
Person-to-Person Transmission of Nipah Virus in a Bangladeshi Community

Emily S. Gurley,* Joel M. Montgomery,† M. Jahangir Hossain,* Michael Bell,† Abul Kalam Azad,‡ Mohammed Rafiqul Islam,‡ Mohammed Abdur Rahim Molla,‡ Darin S. Carroll,† Thomas G. Ksiazek,† Paul A. Rota,† Luis Lowe,† James A. Comer,† Pierre Rollin,† Markus Czub,§¶ Allen Grolla,§ Heinz Feldmann,§¶ Stephen P. Luby,*† Jennifer L. Woodward,# and Robert F. Breiman*†

An encephalitis outbreak was investigated in Faridpur District, Bangladesh, in April–May 2004 to determine the cause of the outbreak and risk factors for disease. Biologic specimens were tested for Nipah virus. Surfaces were evaluated for Nipah virus contamination by using reverse transcription–PCR (RT-PCR). Thirty-six cases of Nipah virus illness were identified; 75% of case-patients died. Multiple peaks of illness occurred, and 33 case-patients had close contact with another Nipah virus patient before their illness. Results from a case-control study showed that contact with 1 patient carried the highest risk for infection (odds ratio 6.7, 95% confidence interval 2.9–16.8, $p < 0.001$). RT-PCR testing of environmental samples confirmed Nipah virus contamination of hospital surfaces. This investigation provides evidence for person-to-person transmission of Nipah virus. Capacity for person-to-person transmission increases the potential for wider spread of this highly lethal pathogen and highlights the need for infection control strategies for resource-poor settings.

Nipah virus was first identified as the pathogen responsible for outbreaks of encephalitis in Malaysia and Singapore from October 1998 to June 1999 (1–6). Fever (97%), headache (61%), and reduced consciousness (55%) were the most common symptoms in Malaysia; case-fatality rate was 40% (7). Most case-patients lived on pig farms

(95% in Malaysia) (1) or worked in abattoirs (100% in Singapore) (4,8). Serologic and reverse transcription–PCR (RT-PCR) testing of blood and urine from pteropid fruit bats in Malaysia and Cambodia showed Nipah virus infection, which suggested that these animals were reservoir hosts (9–11). During this outbreak, Nipah viruses were also isolated from human respiratory secretions and urine (2); however, 2 studies did not find evidence of nosocomial transmission (12,13).

Subsequent investigations in India and Bangladesh have suggested that Nipah virus may have been transmitted from person to person. During an outbreak in Siliguri, India, in 2001, 45 (75%) of 60 patients, many of them health-care workers, had a history of hospital exposure to patients infected with Nipah virus (14). A case-control study conducted during an outbreak in Meherpur District, Bangladesh, in 2001 showed that persons who lived with or cared for patients during the patient's illness were more likely to become infected with Nipah virus, and patients were more likely to have reported touching secretions of other patients; however, this finding could not be differentiated from common environmental exposures (15). During an outbreak in Rajbari District, Bangladesh, in January 2004, case-patients were more likely than controls to have had contact with another patient with Nipah virus illness (16). Pteropid bats were also suspected to be the reservoir for the virus in Bangladesh (9–11,15).

On April 5, 2004, the ICDDR,B and the Institute for Epidemiology and Disease Control Research were alerted to a cluster of 5 persons with fever, headache, confusion, and loss of consciousness in Faridpur District, in western Bangladesh. Nipah virus was the suspected cause of the

*ICDDR,B, Dhaka, Bangladesh; †Centers for Disease Control and Prevention, Atlanta, Georgia, USA; ‡Ministry of Health and Family Welfare, Dhaka, Bangladesh; §Public Health Agency of Canada, Winnipeg, Manitoba, Canada; ¶University of Manitoba, Winnipeg, Manitoba, Canada; and #University of Texas School of Public Health, Houston, Texas, USA

outbreak and an investigation began on April 6, 2004. Investigation goals were to identify a reservoir host(s), define the magnitude of the epidemic, and determine the principal modes of transmission. This report addresses the last 2 goals.

Methods

Finding and Defining Cases

We defined suspected case-patients with Nipah virus illness as persons with fever and altered mental status (serious illness) residing or working in the outbreak area or persons who had fever and cough or headache (mild illness) and who were contacts of patients with Nipah virus infection or resided in the outbreak area. Suspected case-patients were identified by visiting area hospitals, conducting door-to-door visits to all homes in the affected area, and tracing contacts of patients with Nipah virus illness. A history of illness and general information about exposures were obtained for each suspected case-patient. Friends and relatives of deceased case-patients served as proxy informants for interviews, and guardians were included in interviews of children <13 years of age. All those who died in the outbreak areas during this time were considered suspected case-patients.

A probable case-patient was defined as a patient with fever and mental status changes who lived or worked in the same village as a confirmed case-patient and from whom either serum or cerebrospinal fluid (CSF) was not available (i.e., because the patient died before specimen could be collected) or from whom a negative result was obtained from a sample collected <10 days after illness onset but collection of subsequent specimens was impossible (17). A laboratory-confirmed case of Nipah virus infection was defined by evidence of acute infection shown by immunoglobulin M (IgM) to Nipah virus in serum or CSF. To evaluate the possibility of asymptomatic infections, we asked persons with a history of close contact with a patient with Nipah virus-like illness to provide a blood specimen for serologic testing (7,13).

Specimen Collection and Laboratory Testing

Acute-phase blood specimens, throat swabs, saliva, and urine were collected from persons with suspected cases. When possible, hospitalized patients underwent lumbar puncture and chest radiography. Convalescent-phase blood specimens were collected from all persons with suspected cases >10 days after illness onset. Acute- and convalescent-phase serum and CSF were tested with an IgM capture enzyme immunoassay (EIA) for IgM and an indirect EIA for IgG by using Nipah virus (Malaysian prototype) antigen S (18) at the Centers for Disease Control and Prevention in Atlanta, Georgia, USA. Acute-phase serum, CSF, throat

swabs, saliva, and urine were also tested by RT-PCR for viral RNA. RNA was extracted from specimens by using the acid guanidinium-phenol method (19). RT-PCR was performed by using a primer set to detect the nucleocapsid gene as described (20), the Superscript One Step RT-PCR Kit (Invitrogen, Carlsbad, CA, USA), and standard reaction conditions (21). Primers used were NVNF-4: 5'-GGA GTT ATC AAT CTA AGT TAG-3' and NVNBR4: 5'-CAT AGA GAT GAG TGT AAA AGC-3'. PCR products were subjected to electrophoresis on 2% agarose gels and visualized by staining with ethidium bromide. Positive PCR results were confirmed by sequence analysis of PCR products.

Case-Control Study

We conducted a case-control study to identify risk factors for transmission of Nipah virus infection. Persons meeting either the probable or confirmed case definition were enrolled as case-patients. All healthy persons including household members and neighbors (residing within 150 m of a case household), were eligible for participation as control participants. Controls were randomly selected from a list of names generated by a census of all households in the affected community and matched to a case-patient (6:1) by sex and age (± 2 years; all case-patients and controls were >4 years of age). They were given a 2-part questionnaire in Bengali. The first part focused on environmental exposures and established whether the participant had had contact with a specific probable or confirmed case-patient(s). The second part focused on types of contact with specific case-patients to elucidate possible modes of transmission from person to person. Proxy interviews were conducted with guardians and companions for case-patients who had died or who were unable to respond and for all case-patients and controls <13 years of age.

We calculated odds ratios (ORs) and 95% confidence intervals (CIs) by using conditional univariate logistic regression that accounted for matched enrollment of case-patients and controls (22). To evaluate independent risk factors, we tested all variables with a *p* value <0.1 from univariate analyses in conditional stepwise forward multivariate logistic regression. Associations were considered statistically significant if *p* value was <0.05. All statistical analyses were performed with SAS version 9.0 (SAS Institute Inc., Cary, NC, USA).

Environmental Study

In early May 2004, 5 weeks after the outbreak was first recognized, environmental surfaces believed to have a high risk for contamination with Nipah virus were selected for sampling. These included surfaces within hospitals where Nipah virus patients received care, surfaces within homes of confirmed case-patients, trees where bats foraged, date

palm sap collection pots, and fruits that may have been in contact with fruit bats. Surfaces were rolled with sterile, cotton-tipped applicators, which were stored in 500 μ L Dulbecco modified Eagle medium supplemented with an antibiotic-antimycotic solution. Viral RNA was extracted from 140 μ L of the resulting suspension by using a Viral RNA Minikit (QIAGEN, Mississauga, Ontario, Canada) and eluted in a volume of 50 μ L. Nipah virus RNA was detected by using the LightCycler RNA Amplification Kit SYBR Green I (Roche, Laval, Quebec, Canada) with primers for the nucleoprotein gene (NPF: 5'-ATCAATCGTG-GTTATCTTGAAC-3' and NPR: 5'CCTCTTCGTCGA-CATCTTGATC-3') and with thermocycling and real-time detection performed on a SmartCycler II reaction block (Cepheid, Sunnyvale, CA, USA). Positive results were later confirmed by direct sequencing of amplified products, and sequences were compared with those obtained from patient samples (23).

Participants and Ethical Considerations

Because this was an outbreak investigation, protocols did not undergo formal institutional review. The Bangladesh Ministry of Health and Family Welfare requested this investigation and reviewed and approved all protocols. Participation in these studies was strictly voluntary and informed consent was obtained from all participants; for those <18 years of age, individual and parental consent were obtained.

Results

Defining Cases

We identified and collected specimens from 210 suspected case-patients, of whom 32 had fever and altered mental status. Thirty-six case-patients were identified, including 4 who had a febrile illness without altered mental status. Twenty-three (64%) were laboratory confirmed and 13 (36%) were classified as probable; 27 (75%) died. Serum specimens were available from 27 of 36 case-patients, 4 of which had no detectable antibodies to Nipah virus. These specimens were collected <6 days after illness onset. All 4 patients died before a second specimen could be obtained; therefore, they were classified as probable case-patients. Nine patients who had encephalitis-like illnesses and who resided in the outbreak area died before diagnostic specimens could be collected; they were also classified as probable case-patients. No asymptomatic cases of Nipah virus infection were documented in contacts of Nipah virus patients who consented to provide a blood specimen ($n = 20$). Results of sequencing RT-PCR products from throat swab, saliva, and urine samples of 9 patients were consistent with serologic data, which indicated that Nipah virus was the etiologic agent of this outbreak (23).

Contact Tracing

Probable and confirmed case-patients were identified in 7 villages in Faridpur District; dates of onset of illness ranged from February 19 to April 17, 2004 (Figure 1). Thirty-three (92%) of the 36 case-patients had had close contact with another ill person before they became ill (Figure 2). Five cases (patients A, B, F, G, and EE) appeared to be associated with secondary and tertiary person-to-person transmission of Nipah virus. We present some examples of close contact.

Four of Patient A's caregivers who resided in his village became ill after their contact with him; they were his mother (patient D), his son (patient E), his aunt (patient B), and a neighbor (patient C). They became ill 15–27 days after patient A became ill. Patient B received care from her brother (patient F), who lived in a village \approx 30 minutes from her village and who became ill 13 days after patient B.

Patient F became ill after he had returned to his village. As shown in Figure 2, 22 (61%) of 36 cases in this outbreak had contact with patient F before they became ill. Patient F was a local religious leader. Many persons in his family and his followers had close contact with him during his illness. Eight (80%) members of his 10-person household became infected with Nipah virus 6–13 days after his illness onset. Two of patient F's brothers (patients T and U), both of whom lived \approx 2 hours away, visited him for 6 hours on the day he died. They developed serious illness 7 (patient T) and 11 days (patient U) days later, and both died. Patients Y (patient F's daughter) and Z (patient F's son-in-law), who lived 1 hour away from patient F, became ill \approx 1 week after 1 multiple-hour visit with patient F late during his illness; this was their only reported contact with a Nipah virus patient. In 11 other close contacts of patient F, including family and religious followers, Nipah virus infection developed 6–14 days after his illness onset.

Patient G, a follower of patient F, moved to his family's house in an adjacent village to receive care after becoming ill. Patient H became ill \approx 9 days after physically supporting patient G while walking to patient G's family's house. A rickshaw driver, patient I, who helped carry

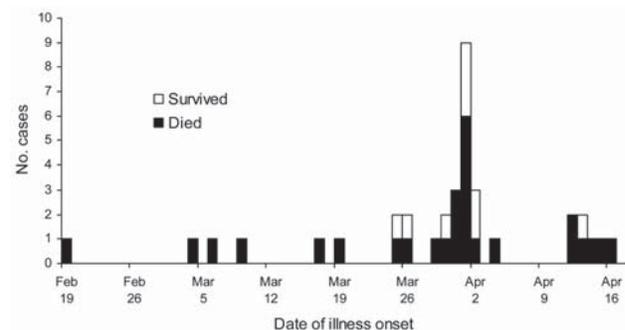


Figure 1. Dates of illness onset during a Nipah virus outbreak in Faridpur District, Bangladesh, 2004.

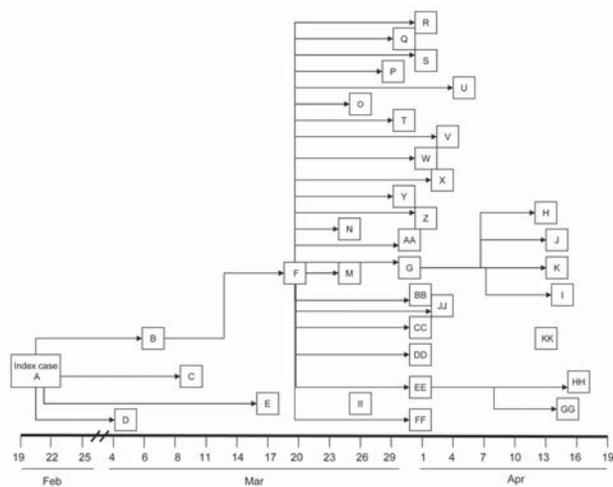


Figure 2. Chain of person-to-person transmission with dates of onset of illness during a Nipah virus outbreak, Faridpur District, Bangladesh, 2004. Letters identify individual patients. Patients KK and II had no known contact with any ill patient before their illness.

and transport patient G to the hospital as his condition deteriorated, became sick 10 days after this exposure and later died. Patient G's father (patient J) and sister-in law (patient K) cared for him during his illness; both became severely ill 2 weeks after patient G's illness onset; only patient K survived.

Patient GG visited his wife and daughter, both hospitalized as suspected Nipah virus patients, at the local healthcare facility. Patient GG spent the night in the hospital and shared a bed with patient EE (a common practice in Bangladesh), who was a male friend, and a suspected Nipah virus case-patient. Although tests on patient GG's wife and daughter did not detect Nipah virus infection, patient EE's infection was laboratory confirmed. Patient GG became ill 10 days after this contact and died.

Patients II and KK had no known contact with any ill patient before their illness and were distinct from other case-patients in that they were not friends, relatives, or followers of patient F and lived outside the affected villages. No cases of Nipah virus illness among healthcare workers were reported to authorities during this outbreak.

Case-Control Study

Thirty-four of the 36 case-patients were enrolled in the study and matched to 6 controls each ($n = 204$) by age and sex (Table). Two case-patients (patients JJ and KK; Figure 2) were not included in the study because of logistic constraints. Patient JJ was away from her home when the questionnaire was administered. Patient KK was a policeman who had 2 residences and traveled frequently while on duty; we were therefore unable to identify appropriate proxies for his interview.

Ten variables were significantly associated with Nipah virus infection in univariate analysis (online Appendix Table 1, available from www.cdc.gov/EID/content/13/7/1031-appT1.htm). Having had close contact (touching or receiving a cough or sneeze in the face) with patient F placed a person at greatest risk of acquiring Nipah virus infection (OR 6.7, 95% CI 2.9–16.8, $p < 0.001$). Having had any contact with someone who later died, had a fever, was unconscious, or had respiratory difficulties was also associated with illness. Having avoided any contact with someone who later died was negatively associated with illness. Having had a household member harvest date palm sap was the only environmental exposure associated with an increased risk for infection. Having visited the home village of the index case-patient was also associated with illness. However, the only exposure variable that remained significant in multivariate analysis was having had contact with patient F (OR 5.6, 95% CI 1.79–17.24, $p = 0.003$).

Univariate analysis of risk factors specifically associated with types of contact with patient F was conducted. Fourteen of 42 exposure variables were associated with illness (online Appendix Table 2, available from www.cdc.gov/EID/content/13/7/1031-appT2.htm). Having had close body contact and having spent longer periods of time with patient F were associated with illness. Having kept a certain distance from patient F and having washed hands after contact with him were negatively associated with illness (online Appendix Table 2). Despite multivariate analysis of risk factors associated with type of contact with patient F, insufficient sample size ($n = 50$) resulted in overfitting of the model and spurious results.

Environmental Study

A total of 468 environmental specimens were collected by swabbing; 137 from walls, bed frames, mattresses, and floors of 2 Faridpur hospitals; 57 from surfaces and utensils of case-patient residences; 150 from trees where bats forage and from fruits; 98 from bat excreta; and 26 from other sites. Eleven positive specimens were collected from

Table. Characteristics of 34 case-patients infected with Nipah virus in case-control study, Bangladesh, April–May 2004

Characteristic	No. (%) case-patients
Sex	
Male	20 (58.8)
Female	14 (41.2)
Age group, y	
1–15	4 (11.8)
16–24	1 (2.9)
25–40	19 (55.9)
41–60	9 (26.5)
>60	1 (2.9)
Adults vs. children	
≤15 y of age (children)	4 (11.8)
>15 y of age (adults)	30 (88.2)

the surrounding wall and bed frame where a confirmed case-patient (patient Z) had been hospitalized on April 6, \approx 5 weeks before environmental samples were collected. No other patients with encephalitis were known to have used that bed after patient Z's hospitalization. The wall and bed frame were visibly soiled, and hospital staff reported that they had not been cleaned since the outbreak. Samples from these areas showed evidence of Nipah virus RNA. Sequences of PCR products were identical to sequences of Nipah viruses isolated from patient HH during the outbreak (18). No samples from case-patient residences, bat-feeding sites, or fruits were positive.

Discussion

This investigation provides compelling evidence for person-to-person transmission of Nipah virus. Exposure histories of infected patients and the epidemiologic curve, which demonstrates multiple peaks of illness onset during this outbreak, suggest that Nipah virus was transmitted by person-to-person contact. Contact tracing documented Nipah virus illness after brief, yet close contact, with other persons infected with Nipah virus. Findings from the case-control study, which showed a 6-fold increased risk for infection for those who reported contact with patient F, a negative association with illness after handwashing, and specific exposures to ill persons linked to transmission, confirm that exposure to ill persons spread the outbreak.

Person-to-person transmission of Nipah virus is biologically plausible. Other paramyxoviruses that infect humans, including human parainfluenza viruses 1–4, measles virus, and mumps virus, are also transmitted from person to person. Nipah virus has been isolated from human respiratory secretions, including those of cases from this outbreak (2,23). Furthermore, we identified that direct exposure to respiratory secretions of patients with Nipah virus illness was associated with infection during this outbreak.

The number of villages affected by the outbreak increased as persons traveled in and out of the affected areas to visit family members. This movement led to new infections in caregivers from other villages and increased the number of villages affected. Similar to transmission of severe acute respiratory syndrome (24), transmission of Nipah virus infection was not associated with all case-patients; however, 1 case-patient, patient F, was associated with 22 subsequent Nipah virus infections. Although host biologic factors may have resulted in increased viral shedding, leading to higher attack rates, the social status of patient F in the community enabled closer contact with more persons during his illness and more opportunity to transmit infection.

During the outbreak in Siliguri, India, 33 healthcare workers and hospital visitors became ill after exposure to hospitalized patients with Nipah virus illness, suggesting

nosocomial infection (14). In Malaysia and Singapore, contact with pigs was associated with infection; healthcare worker studies showed that the risk for nosocomial transmission was low (6,8,12,13,25). Absence of person-to-person transmission in Malaysia and Singapore could be due to differences in patient care practices, host susceptibility factors, or strain variation (23).

Detection of Nipah virus RNA on hospital surfaces demonstrates that infected patients shed virus into the environment, which could provide an opportunity to transmit Nipah virus to others. However, how long the virus will remain infectious in the environment is not known, and no evidence from this investigation indicates that type of transmission occurred. Despite the absence of healthcare worker infection during this outbreak, enhanced infection control practices, such as patient isolation and use of gloves and masks, likely had little protective effect because they were not implemented until late in the outbreak. This outbreak provides evidence that 1 person (patient GG) was infected during a hospital visit while sharing a bed with a confirmed case-patient. Nosocomial transmission of Nipah virus was reported during the outbreak in Siliguri, India (14). Efforts are needed to develop and disseminate reasonable guidelines for infection control and prevention for healthcare facilities and communities in resource-poor settings, especially when one considers our finding that handwashing prevents disease transmission.

Transmission of Nipah viruses to humans during this outbreak appears to have been bimodal. Fruit bats continue to be the only identified primary reservoir for the virus (9–11,15). In contrast with the Malaysia and Singapore experience, no intermediate hosts have been identified in Bangladesh (15). During this outbreak, the introduction of virus into human(s) from an unknown initial source appears to have been followed by person-to-person transmission. Three case-patients had no known contact with a sick patient before onset of illness; these case-patients may have been infected through exposure to virus-contaminated bat saliva, urine, or feces or through contact with some unknown intermediate host. These cases provide further evidence that sporadic cases of Nipah virus infection continue to occur in Bangladesh (26).

Selection of probable case-patients in this study could have been biased toward finding person-to-person transmission because probable case-patients by definition lived in the same area as confirmed case-patients, which increased the likelihood that they had had contact with one another. To rule out the possibility that this had an effect on our case-control study findings, we analyzed our data by using only confirmed cases; there were no major differences in findings compared with analysis that used confirmed and probable cases. Our findings are also limited by recall bias. Family and friends were often asked to provide information

about a deceased patient weeks after their illness (data for 5 patients were collected >1 month after illness onset). When possible, medical records were used to supplement patient reports of illness history, but often these records were incomplete or nonexistent. However, the investigation began just days after onset of illness in most patients, which provided for optimal recall of events. In addition, case-patients or their proxies might have more carefully considered their exposures than controls. However, all community members were aware of, and concerned by, the outbreak, and we believe it is unlikely that any control did not remember their exposure to case-patients in their community.

Capacity for person-to-person transmission increases the risk for wider spread of this highly lethal pathogen. In an impoverished, densely populated country such as Bangladesh, a lethal virus could rapidly spread before effective interventions are implemented. This spread would provide the seed for a substantial regional or global public health problem and highlights the need for local surveillance, outbreak detection and response, and rapid laboratory diagnostics. Sustained, long-term research is needed to characterize the reservoir of the virus and mechanisms for animal-to-animal, animal-to-human, and human-to-human transmission; clarify climatologic and other environmental factors linked to transmission; and define viral epitopes potentially linked to virulence and transmission. Effective infection control practices appropriate for resource-constrained healthcare systems and communities are urgently needed.

Acknowledgments

We gratefully acknowledge the invaluable comments and suggestions of L. Blum and the expert technical contributions of J. Mills, J. Betts, D. Cannon, K. Dubois, T. Stevens, and P. Stockton. We also thank the staff of Faridpur Medical College Hospital and the Civil Surgeon of Faridpur District for support during this investigation.

This study was supported by the US Centers for Disease Control and Prevention, the Public Health Agency of Canada, and the Institute for Epidemiology, Disease Control and Research, Bangladesh.

Ms Gurley works with the Program on Infectious Diseases and Vaccine Sciences at the ICDDR,B, and has participated in 3 Nipah virus outbreak investigations in Bangladesh. Her research interests include infectious disease and disease prevention and control in resource-poor settings.

References

1. Chua KB, Goh KJ, Wong KT, Kamarulzaman A, Tan PS, Ksiazek TG, et al. Fatal encephalitis due to Nipah virus among pig-farmers in Malaysia. *Lancet*. 1999;354:1257–9.
2. Chua KB, Lam SK, Goh KJ, Hooi PS, Ksiazek TG, Kamarulzaman A, et al. The presence of Nipah virus in respiratory secretions and urine of patients during an outbreak of Nipah virus encephalitis in Malaysia. *J Infect*. 2001;42:40–3.
3. Lam SK, Chua KB. Nipah virus encephalitis outbreak in Malaysia. *Clin Infect Dis*. 2002;34(Suppl 2):S48–51.
4. Paton NI, Leo YS, Zaki SR, Auchus AP, Lee KE, Ling AE, et al. Outbreak of Nipah-virus infection among abattoir workers in Singapore. *Lancet*. 1999;354:1253–6.
5. Chua KB. Nipah virus outbreak in Malaysia. *J Clin Virol*. 2003;26:265–75.
6. Parashar UD, Sunn LM, Ong F, Mounts AW, Arif MT, Ksiazek TG, et al. Case-control study of risk factors for human infection with a new zoonotic paramyxovirus, Nipah virus, during a 1998–1999 outbreak of severe encephalitis in Malaysia. *J Infect Dis*. 2000;181:1755–9.
7. Goh KJ, Tan CT, Chew NK, Tan PS, Kamarulzaman A, Sarji SA, et al. Clinical features of Nipah virus encephalitis among pig farmers in Malaysia. *N Engl J Med*. 2000;342:1229–35.
8. Chew MH, Arguin PM, Shay DK, Goh KT, Rollin PE, Shieh WJ, et al. Risk factors for Nipah virus infection among abattoir workers in Singapore. *J Infect Dis*. 2000;181:1760–3.
9. Yob JM, Field H, Rashdi AM, Morrissy C, van der Heide B, Rota P, et al. Nipah virus infection in bats (order Chiroptera) in peninsular Malaysia. *Emerg Infect Dis*. 2001;7:439–41.
10. Chua KB, Koh LC, Hooi PS, Wee KF, Khong JH, Chua BH, et al. Isolation of Nipah virus from Malaysian Island flying-foxes. *Microbes Infect*. 2002;4:145–51.
11. Olson JG, Rupprecht C, Rollin PE, An US, Niezgodna M, Clemins T, et al. Antibodies to Nipah-like virus in bats (*Pteropus lylei*), Cambodia. *Emerg Infect Dis*. 2002;8:987–8.
12. Mounts AW, Kaur H, Parashar U, Ksiazek TG, Cannon D, Arokiasamy JT, et al. A cohort study of health care workers to assess nosocomial transmissibility of Nipah virus, Malaysia, 1999. *J Infect Dis*. 2001;183:810–3.
13. Chan KP, Rolling PE, Ksiazek TG, Leo YS, Goh KT, Paton NI, et al. A survey of Nipah virus infection among various risk groups in Singapore. *Epidemiol Infect*. 2002;128:93–8.
14. Chadha MS, Comer JA, Lowe L, Rota PA, Rollin PE, Bellini WJ, et al. Nipah virus-associated encephalitis outbreak, Siliguri, India. *Emerg Infect Dis*. 2006;12:235–40.
15. Hsu VP, Hossain MJ, Parashar UD, Ali MM, Ksiazek TG, Kuzmin I, et al. Nipah virus encephalitis reemergence, Bangladesh. *Emerg Infect Dis*. 2004;10:2082–7.
16. Montgomery JM, Hossain MJ, Gurley E, Carroll DS, Croisier A, Bertherat E, et al. Risk factors for Nipah virus encephalitis in Bangladesh. *Emerg Infect Dis*. In press.
17. Ramasundrum V, Tan CT, Chua KB, Chong HT, Goh KJ, Chew NK, et al. Kinetics of IgM and IgG seroconversion in Nipah virus infection. *Neurologic Journal of Southeast Asia*. 2000;5:23–8.
18. Daniels P, Ksiazek T, Eaton BT. Laboratory diagnosis of Nipah and Hendra virus infections. *Microbes Infect*. 2001;3:289–95.
19. Chomczynski P, Sacchi N. Single-step method of RNA isolation by acid guanidinium thiocyanate-phenol-chloroform extraction. *Anal Biochem*. 1987;162:156–9.
20. Harcourt BH, Tamin A, Ksiazek TG, Rollin PE, Anderson LJ, Bellini WJ, et al. Molecular characterization of Nipah virus, a newly emergent paramyxovirus. *Virology*. 2000;271:334–49.
21. Katz RS, Premenko-Lanier M, McChesney MB, Rota PA, Bellini WJ. Detection of measles virus RNA in whole blood stored on filter paper. *J Med Virol*. 2002;67:596–602.
22. Kleinbaum DG, Klein M. Logistic regression: a self-learning text, 2nd ed. New York: Springer; 2002.
23. Harcourt BH, Lowe L, Tamin A, Liu X, Bankamp B, Bowden N, et al. Genetic characterization of Nipah virus, Bangladesh, 2004. *Emerg Infect Dis*. 2005;11:1594–7.

24. Anderson RM, Fraser C, Ghani AC, Donnelly CA, Riley S, Ferguson NM, et al. Epidemiology, transmission dynamics and control of SARS: the 2002–2003 epidemic. *Philos Trans R Soc Lond B Biol Sci.* 2004;359:1091–105.
25. Tan CT, Tan KS. Nosocomial transmissibility of Nipah virus. *J Infect Dis.* 2001;184:1367.
26. ICDDR,B. Nipah encephalitis outbreak over wide area of western Bangladesh, 2004. *Health and Science Bulletin.* 2004;2:7–11.

Address for correspondence: Emily S. Gurley, Program on Infectious Diseases and Vaccine Sciences, ICDDR,B, GPO 128, Mohakhali, Dhaka 1212, Bangladesh; email: egurley@icddr.org

**EMERGING
INFECTIOUS DISEASES**[®]
March 2006

**EID
Online**
www.cdc.gov/eid

**Search
past issues
EID
Online**
www.cdc.gov/eid

Appendix Table 1. Exposures and activities associated with Nipah virus infection, Bangladesh, April–May 2004*

Exposure or activity	No. (%) with reported exposure or activity		OR (95% CI)†	p value
	Case-patients (n = 34)	Controls (n = 204)		
Bat exposures				
Touched	0	4 (2)	1.1 (0.0–10.6)	1.00
Ate	0	1 (0.5)	6.0 (0.0–234.0)	1.00
Observed an increase in no. of fruit bats in or around house	0	4/202 (2)	1.14 (0.0–9.089)	1.00
Environmental exposures				
Ate date palm sap	11/31 (35)	102 (50)	0.51 (0.198–1.26)	0.163
Harvested date palm sap	2/33 (6)	3 (2)	4.0 (0.33–34.9)	0.30
Anyone in household harvested date palm sap	5 (15)	8/203 (4)	4.3 (1.0–17.1)	0.049‡
Touched or drank directly from a date palm sap collection bowl	6/33 (18)	16 (8)	2.5 (0.74–7.65)	0.15
Ate partially eaten fruit from ground	6 (18)	32 (16)	1.2 (0.35–3.28)	0.94
Travel history				
Visited village of patient F	26/32 (81)	156 (76)	1.3 (0.49–4.07)	0.747
Visited village of index case	15/33 (45)	34 (17)	4.04 (1.76–9.31)	0.0008‡
Exposure to human illness				
Touched person who was unconscious	19/32 (59)	74/200 (37)	2.6 (1.15–6.16)	0.019‡
Touched person who later died	23/31 (74)	73/197 (37)	5.5 (2.14–16.3)	0.0001‡
Touched person who had seizures	4/29 (14)	39/199 (20)	0.69 (0.16–2.29)	0.75
Touched person who had a fever	27/33 (82)	108/202 (53)	3.9 (1.51–12.01)	0.003‡
Touched person who had respiratory difficulties	19/32 (59)	65/197 (33)	3.3 (1.4–8.27)	0.0051‡
In the same room (not touching) with person who				
Was unconscious	24/33 (72)	120/200 (60)	1.9 (0.79–4.84)	0.183
Later died	25/32 (78)	112/200 (56)	3.1 (1.2–9.13)	0.0159‡
Had seizures	6/29 (21)	60/198 (30)	0.62 (0.19–1.66)	0.438
Had a fever	29 (85)	148/203 (73)	2.15 (0.77–7.43)	0.174
Had respiratory difficulties	20/33 (61)	111/200 (56)	1.19 (0.54–2.74)	0.77
No contact with person who				
Was unconscious	9/33 (27)	71/200 (36)	0.65 (0.25–1.55)	0.404
Later died	7/32 (22)	82/201 (41)	0.37 (0.13–0.96)	0.039‡
Had seizures	24/30 (80)	130/199 (65)	2.1 (0.76–6.52)	0.187
Had a fever	5 (15)	54/203 (27)	0.48 (0.14–1.33)	0.19
Had respiratory difficulties	13/33 (39)	88/201 (44)	0.87 (0.38–1.95)	0.87
Contact with a specific case(s)§¶				
Patient A	4 (12)	17 (8)	1.5 (0.34–4.8)	0.70
Patient B	8 (24)	8 (4)	6.5 (2.0–21.2)	0.001‡
Patient F	18 (53)	32 (16)	6.7 (2.9–16.8)	<0.0001‡#
Patient G	3 (9)	7 (3)	2.9 (0.43–15.1)	0.31
Patient EE	2 (6)	15 (7)	0.79 (0.09–3.5)	1.0
Patient II	0	5 (2)	0.89 (0.0–6.5)	0.93
Smoking history				
Currently smoke	13 (38)	61 (30)	1.9 (0.61–6.4)	0.32
Share cigarettes, biri, hukka, or pipe	10/33 (30)	45 (22)	1.9 (0.61–5.9)	0.31

*Values are no. subjects responding affirmatively/total no. responding (%) unless otherwise indicated. OR, odds ratio; CI, confidence interval.

†Exact method using univariate conditional logistic regression.

‡Included in multivariate conditional logistic regression analysis.

§Contact was defined as seeing, talking, eating or drinking, touching, and providing care.

¶Contact with ≥1 of the 34 probable or confirmed case-patients (selected few shown in table).

#Remained in final multivariate model.

Appendix Table 2. Exposures to patient F and association with Nipah virus infection, Bangladesh, April–May 2004*

Exposure	No. (%) with reported exposure or activity		OR (95% CI)†	p value
	Case-patients (n = 18)	Controls (n = 32)		
Contact with patient F				
During illness	16 (89)	31 (97)	0.27 (0.004–5.46)	0.583
Day of death	13 (72)	15 (47)	2.9 (0.74–12.90)	0.149
After death	17 (94)	22 (69)	7.5 (0.903–354.68)	0.069
Patient F activity				
Coughed while nearby	15/17 (88)	17/25 (68)	3.4 (0.559–38.15)	0.252
Unconscious while nearby	15 (83)	30/31 (97)	0.17 (0.003–2.365)	0.268
Vomited while nearby	8 (44)	9/30 (30)	1.8 (0.465–7.39)	0.481
Proximity to patient F				
Always >6 m	0 (0)	2 (6)	0.72 (0.0–9.518)	0.809
Always >1.5 m but <6 m	2 (11)	20 (63)	0.08 (0.008–0.427)	0.008
Always >0.5 m but <1.5 m	4 (22)	3 (9)	2.7 (0.398–21.04)	0.4015
Sometimes <30 cm (1 foot)	1 (6)	5 (16)	0.324 (0.006–3.267)	0.570
Sometimes <15 cm (0.5 feet)	7 (39)	3 (9)	5.9 (1.106–47.79)	0.035
Sometimes touching	6 (33)	1 (3)	14.6 (1.53–735.12)	0.013
Length of time spent with patient F				
<10 min	2 (11)	10 (31)	0.28 (0.026–1.59)	0.205
10–15 min	2 (11)	9 (28)	0.33 (0.03–1.89)	0.298
30 min	0	9 (28)	0.11 (0.0–0.766)	0.022
1–1.5 h	0	7 (22)	0.154 (0.0–1.122)	0.067
3–4 h	2 (11)	1 (3)	3.76 (0.183–235.8)	0.583
>4 h	16 (89)	6 (19)	31.2 (5.35–351.68)	<0.0001
Specific interactions with patient F				
Received sneeze/cough in face from patient	4/13 (31)	2 (6)	6.3 (0.766–80.81)	0.097
Ate with patient	4 (22)	2 (6)	4.2 (0.524–51.05)	0.228
Shared same plate/bowl/cup/glass	6 (33)	1 (3)	8.5 (0.749–449.8)	0.101
Fed patient with a spoon or cup	10 (56)	1 (3)	35.3 (4.064–999.9)	<0.0001
Fed patient with hands	7 (39)	1 (3)	18.5 (2.02–916.63)	0.004
Talked with patient	11 (61)	4 (13)	10.3 (2.25–59.31)	0.0011
Held patient's hands	13 (72)	6 (19)	10.6 (2.443–55.26)	<0.0006
Touched patient's face	12/17 (71)	4 (13)	15.5 (3.18–96.86)	0.0001
Shared a bed with patient	2 (11)	0	4.5 (0.341+ Inf)	0.25
Helped patient walk, sit, or stand	8 (44)	4 (13)	5.4 (1.15–30.18)	0.03
Lifted or carried patient	2 (11)	2 (6)	1.9 (0.12–27.8)	0.91
Cleaned patient's hands with a cloth or clothing	5 (28)	0	14.9 (1.89 + Inf)	0.0081
Cleaned patient's face with a cloth or clothing	4 (22)	0	11.0 (1.29 + Inf)	0.027
Wiped patient's face with hands	3 (17)	0	7.6 (0.775 + Inf)	0.083
Wiped patient's nose/mouth with hands	3 (17)	0	7.6 (0.775 + Inf)	0.083
Helped patient change clothes	1 (6)	0	1.78 (0.046 + Inf)	0.72
Helped patient use the toilet	2 (11)	0	4.5 (0.41+ Inf)	0.249
Cleaned feces from patient's body	0	0	Undefined	NA
Changed patient's bed linens	0	0	Undefined	NA
Washed patient's clothes	1 (6)	0	1.78 (0.046 + Inf)	0.72
Washed patient's bed linens	0	1 (3)	1.78 (0.0–69.3)	1.00
Dried out patient's mouth after death	0	1 (3)	2.3 (0.0–89.14)	1.00
Always/sometimes kept face covered while near patient	0	0	Undefined	NA
Activity after contact with patient F				
Washed hands	2/17 (12)	15/32 (47)	0.196 (0.03–0.895)	0.032

*Values are no. of subjects responding affirmatively/total no. responding (%) unless otherwise indicated. OR, odds ratio; CI, confidence interval; NA, not available.

†Exact method using univariate conditional logistic regression. + Inf, positive infinity, an unknown upper limit for the CI.