



Intradermal and intramuscular vaccination against PRRS with modified live vaccine confer similar immune response in presence of maternally derived antibodies

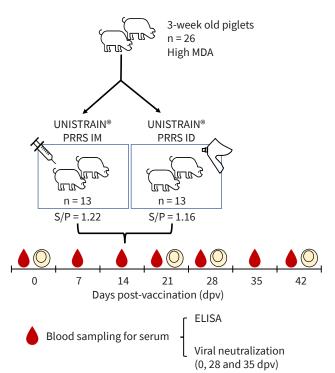
Aguirre, L.1*; Li, Y.1; Baratelli, M.2; Martín-Valls, G.1; Cortey, M.1; Miranda, J.2; Martín, M.1; Mateu, E.1.

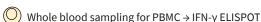
- ¹ Departament de Sanitat i Anatomia Animals, Universitat Autònoma de Barcelona (UAB), 08193 Cerdanyola del Vallès, Spain
- ² HIPRA, Amer (Girona), Spain

Background & Objectives

Maternally-derived antibodies (MDA) may protect piglets against infections during the first weeks of life. However, MDA may also interfere with vaccinations ⁽¹⁾. The present study aimed to assess the development of the immune response against PRRSV after vaccination by either the intradermal (ID) or the intramuscular (IM) route in piglets with high MDA levels.

Materials & Methods





Results

Despite the presence of high MDA levels, vaccination resulted in seroconversion by 14 dpv (average S/P IM: 2.10 ID: 2.13) with high S/P values until the end of the study (average S/P IM: 2.76, ID: 2.94). Both groups developed neutralizing antibodies by 28 dpv that increased by 35 dpv (IM: 4.6±2.8 and ID: $3.9\pm3.4\,\text{Log}_2$) (Fig.1). IFN- γ responses were detected by 21 dpv (IM: 122 and ID: 148 secreting cells/10 6 PBCMs) and further increased by 42 dpv (IM: 159 and ID:189 cells/10 6 PBCMs) (Fig. 2). Differences between groups were non-significant.

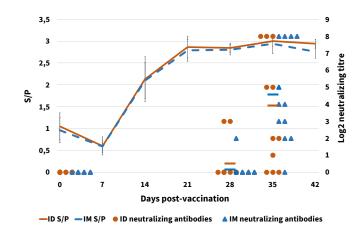


Figure 1. Evolution of S/P ratios as determined by ELISA and virus neutralization in IM and ID groups. The vertical bars indicate the standard deviation, and horizontal bars indicate mean log, neutralization titres.

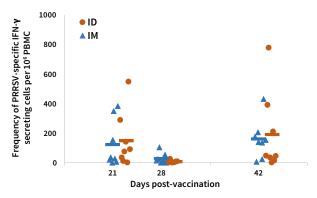


Figure 2. IFN-γ secreting cells frequencies by ELISPOT at 0, 21, 28 and 42 dpv. Horizontal bars indicate mean values for each group and date.

Discussion & Conclusion

The results of the present study suggest that UNISTRAIN® PRRS was able to produce an immune response in piglets despite the presence of high MDA Notably, this response was consistent with previous studies in which the vaccinated animals did not have MDA (2.3). Moreover, intramuscular and intradermal routes of administration produce similar immune responses.

References

- 1. Renson P et al. 2019. Vaccine. 18;37(31):4318-4324.
- 2. Madapong A et al. 2020. Vet. Microbiol. 244.
- 3. Martelli P et al. 2009. Vaccine. 27, 3788-3799.

^{*}Corresponding author: laia.aguirre@uab.cat