 p.i. exhibited mild to higher in the human H2 pandemic when compared to the other H2 human viruses listed here.

Pathogenicity of H2N3 Swine Influenza Viruses in Mice. To test the ability of the H2N3 Sw/4296424 virus to replicate in mice, we inoculated 6- to 7-week-old BALB/c mice intranasally with 102-106 TCID50/ml on days 3 and 5 (SI Table 8) and were negative on day 7 p.i. In the H2N3 inoculated group, virus was isolated from nasal swab samples in 25% (5 of 20) of pigs on day 3, 67% (10 of 15) on day 5, and 20% (2 of 10) on day 7 p.i.; in the contact group, 10% (1 of 10) of samples were positive on days 5 and 7 after contact. In contrast, 100% (10 of 10) of the contact pigs were seropositive after 24 days of contact with inoculated pigs (SI Table 9). Some control pigs had an occasional small focus of mild interstitial pneumonia (Table 2), but they were negative for swine influenza virus infection. All pigs were negative for PRRSV and M. hyopneumoniae by PCR. Our results indicate that the H2N3 virus is pathogenic in pigs and is transmissible among pigs.

Transmissibility of H2N3 Swine Influenza Virus in Ferrets. To cause a pandemic, an emergent influenza A virus must infect humans and be efficiently transmitted among humans. To investigate the potential of the reassortant H2N3 virus to transmit in mammalian systems, we used the ferret contact model. Three 18-week-old ferrets, housed in separate cages, were inoculated with 106-107 TCID50 of the H2N3 virus Sw/2124514. After 24 h, one contact animal was placed in each cage. Nasal washes were taken on days 1, 4, and 7 p.i., and virus was titrated in embryonated eggs. Virus was detected in all inoculated and contact ferrets, but none showed obvious clinical signs (Table 3). These results indicate that the H2N3 influenza virus infected ferrets and was transmitted via contact efficiently.

Serological Investigation of H2N3 Swine Influenza Viruses in Outbreak Farms. To further investigate the spread of the H2N3 viruses, we conducted a limited serological survey of animals associated with the two affected production systems. In the first study, in spring 2007, serum samples were taken from sows from four farms that provided piglets to the nursery farms during the September 2006 outbreak. Ninety percent (54 of 60) were seropositive for the presence of antibodies to Sw/4296424 (SI Table 11). A number of the tested animals were present at the time of the index case, and it is unclear whether they were infected at that time or whether they were infected subsequently. The data do, however, show that the virus was present at both sow and nursery farms and that the virus efficiently transmitted between animals. All sows in this operation were infected with antibodies to Sw/4296424 virus by the HI assay. Of the 30 sows and 90 weaned pigs sampled, 1 of 30 and 26 of 90 were seropositive (SI Table 11), respectively.

Discussion

In this report we characterized reassortant H2N3 viruses isolated from pigs in the United States. Molecular and phylogenetic analysis revealed that the HA, NA, and PA gene segments are similar to

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Table 1. Comparison of amino acids in HA receptor-binding site of human, avian, and swine H2 influenza virus isolates

<table>
<thead>
<tr>
<th>Virus strains</th>
<th>Avian consensus</th>
<th>Mallard/2003/H2N3</th>
<th>Sw/4296424</th>
<th>Sw/2124514</th>
<th>Human consensus</th>
<th>Davis/1/57</th>
<th>Albany/7/57</th>
<th>R/P/S-57</th>
<th>Albany/6/58</th>
<th>Ohio/2/59</th>
<th>Berlin/3/64</th>
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Under HA receptor-binding residues, the H3 numbering system is used for numbers before the slash, and the H2 numbering system is used for numbers after the slash. The Davis/1/57 and Albany/7/57 viruses were isolated earlier in the human H2 pandemic when compared to the other H2 human viruses listed here.

Table 2. Macroscopic and microscopic pneumonia in pigs inoculated with H2N3 virus Sw/4296424 or mock-inoculated

<table>
<thead>
<tr>
<th>Group</th>
<th>Lung lesion score, %</th>
<th>Histopathologic score (0–3)</th>
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<tr>
<td>H2N3 day 3</td>
<td>27.57 ± 7.36</td>
<td>2.23 ± 0.23</td>
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<tr>
<td>Control day 3</td>
<td>0.00 ± 0.00</td>
<td>0.33 ± 0.09</td>
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<tr>
<td>H2N3 day 5</td>
<td>21.86 ± 2.90</td>
<td>2.37 ± 0.11</td>
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<tr>
<td>Control day 5</td>
<td>0.00 ± 0.00</td>
<td>0.56 ± 0.06</td>
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<tr>
<td>H2N3 day 7</td>
<td>21.57 ± 5.02</td>
<td>2.07 ± 0.25</td>
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<tr>
<td>Control day 7</td>
<td>0.00 ± 0.00</td>
<td>0.22 ± 0.11</td>
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Values are the mean ± SEM.
those of AIVs of the American lineage, whereas other gene segments are similar to those of contemporary swine influenza viruses that are triple-reassortant viruses containing human, avian, and swine influenza virus genes. In addition to their potential impact on animal health, these H2N3 viruses have intrinsic properties that make them of considerable concern to public health. These properties include the following: (i) they belong to the H2 subtype as did the 1957 human pandemic strain that disappeared in 1968 (hence, individuals born subsequent to 1968 have little pre-existing immunity to this subtype); (ii) they are circulating in swine, a host shown experimentally to select for mammalian virus traits (7); (iii) they have receptor binding site changes associated with increased affinity for α2,6Gal-linked sialic acid viral receptors; and (iv) they are able to replicate and transmit in swine and ferrets via contact. The latter two points suggest that the swine H2N3 viruses have undergone adaptation to the mammalian host and as such have the ability for sustained transmission. Reinforcing this possibility is the finding in one of the production systems that young pigs born at least 6 months after the index case were seropositive for the virus. Although it is not clear whether the seropositivity in the young animals was due to infection or maternal antibodies, these data suggest that the virus continues to circulate within the affected production systems. The fact that the H2N3 viruses are known to have infected two independent swine production systems and that serologic studies suggest they continue circulating is in contrast to other reports of a wholly AIV infection in North American swine, in which infections have appeared to be self-limiting (19, 20).

Although the genetic factors that are associated with successful zoonotic transmission of influenza viruses remain largely unknown, receptor-binding properties are likely involved. In avian H2 and H3 influenza viruses, HA receptor-binding-site residues corresponding to codon positions 138, 190, 194, 225, 226, and 228 (using the H3 numbering system) are highly conserved (16). In human H2 and H3 viruses, leucine and serine substitutions at residues 226 and 228, respectively, have been shown to accompany their adaptation from avian to human hosts (21). For example, changing the human virus H3 residue L226 to Q226 dramatically changes the receptor-binding specificity of the virus from mammalian to avian virus-like (22). The substitution Q226L was found in both new swine H2N3 isolates, whereas position 228 retains G, which is typical of AIVs (Table 1). This same 226L/228G combination was observed in the first viruses in which infections have appeared to be self-limiting (19, 20).

In April 2006, an outbreak of respiratory disease occurred in pigs at a commercial grower-finisher swine farm. At necropsy, the attending veterinarian observed gross lesions of pneumonia and submitted formalin-fixed and unfixed sections of lung tissue to the Minnesota Veterinary Diagnostic Laboratory (MVDL). At the MVDL, the formalin-fixed tissue was routinely processed for histopathology. Bronchial swab samples from the unfixed lung tissue were suspended in 2 ml of PBS and tested for M. hypovirulentiae by PCR (27). Unfixed sections (≈5 g) of lung were cultured aerobically for bacteria by inoculation on MacConkey, colistin–nalidixic acid, brilliant green, and blood agar plates with and without nitrocefinadine dinitrocellulose (NAD) factor (Staphylococcus epidermidis nurse colonies). In parallel, unfixed sections of lung (≈10 g) were homogenized in Eagle’s minimal essential medium (MEM) containing 4% BSA, 15 μg/ml trypsin, and an antibiotic mixture of neomycin, gentamicin, penicillin, streptomycin, and amphotericin B and were cultivated on MDCK cells. RNA and DNA were also isolated from the homogenate for diagnostic tests for influenza virus nucleoprotein (RT-PCR), PRRSV ORF 6 (RT-PCR), and PCV2 ORF 2 (PCR).

For virus isolation, 10% lung homogenates were centrifuged for 10 min at 640 × g. The supernatant was passed through a 0.45-μm filter to remove any bacterial contamination and was inoculated on monolayers of MDCK cells in 24-well plates. The MDCK cells were maintained in Eagle’s MEM containing 1 μg/ml Tc-pc-trypsin and 0.3% bovine albumin. The plates were incubated at 37°C in a CO2 incubator and were observed daily. After cytopathic effects were observed, infected cells were lysed by freezing and thawing, and virus was serotyped by HI assays with turkey erythrocytes.

In September 2006, another outbreak of respiratory disease occurred in 5- to 6-week-old pigs at a different multisourced commercial swine nursery. Again, gross lesions were consistent with pneumonia, and lung tissue samples were submitted to the MVDL for testing as described above.

**HI Assays.** HI assays were performed for serologic subtyping of H2N3 viruses to determine seroenconversion and to test convalescent serum samples collected from the various swine herds associated with the outbreak. Sera were heat-inactivated at 56°C, treated with a 20% suspension of kaolin (Sigma–Aldrich) to eliminate nonspecific inhibitors, and adsorbed with 0.5% turkey red blood cells. The sera were tested for antibodies against H2N3 swine influenza viruses and reference strains of swine influenza (A/Swine/IA/1973 H1N1, A/Swine/TX/98 H3N2, and A/Swine/NC/2001 variant H1N1) virus by HI assay (28). The ferret sera were tested to determine seroenconversion for H2N3 virus.

**DNA Sequencing, Phylogenetic Analysis, and Subtype Determination.** Viral RNA was prepared from 200 μl of virus suspension with the RNeasy Mini Kit (Qiagen) as directed by the manufacturer. Two-step RT-PCR was conducted by using universal primers as reported in refs. 29 and 30. Each gene segment was amplified.
under standard conditions. PCR products were purified by using a QIAamp Gel extraction kit (Qiagen) and sequenced by using an ABI 3730 DNA Analyzer (Applied Biosystems). Multiple sequence alignments were made by using CLUSTAL W (31), and phylogenetic trees were generated by using the neighbor-joining algorithm in the PHYLIP version 3.57c software package (32). A Mega-blast search of the Influenza Sequence Database was performed. The viral subtype determined by sequencing was compared with those from GenBank. The isolate was plaque-purified, retested, and again subtyped by RT-PCR and sequencing (plaques were uniform in appearance, and two plaques from each isolate were chosen for amplification and sequencing).

Experiments in Pigs. Pigs were obtained from a healthy herd that was free of swine influenza virus and PRRSV. All animal experiments were in compliance with the Institutional Animal Care and Use Committee of the NADC. The inoculation protocol has been described in ref. 33. Briefly, 204-week-old cross-bred pigs were inoculated intratracheally with 2 × 10⁶ TCID₅₀ per pig of Sw/4296424 virus prepared in MDCK cells. Four-week-old control pigs were commingled with inoculated pigs on day 3 p.i. to study transmission efficiency. Twelve control pigs were inoculated with noninfectious cell culture supernatant and from the remaining five inoculated and control pigs on day 27 p.i. were euthanized and was analyzed for seroconversion. Each lung was lavaged with 50 ml of MEM supplemented with TPCK-trypsin and antibiotics. Each dilution (100 μl) was plated on PBS-washed confluent MDCK cells in 48-well plates. After incubation for 1 h at 37°C, 500-μl serum-free MEM supplemented with 1 μg/ml TPCK trypsin and antibiotics was added. All wells were evaluated for cytopathic effects after 48–72 h. Subsequently, plates were fixed with 4% phosphate-buffered formaldehyde and stained as described above.

BALF was tested for the presence of PRRSV and M. hyopneumoniae by diagnostic PCR assays. For PRRSV, the total RNA was isolated from each sample by using the RNeasy Mini Kit (Qiagen). One microgram of the extracted RNA and a primer pair specific for ORF 7 of PRRSV were used in real-time PCR as described in ref. 36. DNA was extracted from BALF for detecting M. hyopneumoniae as described in ref. 36.

Examination of Lungs of Experimental Pigs. At necropsy, lungs were removed in toto. A single experienced veterinarian recorded the percentage of gross lesions of lobes showing the purple-red consolidation typical of swine influenza virus infection. A mean value was determined for the seven pulmonary lobes of each animal (33). Tissue samples from the trachea, the right cardiac pulmonary lobe, and other affected lobes were fixed in 10% buffered formalin, routinely processed, and stained with hematoxylin and eosin for histopathologic examination. Lung sections were given a score of 0 to 3 to reflect the severity of bronchial epithelial injury (33) according to the following criteria: 0.0: no significant lesions; 1.0: a few airways showing (34). The TCID₅₀/ml was calculated for each sample by the method of Reed and Muench (35).

Virus was isolated from nasal swab samples stored at −80°C by thawing and vortexing each sample for 15 sec, centrifuging it for 10 min at 640 × g, and passing the supernatant through 0.45-μm filters to reduce bacterial contamination. An aliquot of 100 μl was plated on confluent, PBS-washed MDCK cells in 48-well plates. After incubation for 1 h at 37°C, 500-μl serum-free MEM supplemented with 1 μg/ml TPCK trypsin and antibiotics was added. All wells were evaluated for cytopathic effects after 48–72 h. Subsequently, plates were fixed with 4% phosphate-buffered formaldehyde and stained as described above.

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Experiments in Mice. Six- to 7-week-old BALB/c mice, bred in the mouse facility of the NADC in Ames, IA, were used for infectivity experiments. All experiments were in compliance with the Institutional Animal Care and Use Committee of the NADC. Animals were weighed and anesthetized with isoflurane USP (Phoenix Pharmaceutical) before intranasal inoculation with 10⁸–10⁹ TCID₅₀ of H2N3 virus (Sw/4296424) in a volume of 50 μl. Weight was recorded once daily, and general health status was observed twice daily. After onset of disease, general health status was observed three times per day. Surviving mice were euthanized on day 14 p.i., and the lungs were collected. The right lung was stored in an Eppendorf tube at −80°C for virus detection, and the left lung was fixed in 10% formalin for histopathologic analysis. Virus detection was done in a 10% tissue homogenate in PBS (homogenized twice for 1 min in a Mini BeadBeater-8; BioSpec Products). The homogenate was centrifuged at 640 × g for 5 min and the supernatant was

| Table 3. Virus titers in nasal washes from H2N3 (Sw/2145514)-inoculated and contact ferrets |
|--------------------------------------|-----------------|-----------------|-----------------|-----------------|-----------------|-----------------|-----------------|
| Animal                             | Virus titer, EID₅₀/ml | HI titer       |                 |                 |                 |                 |                 |
|                                     | Day 1   | Day 4   | Day 7   | Day 21   | Day 28   | Day 21   | Day 28   |
| Inoculated no. 333                 | 10⁶.8   | 10⁵.5   | <10¹⁰   | 1:160    | 1:160    | 1:320    | 1:320    |
| Contact no. 362                    | <10¹⁰   | 10⁶.3   | <10³     | 1:320    | 1:320    | 1:320    | 1:320    |
| Inoculated no. 366                 | 10⁶.0   | 10⁵.5   | <10³     | 1:320    | 1:320    | 1:320    | 1:320    |
| Contact no. 334                    | 10      | 10⁵.8   | <10³     | 1:320    | 1:320    | 1:320    | 1:320    |
| Inoculated no. 368                 | 10⁵.8   | 10⁵.5   | <10³     | 1:320    | 1:320    | 1:320    | 1:320    |
| Contact no. 364                    | <10¹⁰   | 10⁵.8   | 10⁴.3   | 1:320    | 1:320    | 1:320    | 1:320    |
transferred to 1.5-ml reaction tubes for RNA isolation. Real-time RT-PCR was used to detect viral RNA as described in ref. 17.

**Experiments in Ferrets.** Influenza-negative ferrets were obtained through the ferret breeding program and were housed at St. Jude Children’s Research Hospital in compliance with the St. Jude Children’s Research Hospital Animal Care and Use Committee. Infection and transmissibility of H2N3 was tested in six 18-week-old ferrets. Three ferrets were inoculated intranasally with $10^{2.5}$ TCID$_{50}$ of H2N3 (Sw/2124514). Twenty-four hours p.i., one naïve contact animal was introduced into the cage of each inoculated animal. Nasal washes were collected on days 1, 4, and 7 p.i., and virus was titrated in embryonated eggs. Egg 50% infective dose (EID$_{50}$) values were calculated by the Reed–Muench method (35). Serocconversion was determined after 21 days by determining the serum neutralizing antibody titers of the inoculated and contact ferrets, as described in ref. 37.

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