A novel pestivirus species was discovered in a piglet-producing farm in Austria during virologic examinations of congenital tremor cases. The emergence of this novel pestivirus species, provisonally termed Linda virus, in domestic pigs may have implications for classical swine fever virus surveillance and porcine health management.

The genus Pestivirus consists of 4 approved species within the family Flaviviridae (1). Besides bovine viral diarrhea virus 1 (BVDV-1), BVDV-2, border disease virus (BDV), and classical swine fever virus (CSFV), several unassigned strains and tentative species are represented by the so-called atypical pestivirus strains. An atypical pestivirus of swine emerged in 2003 in a commercial pig-breeding farm in Australia and was later termed Bungowannah virus (BV). This well-studied virus caused reproductive disorders, stillbirth, and sudden death in piglets, resulting in the loss of ≈50,000 animals in the 2 affected farms. Because of its marked pathogenicity, BV was considered a threat to global pig health, but this virus or relatives were never found at other locations (2).

Recently, a novel group of porcine pestiviruses, termed atypical porcine pestiviruses (APPVs), was discovered. These viruses were identified in North America (3,4) and subsequently detected in Europe (5–8). There is strong evidence that APPVs are a causative agent behind the type A-II congenital tremor (CT A-II) syndrome of piglets (4). CT A-II is characterized by a generalized shaking of the whole body associated with variable degrees of hypomyelination in the brain and spinal cord. However, hypomyelination is also apparent in the brains of fetuses of sheep, cattle, and pigs after late-gestation state infection with BDV, BVDV, or CSFV (9). We report results of an investigation of CT in piglets on a farm in southeastern Austria in which a novel pestivirus species was discovered.

The Study
A small-scale piglet-producing farm in Styria, in southeastern Austria, reported major piglet losses from CT in January 2015. Animals from this farm were examined and samples were taken by authorized veterinarians. The CT-affected piglets showed a severe lateral shaking and were often incapable of sucking milk, which led to elevated pre-weaning death rates (Video, https://wwwnc.cdc.gov/EID/article/23/7/17-0163-V1.htm). The prevalence of CT varied from 20% to 100% within the affected litters. Litters affected by CT were not used for the production of replacement gilts. The outbreak of CT abruptly stopped in July 2015, when all sows had produced 1 CT-affected litter. Only 22.4 piglets per sow were weaned in that year, compared with an average of 25.8 piglets/sow/year before CT symptoms occurred. Pathological examinations were performed as described previously (8), confirming the presence of typical CT A-II lesions (Figure 1, panel A).

We tested various samples from the farm using an APPV-specific TaqMan probe-based reverse transcription PCR (RT-PCR) (8) but obtained negative results. However, we obtained an amplicon of appropriate length from CT-piglet serum samples by using a novel panpestivirus RT-PCR (PPF 5′-GTKATHCAATACCCTGARGC-3′ and PPR 5′-GGRTTCCAGGARTACATCA-3′), which enables detection of CSFV, BVDV-1, BVDV-2, BDV, BV, and APPV. Surprisingly, sequencing of this RT-PCR product yielded an unknown sequence with a noninterrupted open reading frame. An initial BLAST search (https://blast.ncbi.nlm.nih.gov/Blast.cgi) resulted in “no significant similarity found,” but the translated sequence of 270 aa aligned well with different pestiviruses. The viral RNA was detected in all samples (serum, tonsils, lung, liver, spleen, and central nervous system material) of CT-affected piglets and their littermates from the farm.

After inoculating SK-6 cells with serum samples of affected piglets, we detected the amplification of this unknown pestivirus using RT-PCR. We provisionally termed the agent “Linda” (lateral-shaking inducing neurodegenerative agent) to avoid confusion with other pestiviruses. We determined the full genome of Linda virus (LV) using the standard primer walking RT-PCR approach together with rapid amplification of cRNA ends PCR to identify the ultimate 5′- and 3′-termini using the cultured virus. We determined the length of the LV genome (GenBank accession no. KY436034) to be 12,614 nt, with 381 nt 5′-nontranslated region, an open reading frame of 11,772 nt, and 461 nt of 3′-nontranslated region.

We performed a phylogenetic analysis of LV using CLC Workbench 7.6 (CLCBIO, Aarhus, Denmark), which demonstrated LV’s divergence from other pestiviruses (Figure 1, panel B). We found the identity between...
LV, approved pestiviruses, and APPV to be only 60%, but we found an identity of 68% between LV and BV. Comparison of the pestiviral polyprotein sequences yielded an amino acid identity of 69% between LV and BV and of <54% with all other pestiviruses (Figure 1, panel C). All known cleavage sites of the NS3 protease were present in LV (10). We mapped the NS4A-NS4B cleavage, which takes place at an L/A site in classical pestiviruses and at an L/S site in BV, to the L<sub>2354</sub>/S<sub>2355</sub> motive in LV; the intramolecular NS3 cleavage site (L<sub>1834</sub>/A<sub>1835</sub>) was conserved in LV (11). Each hypothetical mature protein of LV matched best with the mature proteins of BV (Table), even if the identities were relatively low in the structural protein region.

We saw a strong reactivity against LV infected cells when using a BVDV E2-specific antibody termed 6A5 (12). This antibody detected the E1-E2 heterodimer of LV (75 kDa) as well as the E2 monomer (50 kDa) in Western blot analysis, although the reactivity against BVDV-1 E2 was much stronger (Figure 2, panel A). With the help of this antibody, we were able to study the in vitro growth of LV. E2 antigen signals of LV were detectable in the cytoplasm of SK-6 and MDBK cells after infection, but the antigen-positive foci were 10-fold larger in SK-6 than in MDBK cells (Figure 2,
Already in the primary passage on porcine cells, considerably high infectious titers of LV were measured in the supernatant (>10^7 50% tissue culture infectious dose/mL). Using monoclonal antibody 6A5 in immunohistochemical analysis, we detected pestiviral antigen in CT-affected piglets in regions with histological lesions. Specific stains in neurons of the nucleus of the trigeminal nerve (Figure 2, panels D,E), in glial cells in the cerebellar and the cerebral white matter, and in tubular epithelium of the kidneys proved that a pestivirus crossed the blood–brain barrier. As in the case of other pestiviruses, the mechanisms inducing hypomyelination during fetal development remain unclear ([8],[13]).

<table>
<thead>
<tr>
<th>Putative mature protein</th>
<th>Amino acid region</th>
<th>Best BLAST hit (GenBank accession no.)</th>
<th>Alignment coverage, %</th>
<th>E-value</th>
<th>Identity, %</th>
</tr>
</thead>
<tbody>
<tr>
<td>N\textsuperscript{pro}</td>
<td>1–182</td>
<td>BV (YP_008992092.1)</td>
<td>96</td>
<td>4 × 10^{-64}</td>
<td>60</td>
</tr>
<tr>
<td>Core</td>
<td>183–283</td>
<td>BV (YP_008992092.1)</td>
<td>95</td>
<td>3 × 10^{-43}</td>
<td>83</td>
</tr>
<tr>
<td>E\textsuperscript{ns}</td>
<td>284–504</td>
<td>BV (YP_008992092.1)</td>
<td>100</td>
<td>5 × 10^{-117}</td>
<td>74</td>
</tr>
<tr>
<td>E1</td>
<td>505–702</td>
<td>BV (YP_008992092.1)</td>
<td>100</td>
<td>3 × 10^{-42}</td>
<td>67</td>
</tr>
<tr>
<td>E2</td>
<td>703–1077</td>
<td>BV (YP_008992092.1)</td>
<td>98</td>
<td>4 × 10^{-140}</td>
<td>53</td>
</tr>
<tr>
<td>P7</td>
<td>1078–1152</td>
<td>BV (YP_008992092.1)</td>
<td>100</td>
<td>1 × 10^{-25}</td>
<td>60</td>
</tr>
<tr>
<td>NS2</td>
<td>1153–1608</td>
<td>BV (YP_008992092.1)</td>
<td>99</td>
<td>0.0</td>
<td>63</td>
</tr>
<tr>
<td>NS3</td>
<td>1609–2291</td>
<td>BV (YP_008992092.1)</td>
<td>100</td>
<td>0.0</td>
<td>85</td>
</tr>
<tr>
<td>NS4A</td>
<td>2292–2354</td>
<td>BV (YP_008992092.1)</td>
<td>100</td>
<td>2 × 10^{-28}</td>
<td>81</td>
</tr>
<tr>
<td>NS4B</td>
<td>2355–2701</td>
<td>BV (YP_008992092.1)</td>
<td>100</td>
<td>0.0</td>
<td>78</td>
</tr>
<tr>
<td>NS5A</td>
<td>2702–3205</td>
<td>BV (YP_008992092.1)</td>
<td>100</td>
<td>0.0</td>
<td>53</td>
</tr>
<tr>
<td>NS5B</td>
<td>3206–3923</td>
<td>BV (YP_008992092.1)</td>
<td>99</td>
<td>0.0</td>
<td>73</td>
</tr>
</tbody>
</table>

*BV, Bungowannah virus; E, envelope; NS, nonstructural.

Figure 2. Detection of pestivirus E2 protein with monoclonal antibody 6A5 in investigation of piglets with congenital tremor (CT) on a farm in southeastern Austria, 2015. A) Western blot analysis of cells infected with novel virus provisionally termed Linda virus. Total protein of SK-6 cells infected with Linda virus and MDBK cells infected with BVDV-1 (strain NADL) was probed with the pestivirus E2-specific antibody 6A5. The apparent molecular mass of monomeric E2 (LV 50 kDa and BVDV-1 55 kDa) shows that Linda virus E2 has a lower molecular weight than BVDV-1 E2 as a result of fewer N-linked glycosylation sites. In contrast, the mass of E1-E2 heterodimers is comparable (~70 kDa in Linda virus and BVDV-1), because Linda virus E1 has an additional N-linked glycosylation site. B, C) Focus size of Linda virus 48 hours after infection of SK-6 cells (B) and MDBK cells (C) (original magnification ×20). D, E) Detection of pestiviral E2 within neuronal tissue of Linda virus–positive, congenital tremor–affected piglets showing positive signals in neurons of the nucleus of the trigeminal nerve (D) (original magnification ×10) and within glial cells in the cerebellar white matter (E) (original magnification ×20). BVDV, bovine viral diarrhea virus.
Conclusions

A previously unknown pestiviral agent was found in CT-affected piglets on a farm in Austria. We observed a severe hypomyelination in the entire white matter of the spinal cord and detected pestivirus antigen in the brain of the CT-affected piglets, suggesting a causal relationship between infection and lesions. Analyses of the assembled genome allowed an unambiguous assignment of LV within the genus Pestivirus with regard to the presence of pestivirus-specific genes (N and E) and sequence homology to other pestiviruses (14). In contrast to APPV, which hardly infects cultured cells at all, LV could be easily propagated on porcine cell lines without the need for adaptation, similar to what has been reported for BV. We suggest that LV likely shares a common ancestor with BV.

After its description ≈10 years ago, BV was intensively sought worldwide, but the virus has never been detected outside Australia. The broad cell culture tropism of BV led to the hypothesis that the virus recently jumped from another species to porcine hosts (15). The discovery of a related pestivirus with substantial sequence divergence in swine on a different continent suggests that both viruses probably have a porcine origin. The identification of the cross-reacting E2-specific monoclonal antibody 6A5 indicates that a cross-reactivity to related pestiviral proteins exists, which might interfere with the serologic testing for CSFV. Further work is needed to investigate the prevalence and the epidemiology of LV in Europe and to assess its virulence in controlled animal experiments.

Acknowledgments

The authors thank the Austrian farmer and the respective practitioner for cooperation in the investigation and Katharina Buczolich, Hann-Wei Chen, and Petra Kodajova for their excellent technical assistance.

The Animal Health Service of Upper Austria (Tiergesundheitsdienst Oberösterreich – TGD ÖÖ) partially funded this study.

B.L., Lu.S., and T.R. are inventors on a patent application on Linda pestivirus (application no. 62/437, 888). Lu.S. and A.L. examined the animals, provided field samples, and wrote parts of the manuscript. B.L., C.R., L.E.S., and T.R. performed the laboratory work and analyzed the data. S.H., B.R.B., and H.W. performed necropsy, histological, immunohistochemical, and ultrastructural examinations on CT animals and wrote parts of the manuscript. B.L. and T.R. interpreted the results and designed the figures. B.L. and T.R. wrote the manuscript. All authors critically analyzed, revised, and approved the manuscript.

Dr. Lamp is a senior scientist at the Institute of Virology at the University of Veterinary Medicine in Vienna, Austria. His primary research interests relate to the replication of RNA viruses and the molecular mechanisms of virus pathogenesis.

References


Address for correspondence: Benjamin Lamp, Institute of Virology, University of Veterinary Medicine, Veterinaerplatz 1, Building AC, 1210 Vienna, Austria; email: benjamin.lamp@vetmeduni.ac.at