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Efficacy and Duration of Infection Study for RespiSure® and Draxxin® Against a *Mycoplasma hyopneumoniae* Challenge in Swine

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Introduction

Experimental infections with *Mycoplasma hyopneumoniae* (*M. hyopneumoniae*) can persist in the pig's respiratory tract for longer than 200 days post challenge with complete natural clearance confirmed by 254 days following challenge¹. The objective of this study was to evaluate the duration of *M. hyopneumoniae* infection in swine that received either a *M. hyopneumoniae* bacterin vaccination prior to infection, or combination of *M. hyopneumoniae* bacterin prior to infection, and tulathromycin following infection.

Materials and Methods

This study was conducted at commercial swine facilities. All animals enrolled in the study were from a single source, with an established status of being *M. hyopneumoniae* free, based on clinical history and laboratory tests. All pigs were enrolled in the study at weaning, approximately 19 days of age. There were 7 treatment groups (T1 – T7). Two hundred fifty-five (255) animals were designated as negative control animals (T1). Pigs in T2 – T7 were vaccinated for *M. hyopneumoniae* with RespiSure™ at 3 and 6 weeks of age. Eighty (80) animals were enrolled per each of groups (T2-T7). Animals were clinically healthy and were 60 days of age (+/- 5 days) at the time of challenge. All animals were administered an intratracheal challenge of 10 ml of the 1:100 dilution of the challenge material, *M. hyo*-positive lung homogenate obtained from Iowa State University, Ames, IA (*M. hyopneumoniae* strain 232², 1x10⁶ CCU/ml) on study day 35, which was 14 days after last vaccination. Pigs in T3, T5 and T7 were all administered two doses of Draxxin® (tulathromycin) injectable solution (2.5 mg/kg) at various intervals post-challenge (Table 1). Subsets of pigs from all treatment groups had lungs removed at slaughter, and a bronchial swab collected from each animal. Swabs were stored appropriately and transported to the University of Minnesota Veterinary Diagnostic Lab (UofM-VDL) for *M. hyopneumoniae* real time PCR (VetMax™, Applied Biosystems). Any lungs with lesions

suspicious of *M. hyopneumoniae* had tissues collected and transported to the VDL for histopathology examination and further real time PCR.

Table 1. Antibiotic treatment and sample collection schedule for experimental groups

| Treatment | Tulathromycin (Days post-challenge) | Bronchial Swab (Days post-challenge) |
|-----------|-------------------------------------|--------------------------------------|
| 1 | N/A | 133, 161, 189 |
| 2 | N/A | 133 |
| 3 | 90 & 100 | 133 |
| 4 | N/A | 161 |
| 5 | 122 & 132 | 161 |
| 6 | N/A | 189 |
| 7 | 150 & 160 | 189 |

N/A: Not applicable.

Results

The *M. hyopneumoniae* real time PCR results from bronchial swabs are summarized below:

| Day post-challenge | 133 | | 161 | | 189 | |
|--------------------------|-----|-------|-----|-------|-----|-------|
| | n | % Pos | n | % Pos | n | % Pos |
| Control | 63 | 100 | 65 | 43.1 | 93 | 23.7 |
| Vaccination | 65 | 78.5 | 65 | 55.4 | 63 | 34.9 |
| Vaccination & Medication | 69 | 73.9 | 63 | 49.2 | 64 | 15.6 |

Discussion

It has been previously demonstrated that *M. hyopneumoniae* infected pigs can be carriers of the pathogen while convalescent carriers can remain infectious for up to 200 days. Total clearance of *M. hyopneumoniae* occurs by 254 days post-inoculation¹. Under the conditions of this study, the use of *M. hyopneumoniae* vaccination alone or in combination with treatment with tulathromycin did not result in an elimination of the *M. hyopneumoniae* carrier status at 133, 161 or 189 days post-inoculation.

References

- 1 Pieters, et al., 2009. Vet Micro, 134 :261-266.
- 2 Minion et al., 2009. J Bacteriol. 186:7123-7133.