Nosocomial Outbreak of Parechovirus 3 Infection among Newborns, Austria, 2014

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In 2014, sepsis-like illness affected 9 full-term newborns in 1 hospital in Austria. Although results of initial microbiological testing were negative, electron microscopy identified picornavirus. Archived serum samples and feces obtained after discharge were positive by PCR for human parechovirus 3. This infection should be included in differential diagnoses of sepsis-like illness in newborns.

Parechoviruses are small, nonenveloped, single-stranded RNA viruses belonging to the family Picornaviridae. Although most human parechovirus (HPeV) infections cause self-limiting mild respiratory or gastrointestinal symptoms, HPeV type 3 (HPeV3) has been found in 5%–13% of newborns and young infants <3 months of age with late-onset sepsis or encephalitis (1–12). Knowledge of HPeV3 infections originates from single cases or small series of sporadic unrelated infections. One considerable outbreak, affecting ≈200 infants with obviously community-acquired diseases, was recently reported from Australia (1,2). In contrast, we describe a timely confined, and apparent nosocomial, outbreak of HPeV3 infection originating from 1 maternity ward in Austria, affecting one fifth of newborns hospitalized during that period.

The Study
During August 5–September 7, 2014, a total of 9 newborns, 2–27 (median 5) days of age, showed signs of sepsis-like illness (fever and reduced general condition) and were admitted to the Department of Paediatrics and Adolescent Medicine, Medical University of Graz, Graz, Austria. All had been delivered within 2 weeks in the same obstetric unit at the Regional Hospital Feldbach, which is located in a small town (4,500 inhabitants) in southeastern Styria, a region of southern Austria of which Graz is the capital. During these 2 weeks, 44 newborns had been delivered at this obstetric unit (total 1,400/y); 9 (20.5%) born during these 2 weeks became symptomatic (Table; Figure). Diagnostic procedures were performed at the discretion of each attending physician and comprised testing for adenovirus, enterovirus, norovirus, rotavirus, herpesvirus 1, herpesvirus 2, varicella zoster virus, Epstein-Barr virus, cytomegalovirus, human herpesvirus 6, and parvovirus B19. However, no causative agent was identified.

Because the only pediatric department in southern Styria (including Feldbach) is at the Medical University of Graz, all severely ill newborns or infants are admitted there. However, no additional cases of sepsis or sepsis-like illness of unclear etiology were identified in newborns or young infants from that area during these months. The facts that the outbreak was temporally confined and all affected patients had been nursed in the same ward strongly indicated a common source within this unit. Investigations to detect a possible common source in the affected obstetric unit comprised anamnestic and clinical examination of hospital staff members and the newborns’ mothers, surface swabbing (e.g. nursery rooms, baby baths, examination beds, diaper changing tables), microbiological examinations of formula, and analysis of staff roster and occupancy plans. However, a presumed common source or causative agent could not be identified.

To further seek the causative pathogen, we analyzed serum, urine, and nasal secretions from 4 patients (nos. 4, 6, 7, 8; Figure) by negative staining in a transmission electron microscope (Zeiss 906, Oberkochen, Germany) at 80 kv. Although nasal secretions revealed no particles, serum and urine of 2 of the tested patients (nos. 6, 8) contained icosahedral particles, diameter 20–30 nm, resembling picornaviruses. Because of the young ages of the patients, we assumed that the particles were HPeVs. However, by the time we received the electron microscope results indicating picornaviruses, all affected patients had been discharged. We therefore requested fecal samples from all 9 discharged patients for molecular diagnostics of HPeV. We received samples from 4 patients (nos. 5, 6, 7, 9) a median of 29 days (range 9–34) after discharge.

RNA was isolated by using a QIAamp Viral RNA Mini Kit (QIAGEN, Hilden, Germany) according to the manufacturer’s instructions. For HPeV detection, we used a real-time reverse transcription PCR (RT-PCR) selective for the 5’ nontranslated region as described (13,14). Molecular typing of
serum samples were positive for HPeV-3, indicating for 3 other patients. For 4 of the 5 patients tested, archived (nos. 2, 3, 5, 6, 7; Figure). No samples had been retained ness, we retrieved archived serum samples from 5 patients to analyze the presence of this pathogen even during acute illness. Molecular typing that used partial VP3/VP1 cap -id protein regions revealed HPeV3 in all 3 samples. To (online Technical Appendix, http://wwwnc.cdc.gov/EID/BLAST (http://blast.ncbi.nlm.nih.gov//Blast.cgi) algorithm were aligned to reference sequences in GenBank by using a Darmstadt, Germany) and nested PCR primers. Sequences subsequently sequenced by using a BigDye Terminator v.3.1 Cy -quently sequenced by using a BigDye Terminator v.3.1 Cy -Kit (QIAGEN). The resulting fragment (≈300 bp) was subse -ficiency was performed by using a HotStarTaq Master Mix Kit (QIAGEN). Resulting sequences were submitted to GenBank (ac -ession nos. KU556748–KU556754). We fecal samples and 4 serum samples) revealed 100% iden -systemic infection with HPeV3 during the symptomatic phase of disease. Comparison of all 7 sequences (from 3 fecal samples and 4 serum samples) revealed 100% identity. Resulting sequences were submitted to GenBank (ac -cession nos. KU556748–KU556754). Clinical signs and symptoms and laboratory changes for these 9 patients were compatible with those published for patients with HPeV3 infection (/2). All 9 newborns re -covered without complications; no severe, long-term comp -lications were noted 15 months later.

Despite intensive epidemiologic evaluation, we were not able to identify a human (personnel, mothers, siblings, or visitors) or nonhuman (surfaces, formula) source of infection within this ward. However, because the causative agent was identified several weeks after the end of the outbreak, testing for HPeV had not been performed on any such human or environmental specimens. Therefore, we are not able to clearly differentiate common-source infection from person-to-person transmission. Because newborns do not have direct contact with each other, an asymptomatic adult or older sibling might have been the unidentified common source. This hypothesis is in line with the fact that most
Infections with HPeV in adults are asymptomatic. Of the 9 newborns, 4 became symptomatic while still hospitalized; thus, they were certainly infected while in the maternity ward. The other 5 became symptomatic after discharge from the obstetric unit, so they might have acquired the infection while outside the hospital. However, the interval between discharge and readmission was <1 week for all but 1 patient, and no cases of community-acquired sepsis and sepsis-like illness in newborns or infants from that region without association with the affected maternity ward could be identified during that period. Thus, the infection was most likely nosocomial for at least 8 of the 9 patients. Assuming the period of infection for these patients (i.e., during their stay in the maternity ward), we can draw conclusions with regard to the incubation period of HPeV3 infections. The observed intervals between infection and appearance of symptoms were from 1 to 12 days. Only patient 9 became symptomatic 27 days after discharge; this patient might have acquired the infection outside the hospital.

**Conclusions**

For newborns, HPeV3 is a relevant pathogen; febrile illness appears as sepsis. After symptomatic infection, a newborn can shed HPeV3 in feces for at least 1 month. The contagious nature of the virus can lead to nosocomial outbreaks. Thus, timely identification of the causative agent may prevent nosocomial transmission (by isolation and identification of the source) and unnecessary treatment with antimicrobial drugs. For newborns with sepsis-like illness, routine diagnostic considerations should include HPeV3.

Dr. Stenger is assistant professor for Pediatrics and Adolescent Medicine at the Medical University of Graz, Austria, and head of the Working Group for Infectious Diseases of the Austrian Society for Pediatrics and Adolescent Medicine. His main research interest focuses on the management of infectious diseases in children and adolescents with impaired immunity.

**References**


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Technical Appendix

Real-time RT-PCR and Amplification of VP3/VP1 Junction Region for Molecular Typing of Parechoviruses

Screening for parechovirus was done by realtime RT-PCR with primers described in Benshop et al. 2008. Briefly, the Superscript III Platinum One-step Quantitative RT-PCR system by Invitrogen (Darmstadt, Germany) was used as described in the manufactures protocol. For each sample, a doubled reaction mix was prepared containing 300 nM of primers Parecho F31 (NRZ 202) and K30 (NRZ 203) as well as 150 nM of probe HPeV-WT (NRZ-TM2). Reverse transcription was carried out for 15 min at 50°C followed by 2 min 95°C for denaturation. Amplification was done in 50 cycles 15 sec 95 and 35 sec 60°C.

Partial VP1 was amplified by using the One-Step-Reverse Transcription PCR Kit (QIAGEN), followed by a nested PCR with HotStarTaq-Mastermix (QIAGEN). Reverse transcription PCR was conducted in a 12.5-μL reaction that contained 2 μL of RNA, 500 nM of primer NRZ 193 500 nM of NRZ 194 according to the manufacturer’s protocol. The temperature profile used was 43°C for 60 min, (53°C for 60 sec, 55°C 60 sec)x20, 70°C for 15 min and 95°C for 15 min, followed by 35 cycles at 94°C for 30 s, 50°C for 30 s, and 72°C for 90 s. Final elongation conducted at 72°C for 10 min.

Two semi-nested PCRs were conducted by using 1 μL of reverse transcription PCR samples in a 12.5-μL volume that contained 500 nM of primer NRZ 193 and 500 nM of primer NRZ 196 as well as 500 nM of primer NRZ 195 and 500 nM of primer NRZ 194. Amplification was conducted by 35 cycles (94°C for 30 s, 55°C for 40 s, and 72°C for 40 s), and final elongation 72°C for 10 min. The resulting amplification products (368 bp and 364 bp) were visualized by electrophoresis on a 1.5% agarose gel.

Sequencing

Partial VP3/VP1 amplification products were used directly for sequencing. Sequence reaction was conducted by using primers NRZ 194, NRZ 195, and NRZ 196. Resulting
sequences were submitted to GenBank under following accession numbers: KU556748 - KU556754

**Technical Appendix Table.** Primers used for reverse transcription PCR to detect human parechovirus

<table>
<thead>
<tr>
<th>Primer/probe</th>
<th>Sequence, 5’→3’</th>
<th>Position in reference strain AB084913</th>
<th>Reference</th>
<th>Reference name</th>
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<tr>
<td>NRZ 202</td>
<td>CTGGGCGCAGGACGA</td>
<td>429–444</td>
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<td>NRZ 203</td>
<td>GGTACCTCTGCGCATCTTC</td>
<td>569–549</td>
<td>(1)</td>
<td>K30</td>
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<td>NRZ TM2</td>
<td>**VIC-MGB-**AAACACTAGTGTA/(A/T)**GGCC-BBBQ</td>
<td>527–546</td>
<td>(1)</td>
<td>HPeV-WT</td>
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<td>NRZ 193</td>
<td>GAYAATGCAATMTAYACWATYTGGA</td>
<td>2089–2114</td>
<td>(2)</td>
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<tr>
<td>NRZ 194</td>
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<td>2523</td>
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<tr>
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<td>NRZ 196</td>
<td>DGGYCCATCAGCWTGWGCTGA</td>
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**References**
