# Household Transmission of Pandemic (H1N1) 2009, San Antonio, Texas, USA, April-May 2009

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To assess household transmission of pandemic (H1N1) 2009 in San Antonio, Texas, USA, during April 15-May 8, 2009, we investigated 77 households. The index case-patient was defined as the household member with the earliest onset date of symptoms of acute respiratory infection (ARI). influenza-like illness (ILI), or laboratory-confirmed pandemic (H1N1) 2009. Median interval between illness onset in index and secondary case-patients was 4 days (range 1-9 days); the index case-patient was likely to be ≤18 years of age (p = 0.034). The secondary attack rate was 4% for pandemic (H1N1) 2009, 9% for ILI, and 13% for ARI. The secondary attack rate was highest for children <5 years of age (8%-19%) and lowest for adults  $\geq$ 50 years of age (4%-12%). Early in the outbreak, household transmission primarily occurred from children to other household members and was lower than the transmission rate for seasonal influenza.

On April 15 and 17, 2009, the first 2 cases of pandemic (H1N1) 2009 in the United States were identified among children in California; within 10 weeks, the strain was identified in 99 countries or territories (1). Texas was

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the second US state to confirm human transmission of pandemic (H1N1) 2009. On April 24, 2009, the Texas Department of State Health Services reported 2 patients with laboratory-confirmed pandemic (H1N1) 2009 infection in Guadalupe County. The strain was similar to that isolated previously from patients in Mexico and California (2). On June 11, 2009, the World Health Organization raised the pandemic alert to phase 6, indicating that a global pandemic was under way (3).

Characterizing transmission dynamics in various settings, such as households, schools, and the community, is critical to the development of appropriate guidance and public health interventions. Household contacts of persons with seasonal influenza are at increased risk for infection (4-7), but the household transmission characteristics of pandemic (H1N1) 2009 have yet to be fully characterized. This study reports household secondary attack rates and serial time intervals between illness onset in the index case-patient to illness onset in a household contact. We investigated persons with laboratory-confirmed pandemic (H1N1) 2009 and their household contacts in 1 health service region of Texas.

# Methods

#### Population

The Texas Department of State Health Services consists of 11 health service regions. We conducted our investigation in Health Service Region 8, which includes 28 counties in south-central Texas, bordered on the south by Mexico (Figure). Approximately 2.4 million persons live in the region; 1.5 million live in the city of San Antonio (8).

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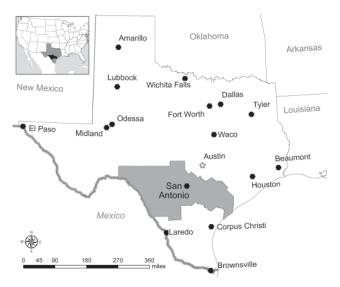


Figure. Texas Department of State Health Services Health Service Region 8 (gray shading), Texas, USA.

The Texas Department of State Health Services provides public health services in counties without local health departments. Within Health Service Region 8, public health services for Bexar County are provided by the San Antonio Metropolitan Health District and for Comal County by the Comal County Health Department.

## **Case Definitions**

We defined a laboratory-confirmed case-patient as a resident of Health Service Region 8 who had a positive respiratory specimen showing nucleic acid sequences unique to pandemic (H1N1) 2009; a real-time reverse transcription-PCR (rRT-PCR) assay was used to detect the virus (9). For persons with no laboratory test performed, we assessed whether they had influenza-like illness (ILI), defined as fever (measured or unmeasured) with either cough or sore throat; or acute respiratory infection (ARI), defined as  $\geq 2$  of the following signs or symptoms: fever, cough, sore throat, and rhinorrhea. The index case-patient was defined as the household member with the earliest symptom onset date of ARI, ILI, or laboratory-confirmed pandemic (H1N1) 2009. A secondary case-patient was defined as a household member with ARI, ILI, or laboratory-confirmed pandemic (H1N1) 2009 and symptom onset 1-9 days after symptom onset in the index case-patient. We chose the maximum interval of 9 days because shedding of seasonal influenza virus uncommonly lasts >8 days (10) and the median incubation period for seasonal influenza is  $\approx 1.4$  days (11). Household members were defined as persons who lived at the same address as a case-patient who had laboratory-confirmed pandemic (H1N1) 2009 infection.

## **Case Finding**

During April 10-May 8, 2009, we identified laboratory-confirmed cases of pandemic (H1N1) 2009 by reviewing 1,167 laboratory records of influenza specimens submitted by healthcare providers for rRT-PCR testing by the regional public health laboratory in San Antonio. We also reviewed 1,251 laboratory records of all specimens submitted by military medical treatment facilities in San Antonio. These specimens were tested for influenza by rRT-PCR at the Epidemiology Laboratory Service of the Department of Defense Global Influenza Surveillance Program at the United States Air Force School of Aerospace Medicine in San Antonio. In addition, we conducted telephone interviews with 540 (67%) of 802 high school students who were reported as absent by their school administrators during April 9-28 in the Texas counties where the first 2 identified case-patients attended school. Respiratory samples were collected from students who reported an acute respiratory illness at the time of interview. Additional casepatients were identified by collecting respiratory samples from nonhousehold contacts of laboratory-confirmed casepatients (i.e., those who had been within 6 feet of someone with ARI for at least 1 hour during the period 1 day before through 7 days after onset of illness in the contact).

## **Household Investigations**

We interviewed case-patients with laboratory-confirmed infection and all household members about the occurrence of illness, receipt of influenza vaccination in the previous 12 months, and medical history. We asked all persons about their use of antiviral medication and reviewed health department pharmacy records where appropriate to ascertain the type, dosage, and timing of antiviral medication and to define whether antiviral medications were prescribed for treatment or prophylaxis. Respiratory samples were collected from household contacts who had an acute respiratory illness at the time of interview; respiratory samples were collected from all members of 9 households identified early in the investigation, regardless of respiratory symptoms.

#### Sample Collection and Laboratory Testing

Nasal wash samples were collected from military servicemen and their household family members, and nasopharyngeal swabs were collected from all others. Nasal wash samples were sent to the Epidemiology Laboratory Service at the United States Air Force School of Aerospace Medicine; nasopharyngeal swabs were sent to the regional public health laboratory in San Antonio. We used rRT-PCR to test all respiratory samples for seasonal influenza (A/H1 and A/H3 influenza viruses). Specimens positive for influenza A but negative for seasonal influenza by rRT-PCR were sent to the Centers for Disease Control and Prevention (CDC) for confirmatory testing for pandemic (H1N1) 2009 (12).

## **Statistical Analysis**

We calculated the serial interval as the number of days from the onset date of illness in the index case-patient to onset date of illness in the secondary case-patient. Secondary household attack rates were calculated by dividing the number of secondary case-patients (excluding the index case-patient) by the total number of household members (excluding the index case-patient). Secondary case-patients for ILI and ARI attack rates also included laboratory-confirmed case-patients. We compared characteristics between groups by using the  $\chi^2$  test or Fisher exact test for categorical data and the Wilcoxon signed-rank test for continuous variables (13).

## Ethics

The collection of information about cases of pandemic (H1N1) 2009 was part of the emergency public health practice response and was not deemed to be research in accordance with the federal human subjects protection regulations (45 Code of Federal Regulations 46.101c and 46.102d) and CDC's Guidelines for Defining Public Health Research and Public Health Non-Research. All protocols pertaining to the pandemic were reviewed for protection concerns and the necessity of Institutional Review Board review by the CDC's National Center for Immunization and Respiratory Diseases (NCIRD) Human Subjects Contact and the NCIRD Associate Director of Science.

## Results

We identified 110 persons with laboratory-confirmed pandemic (H1N1) 2009 infection. We were unable to contact 23 (21%) of these persons, and 3 (3%) did not agree to provide further information. Of 84 persons with laboratory-confirmed pandemic (H1N1) 2009 infection who provided information, 77 (92%) lived with  $\geq$ 1 persons. These 77 households comprised 349 persons; the median household size was 4 persons (range 2–9 persons), including the index case-patient. Seventy five percent of household interviews were conducted  $\geq$ 8 days (range 0–24 days) after the onset of infection in the index case-patient.

From household interviews, we identified an additional 47 persons who reported respiratory symptoms or had laboratory evidence of pandemic (H1N1) 2009 infection: 13 persons with laboratory-confirmed pandemic (H1N1) 2009 infection, 24 persons with ILI, and 10 persons whose illness met the case definition for ARI only. We did not classify 15 of these persons as secondary case-patients: 8 persons had the same date of symptom onset as the index case-patient; we could not establish the date of symptom onset for 3 persons; and 4 persons reported illness onset 10–15 days after the index case-patient. In 1 household where 2 persons had ILI, 1 had a nasopharyngeal swab that was positive for pandemic (H1N1) 2009; the other was positive for influenza A, but the subtype could not be determined, possibly because of the quality of the sample or because 9 days had elapsed between illness and sample collection, thus decreasing viral load. We considered this person to have laboratory-confirmed pandemic (H1N1) 2009 on the basis of an epidemiologic link to another laboratory-confirmed case. In 2 households where secondary case-pateints were identified, nasal swab samples were obtained from members of all 7 households; 1 person, 14 years of age, who did not report any respiratory symptoms, was positive for pandemic (H1N1) 2009 infection.

Among the 97 symptomatic laboratory-confirmed casepatients (84 identified through case finding and 13 through household investigation), illness onset dates ranged from April 11 through May 8, 2009. Eleven (11%) were ≤4 years of age, 61 (63%) 5-18 years of age, 22 (23%) 19-49 years of age, and 3 (3%) >50 years of age. Forty-six (47%) were male. The most common signs and symptoms were fever (93%), cough (91%), rhinnorhea (70%), headache (67%), and sore throat (58%). Vomiting was reported by 26% and diarrhea by 25%. Ninety-two percent of laboratoryconfirmed case-patients met the definition for ARI, and 85% met the definition for ILI. One laboratory-confirmed case-patient was hospitalized: a child who was admitted to the hospital with pneumonia in early April. No deaths occurred. Compared with household contacts who did not have laboratory-confirmed pandemic (H1N1) 2009 or did not report respiratory illness, laboratory-confirmed casepatients (index and secondary) were significantly younger (median age 17 vs. 24 years; p<0.001).

Secondary case-patients were found in 24 (31%) of 77 households; 5 had 2 secondary case-patients, and 1 had 3 case-patients (Table 1). Secondary infections appeared most likely to be transmitted between children (12/32), 38%) or children to adults (10/32, 31%) than from adults to children (6/32, 19%) or adults to adults (4/32, 13%) (p = 0.034). The median serial interval for ARI, ILI, and laboratory-confirmed pandemic (H1N1) 2009 combined was 4 days (range 1–9 days) (Table 1; online Appendix Figure, www.cdc.gov/EID/content/16/4/631-appF.htm). Antiviral treatment was given to the index case-patient of 23 (72%) of 32 secondary case-patients; in these households, the serial interval was 3 days, compared with 5 days when the index case-patient was not given treatment (p = 0.17). Inclusion of 5 household contacts with illness that occurred 10-15 days after symptom onset of the index case-patient did not alter the median serial interval estimate. The median serial interval also remained unchanged when only members of households interviewed >9 days after the onset of symptoms in the household index case-patient were

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included. Limiting the estimate of median serial interval to include only persons with ILI or laboratory-confirmed case-patients reduced the median serial interval to 3 days (range 1–8 days).

The secondary household attack rate was 13% for ARI, 9% for ILI, and 4% for laboratory-confirmed pandemic (H1N1) 2009 (Table 2). Secondary attack rates were highest in children <5 years of age and were higher in children 5–18 years of age than in adults 19–49 and  $\geq$ 50 years of age (Table 2). By household size, secondary attack rates for ARI, ILI, and laboratory-confirmed pandemic (H1N1) 2009 were highest in households with 2–3 persons (ARI 23%, ILI 23%, laboratory-confirmed pandemic [H1N1] 2009 6%) and were lowest in households with 7–9 persons (ARI 9%, ILI 9%, laboratory-confirmed pandemic [H1N1] 2009 2%) (online Appendix Table 1, www.cdc.gov/EID/ content/16/4/631-appT1.htm). The secondary household attack rate did not differ by receipt of seasonal influenza vaccination in the previous 12 months (online Appendix Table 2, www.cdc.gov/EID/content/16/4/631-appT2.htm).

Treatment with antiviral medication was given to 77% of index case-patients (57/74 of persons with ARI, ILI, and laboratory-confirmed pandemic [H1N1] 2009 combined for whom data were available) and 72% of secondary cases (23/32 of ARI, ILI, and laboratory-confirmed pandemic [H1N1] 2009 combined); 90% took oseltamivir; 7% took zanamivir; and 3% took an unknown type of antiviral medication. Neither the age of the index case-patient, household

| Texas, April-                 |               |          | ld case-patients with ARI, IL | -             |        | . , .                | 1011 0,      |
|-------------------------------|---------------|----------|-------------------------------|---------------|--------|----------------------|--------------|
|                               |               | Index ca | se-patients                   | Se            | Serial |                      |              |
| Household                     | Date of onset | Age, y   | Case definition               | Date of onset | Age, y | Case definition      | interval, d† |
| А                             | Apr 18        | 14‡      | A, no subtype                 | Apr 25        | 21     | Pandemic (H1N1) 2009 | 7            |
| В                             | Apr 19        | 5        | Pandemic (H1N1) 2009          | Apr 21        | 9      | Pandemic (H1N1) 2009 | 2            |
| С                             | Apr 22        | 18       | Pandemic (H1N1) 2009          | Apr 25        | <1     | Pandemic (H1N1) 2009 | 3            |
| D                             | Apr 26        | 1        | Pandemic (H1N1) 2009          | May 4         | 27     | Pandemic (H1N1) 2009 | 2            |
| E                             | Apr 26        | 16       | Pandemic (H1N1) 2009          | Apr 27        | 51     | Pandemic (H1N1) 2009 | 1            |
|                               |               |          |                               | Apr 27        | 8      | ILI                  | 1            |
| F                             | Apr 27        | <1       | Pandemic (H1N1) 2009          | Apr 29        | 22     | Pandemic (H1N1) 2009 | 2            |
|                               |               |          |                               | May 6         | 47     | ARI                  | 9            |
| G                             | Apr 27        | 16       | Pandemic (H1N1) 2009          | May 1         | 16     | Pandemic (H1N1) 2009 | 4            |
|                               |               |          |                               | May 1         | 14     | Pandemic (H1N1) 2009 | 4            |
| Н                             | Apr 29        | 6        | Pandemic (H1N1) 2009          | Apr 3         | <1     | Pandemic (H1N1) 2009 | 1            |
| I                             | May 3         | 33       | Pandemic (H1N1) 2009          | May 7         | 15     | Pandemic (H1N1) 2009 | 4            |
|                               |               |          |                               | May 8         | 14     | Pandemic (H1N1) 2009 | 5            |
| Subtotal no. case-patients 9  |               |          |                               |               |        | 13                   | 3 (1–9)      |
| J                             | Apr 20        | 17       | Pandemic (H1N1) 2009          | Apr 26        | 14     | ARI                  | 6            |
| К                             | Apr 24        | 71       | Pandemic (H1N1) 2009          | Apr 27        | 65     | ILI                  | 3            |
| L                             | Apr 25        | 16       | Pandemic (H1N1) 2009          | Apr 27        | 16     | ILI                  | 2            |
| Μ                             | Apr 25        | 12       | Pandemic (H1N1) 2009          | Apr 28        | 30     | ARI                  | 3            |
|                               |               |          |                               | Apr 28        | 33     | ARI                  | 3            |
|                               |               |          |                               | Apr 30        | 6      | ARI                  | 5            |
| Ν                             | Apr 26        | 30       | Pandemic (H1N1) 2009          | May 1         | 28     | ARI                  | 5            |
| 0                             | Apr 27        | 33       | Pandemic (H1N1) 2009          | May 5         | 53     | ARI                  | 8            |
| Р                             | Apr 28        | 25       | Pandemic (H1N1) 2009          | May 4         | 14     | ILI                  | 6            |
| Q                             | Apr 29        | 1        | Pandemic (H1N1) 2009          | May 1         | 21     | ILI                  | 2            |
|                               |               |          |                               | May 2         | 2      | ILI                  | 3            |
| R                             | Apr 29        | 8        | Pandemic (H1N1) 2009          | May 3         | 44     | ARI                  | 4            |
| S                             | May 1         | 6        | Pandemic (H1N1) 2009          | May 3         | 45     | ILI                  | 2            |
| Subtotal no. case-patients 10 |               |          |                               |               |        | 13                   | 3 (1–8)      |
| Т                             | Apr 17        | 11       | ILI                           | Apr 21        | 18     | Pandemic (H1N1) 2009 | 4            |
| U                             | Apr 18        | 48       | ILI                           | Apr 26        | 10     | Pandemic (H1N1) 2009 | 8            |
| V                             | Apr 23        | 53       | ILI                           | Apr 26        | 42     | Pandemic (H1N1) 2009 | 3            |
| W                             | Apr 24        | 5        | ILI                           | Apr 29        | <1     | Pandemic (H1N1) 2009 | 5            |
| Х                             | Apr 28        | 26       | ILI                           | May 2         | 7      | Pandemic (H1N1) 2009 | 4            |
|                               |               |          |                               | May 3         | 4      | ILI                  | 5            |
| Subtotal no.                  | case-patients |          | 5                             | 2             |        | 6                    | 4.5 (3–8)    |
| Total no. case-patients 24    |               |          |                               |               |        | 32                   | 4 (1–9)      |

\*ARI, acute respiratory infection; ILI, influenza-like illness (fever measured or subjective and cough or sore throat).

†Median (range) number of days between symptom onset of the index and secondary case-patients.

<sup>‡</sup>The influenza virus from this person could not be subtyped, possibly because of the quality of the sample or the length of time from symptom onset to sample collection. We considered this case-patient to have been infected with pandemic (H1N1) 2009.

#### Household Transmission of Pandemic (H1N1) 2009

| 2. index<br>2-patients<br>7<br>50<br>17<br>3<br>77<br>6<br>50 | Secondary<br>case-patients, A<br>5<br>13<br>11<br>3<br>32<br>5<br>44 | Not ill, B<br>23<br>83<br>96<br>22<br>224<br>23        | Total household<br>contacts, A + B        28        96        107        25        256        28 | members not<br>included<br>1<br>3<br>3<br>1<br>8<br>2  | Secondary attack<br>rate (A/A + B), %<br>18<br>14<br>10<br>12<br>13             |
|---|--|--|--|--|---|
| 17<br>3<br>77<br>6  | 13<br>11<br>3<br>32<br>5   | 83<br>96<br>22<br>224                                  | 96<br>107<br>25<br>256   | 1<br>3<br>1<br>8   | 14<br>10<br>12<br>13  |
| 17<br>3<br>77<br>6  | 13<br>11<br>3<br>32<br>5   | 83<br>96<br>22<br>224                                  | 96<br>107<br>25<br>256   | 1<br>3<br>1<br>8   | 14<br>10<br>12<br>13  |
| 17<br>3<br>77<br>6  | 11<br>3<br>32<br>5   | 96<br>22<br>224  | 107<br>25<br>256   | 3<br>3<br>1<br>8   | 10<br>12<br>13  |
| 3<br>77<br>6  | 3<br>32<br>5   | 22<br>224  | 25<br>256  | 3<br>1<br>8<br>2   | 12<br>13  |
| 6   | 32<br>5  | 224  | 256  | 1 8 2  | 13  |
| 6   | 5  |  |  | 8  |   |
| •   | -  | 23   | 28   | 2  | 10  |
| •   | -  | 23   | 28   | 2  | 10  |
| 50  | 4.4  |  |  | -  | 18  |
|   | 11   | 86   | 97   | 2  | 11  |
| 18  | 6  | 102  | 108  | 1  | 6   |
| 3   | 2  | 23   | 25   | 1  | 8   |
| 77  | 24   | 234  | 258  | 6  | 9   |
| mic (H1N1) 200  | )9   |  |  |  |   |
| 8   | 2  | 26   | 28   | 0  | 7   |
| 51  | 5  | 92   | 97   | 1  | 5   |
| 16  | 3  | 108  | 111  | 0  | 3   |
| 2   | 1  | 26   | 27   | 0  | 4   |
| 77  | 11   | 252  | 263  | 1  | 4   |
|   | mic (H1N1) 200<br>8<br>51<br>16<br>2<br>77                           | mic (H1N1) 2009<br>8 2<br>51 5<br>16 3<br>2 1<br>77 11 | mic (H1N1) 2009<br>8 2 26<br>51 5 92<br>16 3 108<br>2 1 26<br>77 11 252                          | mic (H1N1) 2009<br>8 2 26 28<br>51 5 92 97<br>16 3 108 111<br>2 1 26 27<br>77 11 252 263<br>ILI, influenza-like illness (fever measured or subjective and cough or sore thro | mic (H1N1) 2009<br>8 2 26 28 0<br>51 5 92 97 1<br>16 3 108 111 0<br>2 1 26 27 0 |

Table 2. Household secondary attack rates for ARI, ILI, and laboratory-confirmed pandemic (H1N1) 2009, by age group, Region 8, Texas, April–May 2009\*

size, nor diagnosis of the index patient (with ARI, ILI, or laboratory-confirmed pandemic [H1N1] 2009) were predictive of treatment with antiviral medication. The secondary household attack rates for ARI, ILI, and laboratoryconfirmed pandemic (H1N1) 2009 combined in households where the index case-patient was given antiviral treatment was 12% compared with 16% in other households (p = 0.64) (Table 3). Antiviral prophylaxis was given to 39% of household contacts (92/235 with data available) (Table 3), and the secondary attack rate of ARI, ILI, and laboratory-confirmed pandemic (H1N1) 2009 combined was 14% (12/83) in households where the index patient took treatment, compared with 66% (6/9) (p = 0.003) in households where the index patient did not take treatment (Table 3).

## Discussion

During an outbreak of pandemic (H1N1) 2009 in the San Antonio, Texas, area, we identified 97 persons with laboratory-confirmed infection in 77 households. The epidemiologic and clinical features were similar to summary reports from the United States (14,15) and other countries (15,16). Nearly one third of households had secondary

case-patients who also had respiratory illness, with a median of 4 days between onset of illness in the index casepatient and household members, a finding similar to that for seasonal influenza (17).

The secondary attack rate was 4% for laboratoryconfirmed pandemic (H1N1) 2009, 9% for ILI, and 13% for ARI. In general, these rates are lower than for seasonal influenza and lower than anticipated for a pandemic strain, although rates vary from 13% to 30%, depending on influenza subtype and year and pandemic period (4,5,18-21). The highest proportion of laboratory-confirmed pandemic (H1N1) 2009 and secondary attack rates occurred in children, a finding consistent with the epidemiology of seasonal and pandemic influenza, where we know children experience higher rates of illness (4,5,7) and higher secondary attack rates (19). Adults may have some cross-protection against pandemic (H1N1) 2009 from antibodies developed during infections with seasonal influenza A virus (H1N1) (22-24).

Four randomized controlled trials of zanamivir and oseltamivir for seasonal influenza have shown that these antiviral medications reduce but do not eliminate viral shedding

Table 3. Household secondary attack rates for ARI, ILI, and laboratory-confirmed pandemic (H1N1) 2009, by antiviral medication treatment and prophylaxis, Region 8, Texas, April–May 2009\*

|   |              | Index case-patient rec | eived antiviral treatmer | nt (attack rate, %) |
|---|--------------|------------------------|--------------------------|---------------------|
| Type of contact                         | No. contacts | Yes                    | No                       | p value†            |
| All contacts of index case-patients     | 235          | 22/185 (12)            | 8/50 (16)                | 0.64                |
| Contacts who took antiviral prophylaxis | 92           | 12/83 (14)             | 6/9 (67)                 | 0.003               |

\*ARI, acute respiratory infection; ILI, influenza-like illness (fever measured or subjective and cough or sore throat).

+Fisher exact test comparing the secondary attack rate for any treatment to no antiviral treatment. Data about antiviral medication were missing for 2 index case-patients and 15 contacts.

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and are effective in preventing disease among household contacts, especially if taken within 48 hours of illness onset in the index case-patient (19,20,25,26). We found that secondary attack rates for all households were lower when the index case-patient received treatment, although this difference was not significant. The role of prophylaxis in the absence of treatment of the index case-patient was difficult to determine; our investigation included only a small number of such persons. Nevertheless, because most index and secondary case-patients received antiviral treatment, household secondary attack rates may have been reduced.

Our investigation has several limitations. Because early case finding was most intensive among high school children associated with school outbreaks, our cohort may have been biased in favor of households where the index case-patients were children; however, this would not explain a lower secondary attack rate among adult household contacts. We did not assess the role of mild or asymptomatic pandemic (H1N1) 2009 infection because we collected respiratory samples only; serologic assays to detect influenza antibodies are the most sensitive method for detecting asymptomatic infection, but virus assays for pandemic (H1N1) 2009 were not available at the time of the investigation. Volunteer challenge studies with seasonal influenza viruses have found that up to 30% of infected persons may be asymptomatic and could be identified through serologic testing (10). Because 25% of household interviews were conducted <8 days after onset of illness of the index case-patient, we may have underestimated the secondary attack rate if these households had secondary case-patients with long serial intervals. However, when we restricted our analysis to persons interviewed  $\geq 8$  days after onset of symptoms in the index case-patient, we found no difference in the median serial interval or distribution of attack rates by age. Conversely, household members interviewed >8 days after onset of illness in the index case-patient may have had incomplete recall of acute respiratory infections. Finally, some of the secondary illnesses may have been acquired in the community, leading to overestimate of household secondary attack rates.

We found that pandemic (H1N1) 2009 disproportionately affected children, who in turn posed a risk for secondary household transmission, especially to their caregivers and siblings. The Advisory Committee on Immunization Practices (2009) recommends that children 6–18 years of age and caregivers of infants be included as initial target groups for the new pandemic (H1N1) 2009 vaccine (27), which may reduce household transmission. As pandemic (H1N1) 2009 continues to spread internationally, ongoing investigations are needed to shed further light on transmission dynamics, to monitor epidemiologic changes over time, and to assess the effectiveness of public health interventions.

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#### References

- World Health Organization. Influenza A (H1N1)—update 52. Laboratory-confirmed cases of new influenza A (H1N1) as officially reported to WHO by States Parties to the International Health Regulations (2005) [cited 2009 Jun 22]. http://www.who.int/csr/ don/2009\_06\_22/en/index.html
- Centers for Disease Control and Prevention. Update: swine influenza A (H1N1) infections—California and Texas, April 2009. MMWR Morb Mortal Wkly Rep. 2009;58:435–7.
- World Health Organization. Transcript of statement by Margaret Chan, Director-General of the World Health Organization. 11 June 2009 [cited 2009 Jul 7]. http://www.who.int/mediacentre/influenzaAH1N1 presstranscript 20090611.pdf
- Fox JP, Cooney MK, Hall CE, Foy HM. Influenza virus infections in Seattle families, 1975–1979. II. Pattern of infection in invaded households and relation of age and prior antibody to occurrence of infection and related illness. Am J Epidemiol. 1982;116:228–42.
- Fox JP, Hall CE, Cooney MK, Foy HM. Influenzavirus infections in Seattle families, 1975–1979. I. Study design, methods and the occurrence of infections by time and age. Am J Epidemiol. 1982;116:212– 27.
- Foy HM, Cooney MK, Allan I. Longitudinal studies of types A and B influenza among Seattle schoolchildren and families, 1968–74. J Infect Dis. 1976;134:362–9.
- Taber LH, Paredes A, Glezen WP, Couch RB. Infection with influenza A/Victoria virus in Houston families, 1976. J Hyg (Lond). 1981;86:303–13.
- Texas Department of State Health Services. Health Service Region
  2009 [cited 2009 Jun 22]. http://www.dshs.state.tx.us/region8/ default.shtm
- Centers for Disease Control and Prevention. Interim guidance on case definitions to be used for investigations of novel influenza A (H1N1) cases. 2009 [cited 2009 Jun 27]. http://www.cdc.gov/ H1N1flu/guidance/
- Carrat F, Vergu E, Ferguson NM, Lemaitre M, Cauchemez S, Leach S, et al. Time lines of infection and disease in human influenza: a review of volunteer challenge studies. Am J Epidemiol. 2008;167:775– 85. DOI: 10.1093/aje/kwm375
- Lessler J, Reich NG, Brookmeyer R, Perl TM, Nelson KE, Cummings DA. Incubation periods of acute respiratory viral infections: a systematic review. Lancet Infect Dis. 2009;9:291–300. DOI: 10.1016/S1473-3099(09)70069-6
- 12. Centers for Disease Control and Prevention. CDC protocol of realtime RT-PCR for influenza A (H1N1) 2009 [cited 2009 Jun 23]. http:// www.who.int/csr/resources/publications/swineflu/realtimeptpcr/ en/index.html

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- Altman DG. Practical statistics for medical research. London: Chapman and Hall/CRC Press; 1999.
- Novel Swine-Origin Influenza A (H1N1) Virus Investigation Team, Dawood FS, Jain S, Finelli L, Shaw MW, Lindstrom S, Garten RJ, et al. Emergence of a novel swine-origin influenza A (H1N1) virus in humans. N Engl J Med. 2009;360:2605–15. DOI: 10.1056/ NEJMoa0903810
- Centers for Disease Control and Prevention. Update: novel influenza A (H1N1) virus infection—Mexico, March–May, 2009. MMWR Morb Mortal Wkly Rep. 2009;58:585–9.
- Health Protection Agency; Health Protection Scotland; National Public Health Service for Wales; HPA Northern Ireland Swine influenza investigation teams. Epidemiology of new influenza A (H1N1) virus infection, United Kingdom, April–June 2009. Euro Surveill. 2009;14. pii: 19232.
- Cowling BJ, Fang VJ, Riley S, Malik Peiris JS, Leung GM. Estimation of the serial interval of influenza. Epidemiology. 2009;20:344–7. DOI: 10.1097/EDE.0b013e31819d1092
- Woodall J, Rowson KE, Mc DJ. Age and Asian influenza, 1957. BMJ. 1958;2:1316–8. DOI: 10.1136/bmj.2.5108.1316
- Hayden FG, Belshe R, Villanueva C, Lanno R, Hughes C, Small I, et al. Management of influenza in households: a prospective, randomized comparison of oseltamivir treatment with or without postexposure prophylaxis. J Infect Dis. 2004;189:440–9. DOI: 10.1086/381128
- Welliver R, Monto AS, Carewicz O, Schatteman E, Hassman M, Hedrick J, et al. Effectiveness of oseltamivir in preventing influenza in household contacts: a randomized controlled trial. JAMA. 2001;285:748–54. DOI: 10.1001/jama.285.6.748
- Blumenfeld HL, Kilbourne ED, Louria DB, Rogers DE. Studies on influenza in the pandemic of 1957–1958. I. An epidemiologic, clinical and serologic investigation of an intrahospital epidemic, with a note on vaccination efficacy. J Clin Invest. 1959;38:199–212. DOI: 10.1172/JCI103789

- Centers for Disease Control and Prevention. Serum cross-reactive antibody response to a novel influenza A (H1N1) virus after vaccination with seasonal influenza vaccine. MMWR Morb Mortal Wkly Rep. 2009;58:521–4.
- Cate TR, Kasel JA, Couch RB, Six HR, Knight V. Clinical trials of bivalent influenza A/New Jersey/76-A/Victoria/75 vaccines in the elderly. J Infect Dis. 1977;136(Suppl):S518–25.
- Dolin R, Wise TG, Mazur MH, Tuazon CU, Ennis FA. Immunogenicity and reactogenicity of influenza A/New Jersey/76 virus vaccines in normal adults. J Infect Dis. 1977;136(Suppl):S435–42.
- Hayden FG, Gubareva LV, Monto AS, Klein TC, Elliot MJ, Hammond JM, et al. Inhaled zanamivir for the prevention of influenza in families. Zanamivir Family Study Group. N Engl J Med. 2000;343:1282–9. DOI: 10.1056/NEJM200011023431801
- Monto AS, Pichichero ME, Blanckenberg SJ, Ruuskanen O, Cooper C, Fleming DM, et al. Zanamivir prophylaxis: an effective strategy for the prevention of influenza types A and B within households. J Infect Dis. 2002;186:1582–8. DOI: 10.1086/345722
- Centers for Disease Control and Prevention. Use of influenza A (H1N1) 2009 monovalent vaccine: recommendations of the Advisory Committee on Immunization Practices (ACIP), 2009. National Center for Immunization and Respiratory Diseases. MMWR Recomm Rep. 2009;58(RR-10):1–8.

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