Coronaviruses cause respiratory and gastrointestinal diseases in diverse host species. Deltacoronaviruses (DCoVs) have been identified in various songbird species and in leopard cats in China. In 2009, porcine deltacoronavirus (PDCoV) was detected in fecal samples from pigs in Asia, but its etiologic role was not identified until 2014, when it caused major diarrhea outbreaks in swine in the United States. Studies have shown that PDCoV uses a conserved region of the aminopeptidase N protein to infect cells derived from multiple species, including humans, pigs, and chickens. Because PDCoV is a potential zoonotic pathogen, investigations of its prevalence in humans and its contribution to human disease continue. We report experimental PDCoV infection and subsequent transmission among poultry. In PDCoV-inoculated chicks and turkey poults, we observed diarrhea, persistent viral RNA titers from cloacal and tracheal samples, PDCoV-specific serum IgY antibody responses, and antigen-positive cells from intestines.
In vivo studies could validate cell culture susceptibility findings and determine whether PDCoV causes infection and disease in species other than pigs. Virus cross-species transmission among hosts plays a major role in the evolution and diversification of viruses, appearing in many instances to be preferential to coevolving within an initial host (20). CoVs already have demonstrated a propensity for crossing species barriers, both in animal-to-animal spread and animal-to-human spread. Initial evidence of CoVs jumping from mammalian to avian species was reported from bovine CoV infecting turkeys but not chickens (21,22). As a zoonotic CoV transmission, SARS-CoV is believed to have jumped from bats or palm civets (Paguma larvata) to humans in 2002, causing 8,098 cases in 37 countries and 774 deaths (11). MERS-CoV emerged more recently, jumping from dromedary camels (Camelus dromedarius) to humans, and has caused 1,879 cases of respiratory illness in humans and 666 deaths (10).

Understanding how cross-species transmission of CoVs occurs is critical to our ability to predict which viruses might be on the verge of SARS- or MERS-like pandemics. In addition, studies of CoV cross-species transmission can inform development of novel therapeutics and strategies to combat CoVs in susceptible animal hosts before they pose an imminent human health threat. We conducted experiments to determine the prevalence of PDCoV infection in and transmissibility among poultry.

Materials and Methods

Animals
We obtained 25 fourteen-day-old chickens (Gallus gallus domesticus) and 25 fourteen-day-old turkey pouls (Meleagris gallopavo) from the specific pathogen-free flock of the Ohio Agricultural Research and Development Center of The Ohio State University (Wooster, Ohio, USA). This flock has no prior exposure to swine or to PDCoV, PEDV, or TGEV. After acclimating in Biosafety Level 2 (BSL-2) facilities for 1 day, all birds appeared healthy with no evidence of diarrhea or other clinical signs. Animal protocols used in this study were approved by the Institutional Laboratory Animal Care and Use Committee of The Ohio State University.

Titration of PDCoV
We determined PDCoV titers from intestinal contents of pigs by 50% tissue culture infectious dose (TCID₅₀) assay (23). We seeded LLC porcine kidney (LLC-PK) cells at 5 × 10⁴ cells/well in a 96-well plate (BD Biosciences, https://www.bdbiosciences.com). We washed 100% confluent monolayers once with 200 µL of maintenance media (MM): minimal essential medium (MEM), 4-(2-hydroxyethyl)-1-piperazineethanesulfonic acid (HEPES), GlutaMAX (GIBCO, https://www.thermofisher.com) consisting of MEM with 1% antibiotic-antimycotic solution, 1% nonessential amino acids, and 1% HEPES. We inoculated 100 µL from 10-fold dilutions of PDCoV in 8 replicates per dilution. Each plate included 1 row of negative control MM only with 5 µg/mL of Trypsin (Corning, https://www.corning.com). After absorption for 1 h, we added another 100 µL of MM with 5 µg/mL of Trypsin to each well. We monitored cytopathic effects for 3–7 days, calculated virus titers after immunofluorescent (IF) staining by using the Reed-Muench method (24), and expressed results as log₁₀ TCID₅₀/mL (23).

Study Design

Birds were floor housed in a temperature-controlled BSL-2 containment room with wood litter shavings and provided ad libitum access to food and water. In consecutive experiments of pouls and chicks, we randomly divided the flock of 25 birds into 2 groups, 15 uninfected and 10 infected birds. Each group was housed separately and inoculated through the choanal cleft. The uninfected group was inoculated with 200 µL of unfiltered, undiluted small intestine contents (SIC) from an uninfected gnotobiotic pig (GP-8). The infected group was inoculated with SIC from a PDCoV-infected pig (DC175) with 6.87 log₁₀ TCID₅₀/mL. One poult in the uninfected group died of unknown causes unrelated to known pathogens before inoculation.

After inoculation, we observed chicks and pouls for clinical signs 2 times each day. We scored fecal consistency as follows: 0, solid; 1, pasty; 2, semiliquid; and 3, liquid (25). We considered a fecal consistency score of >2 as diarrhea. At 2 days postinoculation (dpi), we randomly assigned 5 birds from each uninfected group as sentinels and allowed them to come into contact with the infected group for the duration of the experiment. We recorded body weights and collected cloacal swab, tracheal swab, and serum samples at 2, 4, 7, 9, 11, and 14 dpi. Except for sentinel birds, we euthanized 2 chicks and 2 pouls from each group at 3 and 7 dpi for blood and tissue collection. We concluded the study at 14 dpi and euthanized the remaining 33 birds, including sentinels, for blood and tissue collection.

Serum Antibody Detection
We modified and optimized an ELISA (26) to detect PDCoV-specific IgY antibodies in serum from
PDCoV-inoculated chicks and poults. We added 50 µL of serum diluted 1:1,000 to the PDCoV antigen-coated and mock antigen-coated wells and incubated for 90 min at 37°C, then added 100 µL of biotin-conjugated antichicken IgY (Invitrogen Goat anti-Chicken IgY [H+L] Secondary Antibody, Biotin; ThermoFisher, https://www.thermofisher.com) or biotin-conjugated antiturkey IgY (Goat Anti-Turkey IgY [H+L] Biotin pAb; Cell Sciences, https://www.cellsiences.com) at a dilution of 1:10,000 and incubated at 37°C for 1 h. We added 100 µL of HRP-Conjugated Streptavidin (ThermoFisher) to each well at a dilution of 1:5,000 and incubated at 37°C for 1 h. We washed wells with phosphate buffered saline solution with 0.05% Tween-20 (×5) between each step. We added 3,3′,5′-tetramethylbenzidine substrate (SureBlue TMB 1-Component Microwell Peroxidase Substrate; Seracare, https://www.seracare.com), then added 100 µL of 0.3 mol/L sulfuric acid to stop the reaction. We read plates at an absorbance of 450 nm by using a SpectraMax F5 (Molecular Devices, https://www.moleculardevices.com) plate reader. We conducted statistical analysis by using Prism software (GraphPad, https://www.graphpad.com). We used analysis of variance to compare multiple groups and a 1-tailed Student t-test to compare groups of 2.

**Histopathology and IF Staining**

We examined gross tissues from small intestines, duodenum to ileum, and large intestines, cecum and colon, as well as other organs, including bursa, lung, liver, kidney, proventriculus, and spleen, and then fixed tissues in 10% neutral formalin for 1–2 days at room temperature for histopathology (27). We embedded, sectioned, and then stained samples with hematoxylin and eosin for light microscopy examination. We measured mean jejunal or ileal ratios of villus height and crypt depth (VH:CD) by using MetaMorph software (MetaMorph, Inc., https://www.metamorphsoftware.com), as described previously (18). We tested prepared tissues by IF staining to detect PDCoV antigen using a polyclonal rabbit antiserum against PDCoV (provided by E. Nelson, South Dakota State University, Brookings, SD, USA) (28). We also tested tissues from a PDCoV-infected pig for comparison.

**Real-Time Reverse Transcription PCR**

We suspended cloacal swabs, tracheal swabs, SIC, and large intestine contents (LIC) in 1–4 mL MEM as a 10% suspension. We extracted RNA by using GenCatch Viral RNA Miniprep Kit (Epoch Life Science, https://www.fishersci.com). We further processed samples containing fecal matter by using OneStep PCR Inhibitor Removal Kit (Zymo Research Corporation, https://www.zymoresearch.com).

We determined viral RNA titers by real-time reverse transcription-PCR (rRT-PCR), as reported previously (23). In brief, we amplified a 541-bp fragment of the M gene that covered the quantitative RT-PCR-amplified fragment. We designed 5′-CCGCTAACGTGGATACTATGT-3’ and 5′-CCGCGCTTTGAGTGTAT-3’ primers according to the sequence of a strain from the United States, Illinois121/2014 (GenBank accession no. KJ481931). We purified the PCR products by using a QIAquick PCR Purification Kit (QIAGEN Inc., https://www.qiagen.com), sequenced, and then used these as the template to construct a quantitative RT-PCR standard curve. The detection limit of the rRT-PCR was 10 genomic equivalents (GEs)/reaction, which corresponded to 4.6 log_{10} GE/mL of PDCoV in cloacal and tracheal samples.

**Results**

**Clinical Signs**

By 2 dpi, 70% of infected chicks had diarrhea and fecal scores >2; that percentage decreased to 17% by 9 dpi (Table 1). By 14 dpi, most (5/6) of the remaining infected chicks had normal feces. The 5 uninfected sentinel chicks that cameled with the infected birds at 2 dpi demonstrated mild to moderate diarrhea 2 days after comingling (4 dpi); diarrhea peaked 5 days after comingling (7 dpi). At 14 dpi, only 2 sentinel chicks had abnormal feces. Two chicks in the uninfected group had transient diarrhea during the study but tested negative for known pathogens, including PDCoV.

In poults, 50% exhibited diarrhea at 2 dpi. During the study, the rate of diarrhea increased, and poult(s) did not recover by 14 dpi. (Table 1). Sentinel poult(s) began exhibiting mild to moderate diarrhea 5 days after comingling (7 dpi), and by 9 dpi, 60% were affected. At 14 dpi, all 5 sentinel poult(s) had moderate diarrhea.

Two infected chicks necropsied at 3 dpi had distended gastrointestinal tracts containing a mixture of yellow liquid and gas. Similar but less extensive findings were seen in infected chicks at 7 dpi. No gross pathology was detected in infected chicks or sentinel chicks at 14 dpi. In necropsies of infected poult(s), we observed distended gastrointestinal tracts containing a mixture of yellow liquid and gas at all time points.

**Weights**

Birds were weighed before inoculation and then at 2, 4, 7, 9, 11, and 14 dpi. PDCoV infection greatly affected the chicks’ weight at 2 dpi (Figure 1, panel A). At
4 and 7 dpi, weight gain averages in infected chicks were comparable to those in uninfected birds, but at 9 and 11 dpi, infected chicks had gained much less weight than the uninfected chicks. By 14 dpi, infected chicks rebounded and showed compensatory weight gain at higher levels than uninfected chicks.

Poult weight gain responses differed from those of the chicks. At 2 dpi, infected poult weight gain was severely curtailed; several lost weight, and the average weight gain for the infected group was 0.5 g, compared with almost 10 g for uninfected poult weight gain. By 7 dpi, the infected poult recovered and gained weight at a slightly higher, but not statistically significantly different, rate than the uninfected poult weight gain. This trend continued until the end of the study.

Histopathology and IF Staining
We examined tissue sections by using light microscopy. We noted suspect zymogen depletion in several poult tissues in both the infected and uninfected groups, suggesting possible inanition. We conducted VH:CD measurements of the ileum and jejunum of intestinal tissues from chicks at 14 dpi (Table 2). Infected chicks had a VH:CD ratio of 4.26:1 compared with a ratio of 6.15:1 for uninfected chicks. The VH:CD ratio was lower in sentinel chicks than in uninfected chicks but the difference was not statistically significant.

We could not obtain enough measurements for poult tissues to provide accurate comparisons. IF tissue staining in infected poult tissues demonstrated PDCoV antigen detectable in the epithelial cells lining the villi of the jejunum, although at reduced levels from the ileum and from infected porcine tissue (Figure 2, panel A). We also detected PDCoV antigen in numerous epithelial cells that had sloughed off and remained in the lumen of infected poult tissues when compared with stained tissue sections from uninfected poult tissues (Figure 2, panel B and C). We were unable to visualize a signal in tissues from chicks.

Serum IgY Antibody Responses
We analyzed serum samples collected at 2, 4, 7, 9, 11, and 14 dpi. We used indirect ELISA to test samples

![Figure 1](image-url)

**Figure 1.** Average weight gain/day of (A) chicks and (B) turkey poult tissues in a study of infection and transmission of porcine deltacoronavirus in poultry. Weights were taken at 2, 4, 7, 9, 11, and 14 dpi and differences were averaged by the number of days between time points. Weights for sentinel birds are excluded after 2 dpi. Error bars indicate upper half of SD. Statistically significant values are indicated: *p<0.05; **p<0.01; ***p<0.001. dpi, days postinoculation.
Porcine Deltacoronavirus in Poultry, United States

at 2 and 14 dpi for PDCoV-specific IgY antibodies in all birds from each group, including sentinel birds. We assigned experimental values by averaging 3 replicates. Because we did not have positive controls in chicks and poults, we established a cutoff by using the average final optical density value of uninfected birds at 2 dpi plus 2 SD. We established a separate cutoff value for sentinel birds.

At 14 dpi, infected chicks had increased IgY antibody levels in serum, demonstrating an antibody response to PDCoV (Figure 3, panel A), but sentinel chicks did not have antibody levels demonstrating exposure to PDCoV (Figure 3, panel B). Serum samples from poults exhibited a similar range of IgY values. The average IgY values were much higher in infected birds at 14 dpi compared with infected birds at 2 dpi and uninfected birds (Figure 3, panel C). The IgY greatly increased in sentinel poults at 14 dpi compared with IgY values at 2 dpi, but were still below the cutoff value (Figure 3, panel D).

teRT-PCR on Samples from Chicks

All experimentally infected chicks rapidly shed detectable viral RNA postinoculation, and viral RNA titers remained relatively constant through 11 dpi (Figure 4, panels A, B). Viral RNA from cloacal swabs reached 6.52 log_{10} GE/mL by 2 dpi and remained >6.5 log_{10} GE/mL until 11 dpi, when levels at 7.14 log_{10} GE/mL at 9 dpi, then decreased to 5.82 log_{10} GE/mL at 14 dpi. Despite an absence of noticeable respiratory signs, tracheal swab specimens also showed high levels of PDCoV RNA throughout the study (Figure 4, panel B). PDCoV spread rapidly from infected to naive birds, and all 5 sentinel chicks became positive for PDCoV RNA in both tracheal and cloacal swabs within 2 days of comingling with infected birds (Figure 4, panels C, D).

We calculated titers and viral RNA loads in SIC and LIC from infected and sentinel chicks at 3, 7, and 14 dpi (Figure 5). We used RNA isolated at 14 dpi from SIC of 1 infected and 1 sentinel bird to amplify an ≈1,300-bp portion of the nucleocapsid (N) gene of PDCoV, then gel extracted and sequenced the resulting product. The samples sequenced had >99% identity with the original inoculum, Ohio FD22 strain of PDCoV. We tested infectivity of intestinal contents of infected and sentinel chicks by using TCID50 assay at 7 and 14 dpi (Figure 5, panel A).

rRT-PCR on Samples from Poults

Similar to the results from chicks, results for infected poults showed all had high levels of PDCoV RNA in cloacal and tracheal swabs through 14 dpi (Figure 6, panels A, B). Poults appeared to have higher initial viral loads, averaging 8.07 log_{10} GE/mL by 2 dpi, decreasing to ≈6 log_{10} GE/mL at 4 dpi, and persisting through 14 dpi (Figure 6, panel A). Naive birds also were susceptible to infection, and cloacal and tracheal swab specimens from all sentinel poults were positive for PDCoV RNA within 2 days after comingling with

Table 2. Effect of porcine deltacoronavirus on villus height: crypt depth ratios of ileum or jejunum in experimental chicks

| Characteristics                      | Uninfected, n = 6 | Infected, n = 6 | Sentinel, n = 5 |
|--------------------------------------|-------------------|-----------------|----------------|
| Villous height                       | 370               | 353             | 547            |
| Crypt depth                          | 64                | 84              | 94             |
| Ratio of villous height to crypt depth| 6.15              | 4.26            | 5.87           |

*Values are expressed in millimeters and represent the average of ≤10 measurements/bird at 14 d post inoculation with porcine deltacoronavirus. Bold text indicates statistically significant difference, p = 0.04.
infected poults (Figure 6, panels C, D, E). We calculated titers and viral RNA loads in SIC and LIC from infected and sentinel poults at 3, 7, and 14 dpi (Figure 5, panel B). We tested infectivity of intestinal contents of infected and sentinel poults by using TCID50 assay at 7 and 14 dpi (Figure 5, panel B).

We isolated viral RNA from the SIC of 1 infected and 1 sentinel bird at 14 dpi and used it to amplify an ≈1,300-bp portion of the N gene of PDCoV, then gel extracted and sequenced the resulting product. As we noted in chicks, the samples sequenced had >99% identity with the original inoculum, the Ohio FD22 strain of PDCoV.

**Discussion**

Emerging viruses in at least 2 genera of porcine CoVs have exhibited increased propensity for interspecies transmission (2,29). Porcine APN was identified as a major cell entry receptor for PDCoV (19,30). APN is a protein that exhibits enzymatic activity, peptide processing, cholesterol uptake, and chemotaxis to cell signaling and cell adhesion (31). APN is widely distributed and highly conserved in amino acid sequences across species of the Animalia kingdom (19) and is expressed in a wide range of tissues, including epithelial cells of the kidneys (31), respiratory tract (32,33), and gastrointestinal tract (34).

Our data suggest that chicks and poults are susceptible to infection with PDCoV. In addition, the rapid transmission of PDCoV to the sentinel birds that comingled with infected birds demonstrates that the virus could spread easily. The length of our pilot study did not allow us to determine how long the chicks and poults would be affected by PDCoV or how long they might shed viral RNA. The chicks appeared to recover more rapidly than poults; clinical signs diminished or were completely absent by 14 dpi. However, chicks still were shedding low viral RNA titers at 14 dpi. Poults did not recover by the end of the study and still exhibited gross pathology and mild to moderate diarrhea. PDCoV RNA shedding titers were higher in poults than in the chicks. Cloacal shedding titers in chicks peaked at 9 dpi and then decreased. In poults, cloacal viral RNA shedding titers were multiphasic, peaking at 2 dpi, with additional smaller peaks at 9 and 14 dpi. The rapid onset of viral RNA shedding correlates with previous in vitro data in which the PDCoV S1 domain bound most efficiently to APN of galline origin (19) and cytopathic effects were observed more rapidly in leghorn male hepatoma and DF1 chicken cell lines compared with swine testicular cells (S.P. Kenney, unpub. data).

ELISA results showed that both chicks and poults developed PDCoV antibodies by 14 dpi. Pig infection dynamics have demonstrated a similar serum
neutralizing antibody titer increase at 7–14 days \((12,25)\). Sentinel birds had low or undetectable antibody responses compared with experimentally challenged birds, likely because of the passive infection method and because less time passed between exposure to the virus and the end of the study.

Recent studies demonstrated that PDCoV can infect and kill cells of other species through APN receptors \((19,35)\). PDCoV has been reported to infect commercial chickens in vivo \((36)\). The differences in susceptibility to PDCoV infection between chicks and poults we observed could be related to differences in APN expression levels between the species.

The true incidence rates for PDCoV infection, natural host range, reservoirs, and routes of transmission are still relatively unknown, and no plans for vaccine development have been reported \((37)\). DCoV RNA has been detected in fecal samples from wild birds \((38,39)\), Chinese ferret badgers \((Meles leucurus)\), and leopards \((40)\). In addition to swine, calves have been shown by experimental testing to be susceptible to PDCoV infection \((41)\). These data, coupled with the
PDCoV binding receptor APN being conserved across many species, suggest that the host range for PDCoV is broader than initially expected (19). The close sequence homology between DCoV isolates from mammalian and wild bird species implies a transmission cycle in which PDCoV regularly crosses from wild birds and mammals into animal production systems, including the swine and poultry industries. More epidemiologic data are required to understand the full extent to which DCoVs are threatening food production systems and whether they pose a direct threat to human health.

Our results are consistent with the likelihood that avian species act as potential passthrough or intermediate hosts for PDCoV. In vivo confirmation of avian susceptibility to PDCoV suggests that in vitro data implicating human susceptibility should be evaluated further. Research regarding how PDCoV is adapting and mutating in different species and whether it infects humans is critical to determining if PDCoV poses a pandemic health risk to commercial poultry or humans.

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References
1. Masters PS, Perlman S. Coronaviridae. In: Knipe DM, Howley PM, eds. Fields virology, 6th ed. Philadelphia: Lippincott Williams & Wilkins; 2013. p. 825–58.
2. Chan JF, To KK, Tse H, Jin DY, Yuen KY. Interspecies transmission and emergence of novel viruses: lessons from bats and birds. Trends Microbiol. 2013;21:544–55. https://doi.org/10.1016/j.tim.2013.05.005
3. Schwegmann-Wessels C, Herrler G. Transmissible gastroenteritis virus infection: a vanishing specter. Dtsch Tierarztl Wochenschr. 2006;113:157–9.
4. Song D, Zhou X, Peng Q, Chen Y, Zhang F, Huang T, et al. Newly emerged porcine deltacoronavirus associated with diarrhoea in swine in China: identification, prevalence and full-length genome sequence analysis. Transbound Emerg Dis. 2015;62:575–80. https://doi.org/10.1111/tbed.12399

5. Huang YW, Dickerman AW, Pinedeyro P, Li L, Fang L, Kiehne R, et al. Origin, evolution, and genotyping of emergent porcine epidemic diarrhea virus strains in the United States. MBio. 2013;4:e00377-13. https://doi.org/10.1128/mBio.00377-13

6. Memish ZA, Zumla AI, Al-Hakeem RF, Al-Rabeeah AA, Stephens GM. Family cluster of Middle East respiratory syndrome coronavirus infections. N Engl J Med. 2013;368:2487-94. https://doi.org/10.1056/NEJMoai1303729

7. de Wit E, van Doremalen N, Falzarano D, Munster VJ. SARS-CoV-2 in gnotobiotic pigs. Emerg Infect Dis. 2014;20:662–5. https://doi.org/10.3201/eid2004.131685

8. Lau SK, Fun SK, Lam CS, Lau CC, Tse H, Tsoi HW, Cheng VC, et al. Full-length genome sequence of porcine deltacoronavirus from the United States. Genome Announc. 2014;2:e00218-14. https://doi.org/10.1128/genomeA.00218-14

9. Lau SK, Woo PC, Yip CC, Tse H, Tsoi HW, Cheng VC, et al. Coronavirus HK1 and other coronavirus infections in Hong Kong. J Clin Microbiol. 2006;44:2063–71. https://doi.org/10.1128/JCM.02614-05

10. Fehr AR, Perlman S. Coronaviruses: an overview of their evolution. Virus Res. 2016;226:71–84. https://doi.org/10.1016/j.virusres.2016.05.028

11. Zhang J. Porcine deltacoronavirus: overview of infection dynamics, diagnostic methods, prevalence and genetic evolution. Virus Res. 2012;165:3995–4008. https://doi.org/10.1016/j.virusres.2016.05.028

12. Jung K, Hu J, Everly B, Lu Z, Chepngeno J, Saif LJ. Pathogenicity of 2 porcine deltacoronavirus strains in gnotobiotic pigs. Emerg Infect Dis. 2015;21:650–4. https://doi.org/10.3201/eid2104.141859

13. Viitos-Sillman S, Loy JD, Brodersen B, Kelling C, Doster A, Topliff C, et al. Expression of aminopeptidase N and dipeptidyl peptidase IV in the healthy and asthmatic bronchus. Clin Exp Allergy. 1998;28:110–20. https://doi.org/10.1046/j.1365-2222.1998.00198.x

14. Li W, Hulswit RJJ, Kenney SP, Widjaja I, Jung K, Alhamo MA, et al. Broad receptor engagement of an emerging global coronavirus may potentiate its diverse cross-species transmissibility. Proc Natl Acad Sci U S A. 2018;115:E5135–43. https://doi.org/10.1073/pnas.1802879115

15. Geoghegan JL, Duchène S, Holmes EC. Comparative analysis estimates the relative frequencies of co-divergence and cross-species transmission within viral families. PLoS Pathog. 2017;13:e1006215. https://doi.org/10.1371/journal.ppat.1006215

16. Ismail MM, Tang AY, Saif YM. Pathogenicity of turkey coronavirus in turkeys and chickens. Avian Dis. 2003;47:515–22. https://doi.org/10.1637/5917

17. Ismail MM, Cho KO, Ward LA, Saif LJ, Saif YM. Experimental bovine coronavirus in turkey poults and young chickens. Avian Dis. 2001;45:157–63. https://doi.org/10.2307/1593023

18. Hu H, Jung K, Vlasova AN, Chepngeno J, Lu Z, Wang Q, et al. Isolation and characterization of porcine deltacoronavirus from pigs with diarrhea in the United States. J Clin Microbiol. 2015;53:1537–48. https://doi.org/10.1128/JCM.00031-15

19. Reed LJ, Muench H. A simple method of estimating fifty percent endpoints. Ann. J Epidemiol. 1938;37:493–7.

20. Lelievre C, et al. Experimental infection of conventional nursing pigs and their dams with porcine deltacoronavirus. Emerg Infect Dis. 2015;21:650–4. https://doi.org/10.3201/eid2104.141859

21. Ismail MM, Tang AY, Saif YM. Pathogenicity of turkey coronavirus in turkey poults and young chickens. Avian Dis. 2001;45:157–63. https://doi.org/10.2307/1593023

22. Hu H, Jung K, Vlasova AN, Saif LJ, Saif YM. Experimental infection of gnotobiotic pigs with the cell-culture-adapted porcine deltacoronavirus strain OH-FD22. Arch Virol. 2016;161:3421–34. https://doi.org/10.1007/s00705-016-3056-8

23. Jung K, Kim J, Ha Y, Choi C, Chae C. The effects of transplacental porcine circovirus type 2 infection on porcine epidemic diarrhoea virus-induced enteritis in preweaning piglets. Vet J. 2006;171:445–50. https://doi.org/10.1016/j.tvjl.2005.02.016

24. Okda F, Lawson S, Liu X, Singrey A, Clement T, Hain K, et al. Development of monoclonal antibodies and serological assays including indirect ELISA and fluorescent microsphere immunoassays for diagnosis of porcine deltacoronavirus. BMC Vet Res. 2016;12:95. https://doi.org/10.1186/s12917-016-0716-6

25. Vlasova AN, Saif LJ. Biological aspects of the interspecies transmission of selected coronaviruses. In: Singh SK, editor. Viral infections and global change. Hoboken (NJ): John Wiley & Sons; 2013. p. 393–418.

26. Jønsdóttir HR, Møller GM, Hoogsteden HC, et al. Expression of aminopeptidase N and dipeptidyl peptidase IV in the healthy and asthmatic bronchus. Clin Exp Allergy. 1998;28:110–20. https://doi.org/10.1046/j.1365-2222.1998.00198.x

27. Dijkman R, Jebbink MF, Koekkoek SM, Deijs M, Jønsdóttir HR, Mølkenkamp R, et al. Isolation and characterization of current human coronavirus stains in primary human epithelial cell cultures reveal differences.
34. Kenny AJ, Maroux S. Topology of microvillar membrane hydrolases of kidney and intestine. Physiol Rev. 1982;62:91–128. https://doi.org/10.1152/physrev.1982.62.1.91

35. Wang B, Liu Y, Ji CM, Yang YL, Liang QZ, Zhao F, et al. Porcine deltacoronavirus engages the transmissible gastroenteritis virus functional receptor porcine aminopeptidase N for infectious cellular entry. J Virol. 2018;92:e00318-18. https://doi.org/10.1128/JVI.00318-18

36. Liang Q, Zhang H, Li B, Ding Q, Wang Y, Gao W, et al. Susceptibility of chickens to porcine deltacoronavirus infection. Viruses. 2019;11:E573. https://doi.org/10.3390/v11060573

37. Wang B, Liu Y, Ji CM, Yang YL, Liang QZ, Zhao F, et al. Porcine deltacoronavirus engages the transmissible gastroenteritis virus functional receptor porcine aminopeptidase N for infectious cellular entry. J Virol. 2013;87:6081–90. https://doi.org/10.1128/JVI.03368-12

38. Wang Q, Vlasova AN, Kenney SP, Saif LJ. Emerging and re-emerging coronaviruses in pigs. Curr Opin Virol. 2019;34:39–49. https://doi.org/10.1016/j.co.viro.2018.12.001

39. Hu H, Jung K, Wang Q, Saif LJ, Vlasova AN. Development of a one-step RT-PCR assay for detection of pan-coronaviruses (α-, β-, γ-, and δ-coronaviruses) using newly designed degenerate primers for porcine and avian fecal samples. J Virol Methods. 2018;256:116–22. https://doi.org/10.1016/j.jvirmet.2018.02.021

40. Dong BQ, Liu W, Fan XH, Vijaykrishna D, Tang XC, Gao F, et al. Detection of a novel and highly divergent coronavirus from Asian leopard cats and Chinese ferret badgers in southern China. J Virol. 2007;81:6920–6. https://doi.org/10.1128/JVI.00299-07

41. Jung K, Hu H, Saif LJ. Calves are susceptible to infection with the newly emerged porcine deltacoronavirus, but not with the swine enteric alphacoronavirus, porcine epidemic diarrhea virus. Arch Virol. 2017;162:2357–62. https://doi.org/10.1007/s00705-017-3351-z

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