# Pandemic (H1N1) 2009 Outbreak at Camp for Children with Hematologic and Oncologic Conditions

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An outbreak of influenza A pandemic (H1N1) 2009 occurred among campers and staff at a summer camp attended by children with hematologic and oncologic conditions. The overall attack rate was 36% and was highest among children and adolescents (43%), persons with cancer (48%), and persons with sickle cell disease (82%).

Since it was first identified in April 2009 (1), the influenza A pandemic (H1N1) 2009 virus has sickened >1 million persons in the United States (www.cdc.gov/h1n1flu/surveillanceqa.htm). Because of the novelty of this virus, its transmissibility and severity are still under study.

We investigated an outbreak that occurred at a summer camp in northern rural Louisiana. Study approval was provided by the institutional review boards of Louisiana State University Health Sciences Center, Children's Hospital, and Tulane University Medical Center, New Orleans, Louisiana.

#### The Study

The camp opened July 26, 2009 (day 1), with 101 campers and 116 staff. Campers were children with hematologic or oncologic conditions and their nonaffected siblings. All participants were physically examined and questioned so that anyone with presence of or exposure to potentially communicable diseases could be excluded from attending camp. Campers (8–12/patrol) were grouped in age- and gender-specific patrols (B1–B4 and G1–G4) with assigned staff (4–5/patrol); they slept in bunkhouses and had common bathing facilities. All campers and staff dined together and shared various activities.

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On day 2, fever developed in a healthy camper in patrol G3 and promptly subsided. Fever also developed in a second camper (patrol B2) with sickle cell disease (SCD). This camper was seen at the local hospital, had a negative rapid influenza test (RIT) result, and was sent home. On day 3, fever and cough developed in 4 children (2 from B2); 1 was tested by RIT with a negative result. Evaluation of the entire B2 patrol found no one else symptomatic. On day 4, fever developed in1 child with cancer (B2) and 1 with SCD (G2 patrol); each had positive test results for influenza A virus.

The number of episodes of fever was considered excessive, and because additional campers reported having fever the camp was closed. At the time of dismissal, all campers and staff were provided masks and instructions about cough etiquette and handwashing; a dose of oseltamivir was also administered, and a prescription for oseltamivir was provided.

Ten days after the camp closed, all attendees were contacted (by email, telephone, and regular mail) to gather information about their outcomes. Clinical signs and symptoms of interest were fever (measured or subjective) or chills; cough or sore throat; muscle pain; and nausea, vomiting, or diarrhea. On the basis of an adaptation of the definition by the Centers for Disease Control and Prevention (www.cdc.gov/h1n1flu/clinicians), persons with fever or chills and symptoms in >2 other categories were classified as having influenza-like illness (ILI); persons with fever alone (without an explanation) and symptoms in 1 other category, or no documented fever but symptoms in >2 other categories, were classified as having probable ILI (P-ILI). A case-patient was defined as a person in whom ILI or P-ILI developed within 10 days of closing of the camp.

Questionnaires were returned by 88 (76%) and 77 (76%) of staff and campers, respectively. Mean age was 22.5 (range 14–69) years for staff and 10.5 (range 5–15) years for campers. Of 88 staff, 56 reported no chronic medical condition; a few reported cancer in remission (n = 7), SCD (n = 3), or other conditions (n = 22: 9 asthma, 2 inflammatory bowel disease, 2 unspecified, and 1 each with epilepsy, gall stones, chronic hepatitis C, immune thrombocytopenic purpura, Kartagener syndrome, mental retardation, porphyria, stroke, and type 1 diabetes mellitus). Of 77 campers, 45 (58%) reported no underlying illness, and a few reported cancer (n = 20: 12 leukemia, 2 lymphoma, 4 solid organ tumor, 2 unspecified; 2 were receiving maintenance therapy, the others had completed treatment), SCD (n = 8), or other (n = 4: 2 asthma, 1 immune thrombocy-)topenic purpura, and 1 cardiomyopathy).

Of 165 attendees who returned the questionnaire, 59 (38.5%) reported symptoms: fever (40, 68%), cough (30, 51%), sore throat (21, 36%), muscle pain (16, 27%), nau-

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## DISPATCHES

sea (10, 17%), diarrhea (9, 15%), vomiting (8, 14%), runny nose (6, 10%), and headache (4, 7%). Abdominal pain, weakness, earache, conjunctivitis, and joint stiffness were rare. Pain crisis developed in 2 patients with SCD. Twentyfive patients met the definition of ILI, and 34 met the definition of P-ILI (Figure). Symptoms were more common among campers than staff (46.8% vs. 26.1%; p = 0.009), persons of younger age (Table 1), and persons with cancer or SCD (Table 2). Proportions were compared with 2-tailed  $\chi^2$  or Fisher exact test by using Epi Info version 3.5 (www. cdc.gov/Epiinfo).

Probably because of small numbers, no difference was evident in the attack rate between staff assigned and unassigned to a patrol, between boy and girl campers, between patrols, or between those who recalled or did not recall being in contact with a case-patient. Twenty-six case-patients sought medical attention, and 12 (2 staff and 10 campers) were hospitalized; 8 had SCD and 4 were in active cancer treatment; all were regular-floor admissions; none required oxygen support or other intensive care; mean length of stay was 4.25 (range 3–6) days; and all recovered uneventfully.

Twenty-five persons (18 campers and 7 staff) were tested for influenza: 13 were tested at an outside facility, of whom 3 had positive results (method of testing unknown); 12 were tested at Children's Hospital by RIT (BinaxNOW; Invernass Medical, Waltham, MA, USA) and direct immunofluorescence assay (D<sup>3</sup> Ultra; Diagnostic Hybrids, Athens, OH, USA), and 5 (42%) had positive test results for influenza A virus. Of the 5 positive specimens obtained from patients, 4 were confirmed as pandemic (H1N1) 2009 virus by real-time reverse transcription–PCR.

### Conclusions

Limited preliminary information is available about summer camp outbreaks of pandemic (H1N1) 2009 (2). The closest scenarios (school outbreaks) reported attack rates of 3%–33% in the United States (3–5) and 2%–31%in the United Kingdom (6,7). Differences between studies and groups probably reflect different levels of exposure and surveillance definitions. In the outbreak reported here, the overall attack rate was 36%; infection was more common among campers (47%), children and adolescents (43%), those with cancer (48%), and those with SCD (82%). Because of their underlying condition, the threshold for evaluation and intervention is lower for children with hematologic or oncologic processes, which might account for some increased reporting of symptoms, hospitalization, and extended stay.

Children with SCD were disproportionately affected; 82% of them reported symptoms, and 89% of those symptomatic were hospitalized because of fever or pain crisis. For seasonal influenza, the hospitalization rate is 56× high-

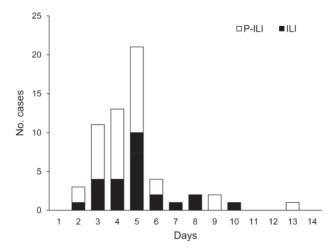


Figure. Outbreak curve of influenza-like illness (ILI) and probable ILI (P-ILI) cases during the 4 days of summer camp for children with hematologic and oncologic conditions (1–4) and the 10 days (5–14) after closing the camp, Louisiana, USA, 2009.

er for children with SCD than those without SCD (8); the same trend seems true for pandemic (H1N1) 2009 infection. Among those with no underlying condition, the influenza attack rate was lower (28%), few (11%) persons sought medical attention, and none were hospitalized. For persons with no underlying condition, the reported symptoms suggest an illness no more severe than seasonal influenza.

A few limitations should be recognized. First, 25% of attendees did not return the questionnaire. Second, we used a clinical definition of ILI; however, although the limitations of the definition are well recognized, it is a well-accepted tool for outbreak investigation. Third, data are based on recall and report by participants. Fourth, only a few persons were tested to determine the causative agent. However, of the 12 patients we tested at Children's Hospital, 5 (42%) had influenza A virus and 4 were confirmed as having pandemic (H1N1) 2009 virus infection. This level of positivity is in accordance with sensitivity reported for available tests (9,10) and suggests that most, if not all, cases of ILI identified were caused by pandemic (H1N1) 2009 virus.

Table 1. Attack rates of influenza-illness according to age and
underlying medical condition at summer camp for children with
hematologic and oncologic conditions, Louisiana, USA, 2009*
Underlying condition, no. positive/no. tested (%)

Age group, y	No†	Yes‡	All§					
5–15	19/54 (35.2)	18/33 (54.5)	37/87 (42.5)					
16–30	10/41 (24.4)	11/28 (39.3)	21/69 (30.4)					
31–69	0/6 (0)	1/3 (33.3)	1/9 (11.1)					
Total	29/101 (28.7)	30/64 (46.9)	59/165 (35.8)					
*By $\chi^2$ test for trend for increasing age groups within each category. to = 0.049.								

‡p = 0.255

§p = 0.027.

Table 2. Risk for influenza and hospitalization according to category of underlying condition at summer camp for children with
hematologic and oncologic conditions, Louisiana, USA, 2009*

Underlying condition	Risk for influenza symptoms			Risk for hospitalization, if symptomatic		
	Staff	Campers	All	Staff	Campers	All
None	10/56 (17.9)	19/45 (42.2)	29/101 (28.7)	0/10 (0)	0/19 (0)	0/29 (0)
Other than cancer or sickle cell disease	7/22 (31.8)	1/4 (25.0)	8/26 (30.8)	0/7 (0)	0/1 (0)	0/8 (0)
Cancer	4/7 (57.1)†	9/20 (45.0)	13/27 (48.1)†	0/4 (0)	4/9 (44.4)‡	4/13 (30.8)‡
Sickle cell disease	2/3 (66.6)†	7/8 (87.5)†	9/11 (81.8)‡	2/2 (100.0)†	<b>6/7 (85.7)</b> §	<b>8/9 (88.9)</b> §
Total	23/88 (26.1)	36/77 (46.8)	59/165 (35.8)	2/23 (8.7)	10/36 (27.8)	12/59 (20.3

\*Values given as no. in category/total no. with condition (%). **Boldface** indicates significance. Data were analyzed by using 2-tailed Fisher exact test for each category compared with persons with no underlying condition.

†p = 0.01–0.05. ‡p = 0.0001–0.01.

\$p<0.0001.

sp<0.0001.

Dr Morrison is on faculty in pediatric hematology-oncology at Louisiana State University Health Sciences Center and Children's Hospital in New Orleans, Louisiana, USA. Her research interests include infections in immunocompromised children.

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