affected during this epidemic. Fourteen patients died; these deaths occurred in Kathmandu, the capital of Nepal, or in different cities in India after these patients were transferred there for better treatment. Fifty pregnant women had acute jaundice, but none of these women died.

The epidemic was presumed to be caused by consumption of contaminated water (3). In February and March 2014, water and sewerage pipelines were damaged in different areas of Biratnagar during construction and repair of roads. A survey conducted by the Department of Community Medicine, B.P. Koirala Institute of Health (Dharam, Nepal), found high levels of coliform bacteria in water supplies from different areas in Biratnagar during the epidemic. Tap water also looked cloudy and visibly contaminated (3).

To obtain more information about the epidemic, the incidence of acute jaundice was determined for 656 prisoners and 75 security personnel at the Biratnagar Jail. The study protocol was approved by the Liver Foundation Nepal. Informed consent was not obtained because identity of patient samples remained anonymous. Acute jaundice was detected among 30 (4.6%) prisoners and 4 (5.3%) security personnel. The same source of consumable water was used by the general population, inmates, and security personnel in Biratnagar.

To identify the causative agent of this epidemic, serum samples from 48 patients were obtained at Koshi Zonal Hospital, the largest government hospital in this zone. Hepatitis A virus RNA and IgM against hepatitis B virus core antigen were not detected in the 48 serum samples. Conversely, IgG, IgM, and IgA against HEV were detected in 47 (97.9%), 45 (94%), and 45 (94%) serum samples, respectively, and HEV RNA was detected in 42 (87.5%) of 48 serum samples, which indicated that the epidemic was caused by HEV.

A partial 412-nt sequence from open reading frame 2 corresponding to nt 5944–6355 of the HEV B1 genome (4) was obtained as reported (5). We obtained 40 HEV isolates from the 42 samples and sequenced partial 412-nt segments. All 40 HEV sequences from the epidemic in Biratnagar segregated into a cluster within genotype 1a (Figure). These sequences showed 99.8% nt identity with each other but only 90.8%–95.4% nt identity with other HEV isolates from Nepal and those from India, Bangladesh, Pakistan, and China.

Compared with previous HEV epidemics in Nepal (6) and other parts of the Indian subcontinent, the local government of Biratnagar and central government of Nepal took steps to contain the reported epidemic. Activities of public and private sectors in Biratnagar ended the epidemic in ≈12 weeks, and no new cases of acute jaundice have been reported in Biratnagar.

Persons in Biratnagar were given information regarding epidemics and ways to contain them. They were instructed by electronic media to use boiled water for consumption. It became clear that additional information regarding about maintaining water and sewerage systems during road construction and repair should also be provided. Because 14 patients died of HEV infection during this epidemic, more preparedness for epidemics of waterborne diseases is required to minimize unnecessary illnesses and deaths.

References

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Human Parvovirus 4 Infection among Mothers and Children in South Africa

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To the Editor: Human parvovirus 4 (PARV4) is a single-stranded DNA virus in the family *Parvoviridae* (1).
In Western countries, IgG against PARV4 is largely found only in persons with risk factors for parenteral infection and is strongly associated with co-infection with bloodborne viruses (2–4). In Africa, transmission seems to be more complicated; reported PARV4 seroprevalence is 4%–37%, even among persons at low risk and with no evidence of HIV or hepatitis C virus (HCV) co-infection (1,5,6).

The clinical significance of PARV4 infection remains uncertain. Infections may be asymptomatic, but a variety of clinical associations have been reported (1,7), including an increased rate of progression to AIDS in persons co-infected with HIV (8). This association raises particular concerns for many African populations in which these viruses are co-endemic.

To characterize the epidemiology of PARV4 infection in South Africa, we studied adults and children from pediatric outpatient clinics in Kimberley, South Africa, during May 2009–August 2013. Of the 157 participants, 90 were HIV-1–infected children, 24 their HIV-negative siblings, and 43 HIV-1–infected mothers (of whom 4 had >1 child enrolled). Approval was given by the Ethics Committee of the Faculty of Health Science, University of Free State, South Africa. Written consent was given by all adults and parents/guardians on behalf of their children.

Blood samples were collected from participants, and serum was tested for evidence of PARV4 infection by using ELISA (in duplicate) to detect IgG against PARV4 viral protein 2 (3,6) and by using PCR to detect PARV4 DNA (9). For 92 patients, HIV RNA loads were available; testing was performed by using the Abbott Laboratories m2000 platform (Abbott Park, IL, USA). For 118 of the HIV-infected patients, CD4+ T-cell counts were ascertained by flow cytometry. Statistical analyses were undertaken by using Prism version 6.0f and online software (http://graphpad.com/quickcalcs/). Confidence intervals were calculated by using the adjusted Wald method (http://www.measuringusability.com/wald.htm).

We detected IgG against PARV4 in 58 (37%) of 157 patients; this proportion is broadly comparable with that reported from other settings in sub-Saharan Africa, including Burkina Faso, the Democratic Republic of the Congo, and a previous cohort of HIV-infected persons in South Africa (5). Although routes of transmission in Africa remain to be characterized, these high seroprevalence rates support the possibility that some PARV4 transmission may be occurring by nonparenteral routes, as suggested by others (5,10).

PARV4 IgG seroprevalence was higher among adults (49%) than children (33%), although this difference did not reach statistical significance (p = 0.07, Fisher exact test; Figure, panel A). We found a significant relationship between increasing age and PARV4 IgG serostatus (R² = 0.59 by linear regression, p = 0.025; Figure, panel B). The numbers in each group are small, and further work is needed to define this association with more confidence. We did not detect any cases of PARV4 viremia, suggesting that chronic viremia or reactivation are probably uncommon, even among HIV-infected patients.

On the basis of previously reported data demonstrating PARV4 viremia in neonates (7), we hypothesized that vertical transmission is possible. To investigate further, we sought evidence of concordance between IgG serostatus of mothers and their children. Maternal PARV4 IgG status did not differ between IgG-positive and IgG-negative children (p = 1.00, Fisher exact test; online Technical Appendix Table 1, 714 Emerging Infectious Diseases • www.cdc.gov/eid • Vol. 21, No. 4, April 2015
Co-infection with Avian (H7N9) and Pandemic (H1N1) 2009 Influenza Viruses, China

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Human Parvovirus 4 Infection among Mothers and Children in South Africa

Technical Appendix

Technical Appendix Table 1. Characteristics of 114 children in Kimberley, South Africa, according to PARV4 IgG serostatus*

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>PARV4 IgG negative, n = 77</th>
<th>PARV4 IgG positive, n = 37</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Median age (IQR), y</td>
<td>7 (3–9)</td>
<td>7 (4–8)</td>
<td>0.87†</td>
</tr>
<tr>
<td>Number of males (%)</td>
<td>38 (49)</td>
<td>12 (32)</td>
<td>0.21‡</td>
</tr>
<tr>
<td>Number with HIV (%)</td>
<td>65 (85)</td>
<td>25 (68)</td>
<td>0.05‡</td>
</tr>
<tr>
<td>Median CD4+ T cell percentage (IQR)§</td>
<td>25 (18–30)</td>
<td>24 (16–30)</td>
<td>0.68‡</td>
</tr>
<tr>
<td>Median HIV viral load (IQR), copies/mL§</td>
<td>120,000 (21,600–365,000)</td>
<td>52,000 (7,700–150,000)</td>
<td>0.13†</td>
</tr>
<tr>
<td>Proportion with PARV4 IgG+ mother (%)¶</td>
<td>15/33 (45)</td>
<td>7/14 (50)</td>
<td>1.00‡</td>
</tr>
</tbody>
</table>

*IQR, interquartile range; PARV4, parvovirus 4.
†Mann-Whitney U test.
‡Fisher exact test.
§CD4+ T cells and HIV viral loads are reported for HIV-infected persons only.
¶47 mother–child pairs represented (from a total of 43 mothers recruited; 4 have 2 children each. Numerators in this row add up to 22 because 1 PARV4-positive mother had 2 children).

Technical Appendix Table 2. Characteristics of 43 HIV-positive adults in Kimberley, South Africa, according to PARV4 IgG serostatus*

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>PARV4 IgG negative, n = 22</th>
<th>PARV4 IgG positive, n = 21</th>
<th>p-value†</th>
</tr>
</thead>
<tbody>
<tr>
<td>Median age (IQR), y</td>
<td>26 (22–35)</td>
<td>33 (28–39)</td>
<td>0.09</td>
</tr>
<tr>
<td>Median CD4+ T cell count (IQR), cells/mm³</td>
<td>318 (261–440)</td>
<td>322 (221–469)</td>
<td>0.77</td>
</tr>
<tr>
<td>Median HIV RNA Load (IQR), copies/mL</td>
<td>23,199 (5,050–89,930)</td>
<td>93,000 (8,400–207,500)</td>
<td>0.15</td>
</tr>
</tbody>
</table>

*IQR, interquartile range; PARV4, parvovirus 4.
†Mann-Whitney U test.