Original Article

Evaluation of safety, humoral immune response and faecal shedding in horses inoculated with a modified-live bovine coronavirus vaccination

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Summary
Equine coronavirus (ECoV) is considered an emerging enteric virus with reported morbidity rates ranging from 10 to 83% and fatality rates ranging from 7 to 27% in adult horses; a vaccine for ECoV is currently not available. This study investigated the safety, humoral response and viral shedding in horses inoculated with a commercially available modified-live bovine coronavirus (BCoV) vaccine. Twelve healthy adult horses were vaccinated twice, 3 weeks apart, either orally, intranasally or intrarectally. Two healthy unvaccinated horses served as sentinel controls. Following each vaccine administration, horses were monitored daily for physical abnormalities whilst the onset and duration of BCoV shedding was determined by quantitative PCR (qPCR) in nasal secretions and faeces. Whole blood was collected every 3 weeks to determine BCoV-specific antibody response. With the exception of transient and self-limiting changes in faecal character observed in seven vaccinated and one control horse, no additional abnormal clinical findings were found in the study horses. Following the first and second vaccine administration, two and one horse respectively, tested qPCR-positive for BCoV in nasal secretions 1-day post intranasal vaccination. No vaccinated horses tested qPCR-positive for BCoV in faeces following each vaccine administration. One of the two horses that shed BCoV seroconverted to BCoV after the first vaccine administration and an additional two vaccinated horses (oral and intrarectal) seroconverted to BCoV after the second vaccine administration. In conclusion, the results show that the modified-live BCoV is safe to administer to horses via various routes, causes minimal virus shedding and results in detectable antibodies to BCoV in 27% of the vaccinates.

Introduction
Equine coronavirus (ECoV) is considered an emerging enteric virus with reported morbidity rates ranging from 10 to 83% and a fatality rates ranging from 7 to 27% in adult horses (Oue et al. 2011; Pusterla et al. 2013; Fielding et al. 2015). Clinical signs include pyrexia, anorexia, lethargy, colic and diarrhoea accompanied by leucopenia secondary to neutropenia and lymphopenia. Recent studies reported the detection of ECoV worldwide in faeces and respiratory secretions of adult horses with pyrogenic and enteric disease (Oue et al. 2013; Pusterla et al. 2013; Miszczak et al. 2014). Coronaviruses are grouped into three different genera: alphacoronavirus, betacoronavirus and gammacoronavirus (Woo et al. 2009). ECoV shares the betacoronavirus-1 genera together with bovine coronavirus (BCoV) and is therefore closely related (Woo et al. 2009). Given the strong homology between BCoV and ECoV, the former could therefore act as a surrogate for ECoV vaccination (Guy et al. 2000; Oue et al. 2011).

Vaccination is the cornerstone of reducing clinical signs of viral disease but at present a vaccine for ECoV is not available. An inactivated BCoV vaccine was recently shown to lead to a measurable BCoV and ECoV antibody response (Nemoto et al. 2017). The recent study used an inactivated adjuvanted BCoV vaccination which probably would lead to a more measurable immune response than a modified-live vaccine but the effect on mucosal immunity is still undetermined in horses. Therefore, investigating a modified-live vaccine could, if safe, change the mucosal immunity and response to infection.

A modified-live bivalent vaccine containing bovine rotavirus and coronavirus is commercially available for the reduction of enteric disease in cattle. Field efficacy studies in calves have shown a significant reduction in the incidence and death from neonatal calf diarrhoea following the use of the modified-live BCoV vaccine (Waltner-Toews et al. 1985).

Based on the close genetic and antigenic relationship between ECoV and BCoV, it was hypothesised that the modified-live BCoV vaccine would be safe to administer to adult healthy horses, would induce a short duration of viral shedding based on the vaccine administration route and probably trigger a measurable humoral immune response in vaccinated horses.

Materials and methods
Fourteen adult, healthy horses housed individually at the Center for Equine Health, School of Veterinary Medicine, University of California at Davis were enrolled in the study. Animal use was approved by the Institutional Animal Care and Use Committee at the University of California at Davis. The study horses were deemed healthy based on normal physical examination and haematological parameters. Further, nasal secretions and faeces from every study horse were determined to be qPCR-negative for BCoV and ECoV and all horses tested seronegative against BCoV and ECoV prior to study commencement. Four study horses per group
were vaccinated either orally, intranasally or intrarectally using a commercially available modified-live bovine rotavirus coronavirus vaccine (CalfGuard). The vaccine vials were reconstituted according to the manufacturer’s recommendations and given either orally, intranasally using a small 6” nasal cannula or intrarectally using a 10 French 10” rubber catheter. To increase the vaccine volume and increase mucosal contact, the oral and intrarectal vaccines were diluted each in 50 mL of PBS prior to administration. Vaccinated horses were given a primary vaccination followed by a booster vaccination 3 weeks later (as per the manufacturer’s label). Two additional horses remained unvaccinated and served as environmental sentinels. Following the first and second vaccination period, all horses were monitored daily, for 7 days, by means of a physical examination. Further, faeces and nasal secretions were collected daily from each of the study horses for the molecular detection of BCoV via qPCR. Briefly, the qPCR assay used was based on the detection of a specific 93 basepair product of the N gene of BCoV (GenBank accession number EU401980.1: oligonucleotides: forward primer BCoV-826fl CCCAATAAAAAATGCACTGTTCA, reverse primer BCoV-919r CACTAGTCCAAAGTTCATCTCC, probe UPL #119 TGGGTGGT). The samples were amplified in a combined thermocycler/fluorometer (7900 HT Fast), with the standard thermal cycling protocol: 2 min at 50°C, 10 min at 95°C and 40 cycles of 15 s at 95°C and 60 s at 60°C. Furthermore, a qPCR assay targeting a universal sequence of the bacterial 16S rRNA gene (faeces) and a qPCR assay targeting the housekeeping gene eGAPDH (nasal secretions) were used as quality control (i.e. efficiency of DNA purification and amplification) and as an indicator of inhibition (Mapes et al. 2007). A standard curve was run for the assay using BCoV plasmids and the amplification efficiency calculated from the slope using the formula $E = 10^{(-1/C)} - 1$. The amplification efficiency was 98.3% for the N gene of ECoV indicating a very high analytical sensitivity. The qPCR assay was able to detect the modified-live BCoV vaccine but unable to detect field sample positive for ECoV.

Whole blood was collected prior to each vaccination and every 3 weeks up to 21 days following the second vaccine administration for the detection of specific antibodies to BCoV. A commercial ELISA for BCoV was utilised to detect specific antibodies to BCoV. Briefly, blood samples were centrifuged for 1 h after collection and serum was aliquoted and stored at −80°C until analysed. The BCoV antibody test manufacturer’s instructions were followed except the secondary antibody was replaced with hors eradish peroxidase conjugated antiequine IgG antibody at a dilution of 1:110,000 and 5 positive and 5 negative equine coronavirus serum samples were used to determine OD cut-off titre. Each serum sample was diluted 1:25 with saline and was added to both the viral antigen and noninfectious BCoV antigen wells. After one-hour incubation and plate washing, the secondary antibody was added and incubated for one additional hour. The plate was washed again, and substrate solution was added for exactly 10 min and then stopped. The plate was read within 10 min of stopping at an OD of 450 nm. The ELISA was able to reliably and repeatedly classify negative and positive equine control serum samples using an OD cut-off of 0.233.

Results

The study group consisted of seven geldings and seven mares with an age range of 11–22 years (median 17 years). One of the oral vaccinated horses was lost to follow-up shortly before the second vaccine administration due to an unrelated disease. Physical examination showed no abnormalities with the exception of transient and self-limiting changes in faecal character (“cow-pie” faeces) in eight horses (Table 1). During the primary vaccination, 2 intranasally vaccinated horses (Horse 1: Days 1–6; Horse 4: Days 6–7), 3 intrarectally vaccinated horses (Horse 5: Days 6–7; Horse 6: Days 6–7; Horse 8: Days 5–7) and one orally vaccinated horse (Horse 10: Day 5) demonstrated “cow-pie” faeces. Following booster vaccination, all the same horses showed “cow-pie” faeces (Horse 1: Days 2, 6 and 7; Horse 4: Day 3; 2 None Positive nasal secretions (2nd day)
4 Positive (2) Positive nasal secretions (2nd day)

Table 1: Specific clinical, molecular and serological findings determined for each study horse during the two vaccination periods

| Vaccination group | Horse | Soft formed faeces (number of days) | BCoV PCR (time post-vaccination) | BCoV Serology | Day 42 |
|-------------------|-------|-----------------------------------|---------------------------------|--------------|-------|
| Nasal             | 1     | Positive (6)                      | Negative                         | Positive (4) | Negative |
|                   | 2     | None                              | Positive nasal secretions (2nd day) | Negative     | Negative |
|                   | 3     | None                              | Negative                         | Positive (2) | Negative |
|                   | 4     | Positive (2)                      | Positive nasal secretions (2nd day) | Positive (1) | Positive |
| Rectal            | 5     | Positive (2)                      | Negative                         | Positive (1) | Negative |
|                   | 6     | Positive (2)                      | Negative                         | Positive (1) | Negative |
|                   | 7     | None                              | Negative                         | Positive (4) | Negative |
|                   | 8     | Positive (3)                      | Negative                         | None         | Positive |
| Oral              | 9     | None                              | Negative                         | None         | Negative |
|                   | 10    | Positive (1)                      | Negative                         | Positive (2) | Negative |
|                   | 11    | None                              | Negative                         | None         | Negative |
|                   | 12    | Positive (1)                      | Negative                         | None         | Negative |
|                   | 13    | None                              | Negative                         | None         | Negative |
|                   | 14    | None                              | Negative                         | None         | Negative |
Horse 5: Day 3; Horse 6: Day 1; Horse 8: Days 1, 5, 6 and 8 and Horse 10: Days 6–7) with the addition of one intranasally vaccinated horse (Horse 3; Days 1 and 7) and one control horse (Horse 13: Day 7). One of the intranasally vaccinated horses had soft formed faeces prior to vaccination and throughout the study (Horse 1).

Following primary nasal vaccination, two horses tested qPCR-positive for BCoV 24 h post-vaccination administration (Table 1). One of the two horses tested qPCR-positive for BCoV 24 h post-vaccine administration in nasal secretions following the second vaccination. All other nasal swabs tested qPCR-negative for BCoV. The faecal samples from all vaccinated and control horses tested qPCR-negative for BCoV following both vaccine administration periods.

A total of three horses seroconverted to BCoV (Table 1). One intranasally vaccinated horse seroconverted after the first vaccine and remained seropositive following the second vaccine administration. This horse was not one of the two horses that tested qPCR-positive in nasal secretions. Further, one orally and one intrarectally vaccinated horse seroconverted after the second vaccine administration.

One horse had a normocytic, normochromic anaemia (28.8%; ref: 30–46%), one horse was lymphopaenic (926/µL; ref: 1600–5800/µL) prior to the second vaccination and three horses had a mild eosinophilia 204–340/µL (ref: 0–200/µL). All other haematological parameters were within reference ranges for the laboratory.

Discussion

The results from this study showed that the modified-live BCoV vaccine was safe to administer via various routes, as none of the vaccines developed clinical signs associated with enteric coronavirus infection (lethargy, anorexia, fever, diarrhoea or colic). This is in agreement with various studies evaluating the safety of BCoV vaccination in cattle (Thurber et al. 1977; Walthner-Toews et al. 1985). The only physical abnormal findings noticed during the study period was mild, transient changes in faecal character in eight study horses. Of those horses affected by changes in faecal character, three were in the intranasal group, three in the intrarectal group, one in the oral group and one control. Without evidence of BCoV faecal shedding in all horses with “cow-pie” faeces, and the inclusion of a control, it is probable that the transient faecal character change was related to dietary or environmental factors. During the study protocol, the faecal samples were not analysed for bovine rotavirus which can have a genetic homology with equine rotavirus (Ghosh et al. 2013). Rotavirus can cause severe diarrhoea in foals but does not cause clinical disease in adult horses and therefore was not deemed necessary to test during the study. It is not possible to definitively rule out the bovine rotavirus as a cause of the soft faeces but given the inclusion of a control horse it seems unlikely.

BCoV shedding was only found in nasal secretions of two intranasally vaccinated horses. The detection of BCoV by qPCR in the two horses 24 h after vaccine administration was attributed to persistence of vaccine virus rather than viral replication in respiratory epithelial cells.

Overall, 27% of vaccinated horses had a measurable serological immune response after two vaccine administrations. Seroconversion to BCoV was determined in one horse from each of the three vaccination routes, indicating antigen detection by the immune system. The study investigated alternative vaccine administration routes in order to determine if a serological response could be achieved whilst bypassing the stomach, known to negatively impact avirulent pathogens due to its low pH. Intrarectal vaccine administration of Lawsonia intracellularis in foals has been previously shown to be more effective than oral vaccination, as this route triggered a weak humoral and long-lasting cell-mediated response (Pusterla et al. 2009). The limited number of horses showing a measurable serological response for the various administration routes could relate to the vaccine antigen mass, inability of the vaccine BCoV strain to replicate in equine epithelial cells or pre-existing neutralising cross-reactive antibodies to ECoV. Unfortunately, the study did not measure mucosal or cell-mediated immunity and detection of peripheral antibodies to BCoV alone is not a measure of protection.

During the study period, no serological diagnostics existed to confirm the seropositive, or seronegative, ECoV status of a horse. An ELISA has since been created, and the samples were retrospectively tested for ECoV antigens (Kooijman et al. 2016). All samples were shown to be negative for antigens against ECoV, and therefore, there should not have been alteration in the response to the vaccine due to previous exposure to ECoV. Therefore, the changes in BCoV serological values are probably due to the administration of the vaccine.

A previous study looked at the response to a BCoV vaccination which was based on an inactivated vaccine rather than the modified-live vaccine used in this study (Nemoto et al. 2017). Various studies have assessed the response in equine cases to both inactivated and modified-live vaccines with the latter showing an improved reduction in clinical signs in Equine Herpes Virus-1 infections (Goodman et al. 2006) whilst in humans a modified-live vaccine leads to an increased IgA concentration on mucosal surfaces (Cox et al. 2004). Overall modified-live vaccinations show a significantly higher immunogenicity than inactivated vaccines (Kollaritsch and Rendi-Wagner 2013) but killed-advantageous vaccines will trigger a more reliable, measurable immune response, although, mucosal immunity response is still undetermined. Therefore, use of the modified-live vaccine may have represented a more efficacious route of vaccination in ECoV.

The limitations of the current study included the small number of horses, the unknown viability of the virus within the oral and nasal cavity and stomach and the unknown naivety of the horses to coronavirus in the study.

Conclusions

As the threat of ECoV amongst adult horses increases, there is a need to develop a means of effective protection. The modified-live BCoV vaccine appears safe in adult horses; however, it triggers a limited seroconversion response. At this time the vaccine cannot be recommended for the prevention of ECoV. Future studies will investigate mucosal and cell-mediated immunity and investigate the efficacy of the BCoV in the prevention of ECoV infection under experimental conditions.

Authors’ declaration of interests

None of the authors of this paper has a financial or personal relationship with other people or organisations that could inappropriately influence or bias the content of this paper.
Ethical animal research
Animal use was approved by the Institutional Animal Care and Use Committee at the University of California at Davis (approval number 18417; date of approval 9 October 2014).

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Authorship
J. Prutton and N. Pusterla contributed to study design, execution, analysis and preparation of the manuscript. S. Barnum (née Mapes) contributed to execution. All authors gave their final approval of the manuscript.

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3Svanovir, Uppsala, Sweden.
4Sigma-Aldrich, St Louis, Missouri, USA.

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