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sp., *Vibrio cholerae*, and *Escherichia coli* (subtypes enterohemorrhagica, enteropatogena, enterotoxigenic and enteroagregatia) by using culture or PCR (8). In 81 (33.3%) of the samples, ≥1 bacterial pathogen was found. The samples originally sent for viral diagnosis (viral cohort, n = 386) had been tested in HUSLAB for norovirus during April–May, 2013 by using reverse transcription quantitative PCR (RT-qPCR) (HUSLAB in-house). Further diagnosis for rotavirus and adenovirus had been requested by physicians from 105 (27.2%) of 386 samples (Dialex MB antigen detection assay, Orion Diagnostica, Espoo, Finland), and for astrovirus from 33 (8.6%) samples (RT-PCR, HUSLAB in-house). A viral pathogen was discovered in 141 (36.5%) samples; in 139, the pathogen was norovirus.

The samples had been sent from diverse locations within Finland, and thus were not from a few isolated outbreaks. No further information on patients and samples was available for either cohort, and not enough samples were left for retrospective analysis of additional pathogens. The Ethics Committee of the Hospital District of Helsinki and Uusimaa approved the study.

BuV DNA was detected by using a new real-time qPCR with the following primers and probe: BuV forward, 5’-ACACGTGTGACAGTG-GATTCAAAACCTT-3’; BuV reverse, 5’-GGTTGTTGTTGATGTGTTAGTTC-3’; BuV qPCR probe, 5’-FAM-CGGAAGAGATTTTGACGATGCYTAGCAA-BHQ1 3’. The detailed qPCR protocol is shown in the online Technical Appendix (wwwnc.cdc.gov/EID/article/20/6/13-1674-Techapp1.pdf). The analytical sensitivity of the RT-qPCR assay was 5–10 copies per reaction.

Of the 629 fecal samples, 7 (1.1%) were positive for BuV DNA, of which 4 were from the bacterial cohort and 3 from the viral cohort. BuV DNA quantity was low in all samples, ranging from 1.9 × 10⁴ to 3.2 × 10⁴ copies per milliliter of fecal supernatant (Table). In contrast to the original discovery of the virus in children with diarrhea (6), all positive samples were from adults (median age 53 years, range 21–89 years). All BuV DNA–positive results were confirmed by repeated BuV qPCR, by amplifying and sequencing another area of the virus, or by both methods (Table): all sequenced amplicons were more similar to the BuV genotype 1 (online Technical Appendix Figure (6)). Two of the BuV-positive samples were from the same patient, taken 4 days apart, and the latter sample also harbored norovirus. The additional 6 BuV-positive samples were negative for the other viral or bacterial pathogens tested.

Seven fecal samples collected from adults in Finland contained BuV DNA, indicating that circulation of the virus is restricted neither to children nor to Africa. However, the low DNA loads in all the positive samples suggest that BuV might not be the primary cause of these cases of gastroenteritis. A known gastroenteritis-inducing pathogen (norovirus) was found in 1 of the 7 BuV-positive samples. We did not observe any clustering of the 7 positive samples into a specific period (Table).

Although the association with gastroenteritis seems weak, BuV might cause symptoms of other types. We did not include feces from healthy subjects for comparison. The identified BuV DNA in our samples could originate from previous or current infections unrelated to gastroenteritis, or be associated with prolonged virus secretion in the respiratory or digestive tracts, a phenomenon shown, e.g., for human bocavirus1 (9,10). Acquisition of the virus from a food source cannot be ruled out, although 1 patient harbored the DNA for at least 4 days, during which a 10-fold increase in viral load was observed.

Overall, this study shows that BuV circulates in northern Europe and can be found in the feces of patients with

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**Table. Samples collected for bacterial and viral testing that were subsequently positive for bocavirus DNA**

<table>
<thead>
<tr>
<th>Sample no.</th>
<th>Sample cohort</th>
<th>Quantity, copies/mL supernatant</th>
<th>Age, y/sex</th>
<th>Pathogens tested for by HUSLAB†</th>
<th>Other pathogens found</th>
<th>Sampling date</th>
<th>Sequenced region, nt, divergence (%) from JX027295‡</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Bacterial</td>
<td>5.2 × 10⁴</td>
<td>21/M</td>
<td>Bacteria</td>
<td>0</td>
<td>2012 Dec 4</td>
<td>VP2, 2786–4495, 0.88</td>
</tr>
<tr>
<td>2</td>
<td>Bacterial</td>
<td>1.9 × 10⁴</td>
<td>38/M</td>
<td>Bacteria</td>
<td>0</td>
<td>2013 Jan 6</td>
<td>VP2, 2786–4495, 0.71</td>
</tr>
<tr>
<td>3‡</td>
<td>Bacterial</td>
<td>1.9 × 10⁴</td>
<td>53/M</td>
<td>Bacteria</td>
<td>0</td>
<td>2013 Jan 11</td>
<td>§</td>
</tr>
<tr>
<td>5</td>
<td>Bacterial</td>
<td>3.7 × 10⁴</td>
<td>46/M</td>
<td>Bacteria</td>
<td>0</td>
<td>2013 Apr 27</td>
<td>VP2, 2786–4495, 0.76</td>
</tr>
<tr>
<td>6†</td>
<td>Viral</td>
<td>3.4 × 10⁴</td>
<td>77/M</td>
<td>Norovirus</td>
<td>0</td>
<td>2013 Apr 19</td>
<td>VP2, 2786–4495, 1.60</td>
</tr>
<tr>
<td>7†</td>
<td>Viral</td>
<td>3.6 × 10⁴</td>
<td>89/F</td>
<td>Norovirus</td>
<td>0</td>
<td>2013 Apr 20</td>
<td>Partial NS, 16–1080, 1.13</td>
</tr>
</tbody>
</table>

*VP2, viral protein 2; NS, nonstructural.
†Samples originally sent to HUSLAB (Helsinki, Finland) for bacterial diagnosis were analyzed for *Salmonella* sp., *Shigella* sp., *Campylobacter* sp., *Yersinia* sp., *Vibrio cholerae*, and *Escherichia coli* (subtypes enterohemorrhagica, enteropatogena, enterotoxigenic, and enteroagregatia) by using culture or PCR. BuV-positives samples could not be analyzed for the presence of pathogens other than those originally tested for because the samples had been discarded.
‡Sequence divergence analyzed by using the DNA distance matrix in BioEdit (www.mbio.ncsu.edu/BioEdit/bioedit.html). The bocavirus sequences were submitted to GenBank (accession nos. KJ461874–KJ461879).
§This sample was positive for bocavirus by quantitative PCR. However, we were not able to amplify another region of the virus from this sample, likely caused by a low amount of the virus in the sample, which had the lowest copy number among the positive samples.
††Samples from the same patient, collected 4 days apart.
gastroenteritis. Despite the absence of known pathogens among 6 of 7 BuVs-shedding patients, the causative role of BuV in gastroenteritis remains uncertain. Serologic studies will help clarify a possible association between BuVs and diarrhea or other diseases.

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Human Granulocytic Anaplasmosis Acquired in Scotland, 2013

To the Editor: Human granulocytic anaplasmosis is a tick-borne disease caused by Anaplasma phagocytophilum, an obligate intracellular gram-negative bacterium that infects granulocytes. The usual clinical signs and symptoms include nonspecific fever, chills, headache, and myalgia. Infection is usually mild or asymptomatic, but severe systemic complications can occur, leading to a need for intensive care and estimated fatality rates of 0.5%–1.0% (1,2).

A. phagocytophilum was first described in 1932 in Scotland as the causative agent of tick-borne fever in sheep (3). Although some clinical cases of human granulocytic anaplasmosis have been reported in Europe, mostly from Slovenia, Sweden, and Poland (4), most cases have occurred in the United States. This difference cannot be explained by the prevalence of the pathogen in ticks or human exposure to the pathogen because the 3% prevalence of A. phagocytophilum among Ixodes ricinus ticks in Europe seems to be nearly as high as that among ticks in the United States (2). The median seroprevalence rate for A. phagocytophilum infection among humans in Europe is 6.2%, reaching up to 21% (2). This incongruence between seroprevalence rate and number of human cases might be associated with underdiagnosis of cases (2), a high rate of asymptomatic disease (5), or cross-reactivities in serologic tests that might lead to overestimation of seroprevalence rate (5).

In August 2013, an immunocompetent 40-year-old man sought treatment for fever (~39°C) and other nonspecific symptoms such as malaise, myalgia, and severe headache 3 days after becoming aware of several tick bites received while on a hiking