A Cross-Sectional Study of Porcine Reproductive and Respiratory Syndrome Virus and *Mycoplasma hyopneumoniae* in Wild Boars Reared in Different Types of Captive Setting in Thailand

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**Abstract**

Porcine reproductive and respiratory syndrome virus and *Mycoplasma hyopneumoniae* are account as economically important pathogens in swine business worldwide. The studies of these two pathogens in Thailand were restricted on domestic pigs in swine industry and the status of these pathogens in wild boars is currently not available. This cross-sectional study aimed at estimating the seroprevalence of antibodies to these agents in captive wild boars. Fifty-two serum samples were collected from wild boars reared in three different types of captive settings including a commercial wild boar farm, a wildlife breeding research station and a zoo. All serum samples were examined for antibodies to porcine reproductive and respiratory syndrome virus and *Mycoplasma hyopneumoniae* by ELISA methods. The results revealed that the highest true prevalence of both pathogens was found in sera derived from wild boars in wildlife breeding research station at 69 and 66% for porcine reproductive and respiratory syndrome virus and *Mycoplasma hyopneumoniae* respectively. This study is the first to reveal seroprevalence of antibodies against porcine reproductive and respiratory syndrome virus and *Mycoplasma hyopneumoniae* in wild boar populations in Thailand. More detailed investigations are critically needed.

**Keywords:** Captive wild boar; *Mycoplasma hyopneumoniae*; PRRSV; Seroprevalence; Thailand

**Introduction**

Porcine reproductive and respiratory syndrome virus (PRRSV) and *Mycoplasma hyopneumoniae* are account as economically important pathogens worldwide [1,2]. Both genotype 1 and genotype 2 of PRRSV were detected in domestic pig farms in all regions of Thailand. Prevalence of PRRSV in domestic pigs in the country varied from 24.6-53.6% and the prevalence was 32.7% in the central region [3] where the density of domestic pig population is high. The recorded pig density in this region according to previous study was between 0.00 and 777.56 heads per square kilometer [4]. Nonetheless, the clusters of PRRSV were not associated with geographic distribution of the virus. The identical viruses were present in different provinces [5]. The PRRSV infected pigs are often co-infected with *M. hyopneumoniae* and other agents causing porcine respiratory disease complex (PRDC) [2,3]. The prevalence of *M. hyopneumoniae* was previously reported at 2.08-45.00% in domestic pigs in Thailand [6]. There was evidence suggesting that both PRRSV and *M. hyopneumoniae* can be transmitted via airborne over long distance up to 9.2 km from the source herd [7]. The exchange of these two agents between domestic pigs and wild boars might exist in Thailand especially in the high density area of pig production in the central region.

Wild boars are reared in Thailand for many purposes such as research activities, exhibition in the zoos and selling as exotic meat. At least three types of captive wild boar settings were recognized in Thailand including wildlife breeding research station, zoo and commercial wild boar farm [8]. However, the studies of PRRSV and *M. hyopneumoniae* in Thailand were restricted on domestic pigs in swine industry. Hence, the epidemiological status of these two pathogens in wild boars (*Sus scrofa*) is currently not available in the country.

Therefore, this preliminary cross-sectional study aimed to initiate infectious disease study in wild boars by estimating seroprevalence of antibodies against PRRSV and *M. hyopneumoniae* in captive wild boars reared in different types captive settings.

**Materials and Methods**

**Sample collection**

Three wild boar captive settings were conventionally selected to represent each type of wild boar rearing. A wildlife breeding research station and a zoo are located in two major pig-producing provinces in the central region namely Ratchaburi province and Chonburi province respectively. These two provinces were recorded as potential foci for PRRSV and *M. hyopneumoniae* in wild boar populations.

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as the areas with highest density of domestic pig population in Thailand [4]. However, the information on wild boar population density is currently not available. Some important characteristics of each studied captive setting are shown in Table 1.

The prevalence of the studied diseases was unknown in these locations and populations. Thus, sample size in this study was calculated based on 50% estimated prevalence with precision of 5% which resulted in highest number of samples required. A statistical package named SAMSIZESIZE was used in our sample size calculation (http://sampsize.sourceforge.net/iface/). Since the total number of animals in all these studied premises is low, all wild boars in each captive setting were required to be tested according to the calculation.

Between February and March 2009, fifty-two wild boars were captured for blood sample collection; 19, 16 and 17 samples were derived from commercial wild boar farm, wildlife breeding research station and the zoo respectively (sampling fraction=95%, 100% and 100%), After collection and processing, serum samples were then securely delivered to serological laboratory of the Monitoring and Surveillance Center for Zoonotic Diseases in Wildlife and Exotic Animals, Faculty of Veterinary Science, Mahidol University within 24 hours.

**Laboratory examination**

All serum samples were serologically examined for antibodies to PRRSV and *M. hyopneumoniae* by ELISA methods according to manufacturer’s instructions (IDEXX Laboratories, Inc., Westbrook, Maine, USA). S/P (sample to positive) ratio was used to determine seropositivity where S is the difference between optical density of the sample and optical density of the negative control and P is the difference between optical density of the positive control and optical density of the negative control. Tested Samples were interpreted as seropositive for both PRRSV and *M. hyopneumoniae* when S/P ratio was equal or higher than 0.4.

**Prevalence calculation**

True prevalence was calculated based on sensitivity and specificity of the tests. The sensitivity and specificity of the ELISA essay for PRRSV are 99.9% and 98.8% and the values are 37.3% and 98.6% respectively for *M. hyopneumoniae* testing.

**Results**

The results of this study revealed that 13 out of 52 tested sera had antibodies against PRRSV (24% true prevalence; 95% CI: 0.14-0.38). The true prevalence as high as 69% (95% CI: 0.43-0.87) was found in wildlife breeding research station in Ratchaburi province. In contrast, relatively low value of true prevalence was observed in commercial wild boar farm in Phetchabun province and zoo in Chonburi province at 4% (95% CI: 0.00-0.24) and 5% (95% CI: 0.00-0.27), respectively. *M. hyopneumoniae* was detected in 4 out of 52 tested samples (18% true prevalence; 95% CI: 0.04-0.46) and *M. hyopneumoniae* positive sera were only found in wildlife breeding research station. The true prevalence at this captive setting was 66% (95% CI: 0.21-1.00). Seroprevalence of PRRSV and *M. hyopneumoniae* in all sampling locations was summarized in Table 2 and geographic locations of sample collecting sites together with true prevalence were shown in Figure 1.

**Discussion**

The highest true prevalence of PRRSV was detected in the wildlife breeding research station in Ratchaburi province, the province

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### Table 1: Some important characteristics of each studied captive setting and the area of study site.

| Parameter | Commercial Wild Boar Farm (VS) | Wildlife Breeding Research Station (KS) | Zoo (KK) |
|-----------|-------------------------------|----------------------------------------|----------|
| Number of wild boars | 20 | 16 | 17 |
| Wild boar replacement strategy | Reproducing within the farm and purchasing from other farms | Reproducing within the station and obtaining from donations | Reproducing within the zoo and obtaining from donations |
| Vaccination against PRRSV and *M. hyopneumoniae* | No | No | No |
| Domestic pig density in the area of study site (heads per square meters)* | 4.50-7.43 | 147.00-777.56 | 63.51-148.99 |
| Prevalence of PRRSV in domestic pigs in the area of study site (%) ** | 53.6 | 35.1 | 30.5 |
| Prevalence of *M. hyopneumoniae* in domestic pigs in the area of study site (%) *** | N/A**** | 2.08-45.00 | 2.08-45.00 |

*Knips [4] **Tummaruk et al. [3], *** Boonsoongnern et al. [6]. **** N/A = not available

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### Table 2: Number of samples tested and number of positive samples as well as seroprevalence of porcine reproductive and respiratory syndrome virus (PRRSV) and *Mycoplasma hyopneumoniae* in wild boars derived from three different types of captive settings including a commercial wild boar farm (VS), a wildlife breeding research station (KS) and a zoo (KK) in Thailand.

| Type of captive setting | Number tested | PRRSV | M. hyopneumoniae |
|------------------------|---------------|-------|-----------------|
|                        | Number positive | Appearance prevalence | True prevalence | 95% CI* | Number positive | Appearance prevalence | True prevalence | 95% CI* |
| Commercial wild boar farm (VS) | 19 | 1 | 0.05 | 0.04 | 0.00-0.24 | 0 | 0 | 0 | 0.00-0.43 |
| Wildlife breeding research station (KS) | 16 | 11 | 0.69 | 0.69 | 0.43-0.87 | 4 | 0.25 | 0.66 | 0.21-1.00 |
| Zoo (KK) | 17 | 1 | 0.06 | 0.05 | 0.00-0.27 | 0 | 0 | 0 | 0.00-0.49 |
| Overall | 52 | 13 | 0.25 | 0.24 | 0.14-0.38 | 4 | 0.07 | 0.18 | 0.04-0.46 |

* 95% CI = 95% Confidence Interval
where co-circulation of multiple clusters from two genotypes of PRRSV in domestic pigs was reported [9]. Besides, Ratchaburi is the province that possessed highest record of domestic pig density in Thailand [4]. With this high density and co-circulation of multiple clusters of PRRSV in the province, it is possible that this particular pathogen was shared between wild boars and domestic pigs reared in this area. An in-depth investigation is needed to explore the genetic linkage of the PRRSV found in domestic pigs and wild boars and further determine whether wild boars can serve as reservoir of the disease for domestic pigs in that location. Nevertheless, only 2% seropositivity of PRRSV was found in wild boars population lived in the wild in a previous study conducted in the US [1]. It is noticeable that prevalence of PRRSV found in captive wild boars in the current study was much higher than the prevalence detected many months earlier. However, antibody detection methods were used instead of detecting the antigens. The current infectious status respectively. Thus, the seropositive results found in this study can reflect the infections happened in these populations in many months earlier. However, antibody detection methods were used instead of detecting the antigens. The current infectious status of the animals in these settings was not known.

The persistence of these two pathogens is likely to allow the diseases to silently infect the animals unnoticeably after the replacement wild boars were introduced into the captive settings. Therefore, it is suggested that the newly introduced animals should be tested for important infected pathogens such as PRRSV and M. hyopneumoniae and should be quarantined before introducing into the herds.

Conclusion

The present study illustrated the picture of PRRSV and M. hyopneumoniae seropositivity in three premises representing three types of wild boar captive settings. More locations and samples are required in order to obtain more accurate infectious status of these diseases in Thailand. Since PRRSV and M. hyopneumoniae are prevalent elsewhere such as USA in feral wild boars that habit in the wilds [1], the future study should also expand to this type of population.

This study is the first to report seroprevalence of antibodies to PRRSV and M. hyopneumoniae in wild boar populations in Thailand. More detailed investigations are critically needed.

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References

1. Baker SR, O’Neil KM, Gramer MR, Dee SA (2011) Estimates of the seroprevalence of production-limiting diseases in wild pigs. Vet Rec 168: 564.

2. Sibila M, Montabre G, Boadella M, Huerta E, Casas-Diaz E, et al. (2010) Serological, pathological and polymerase chain reaction studies on Mycoplasma hyopneumoniae infection in the wild boar. Vet Microbiol 144: 214-218.

3. Tummaruk P, Surapat P, Sirriyakun S, Seeimakram O, Olarnratmanee EO, et al. (2012) Porcine reproductive and respiratory syndrome virus detection in Thailand during 2005-2010 in relation to clinical problems, pig types, regions, and seasons. Trop Anim Health Prod 45: 771-779.

4. Knips V (2004) Review of the livestock sector in the Mekong countries. Livestock Information, Sector Analysis and Policy Branch (AGAL), Food and Agriculture Organization of the United Nations.

5. Thawongnuwech R, Amonsin A, Tatsanakit A, Damrongwatanapokin S (2004) Genetics and geographical variation of porcine reproductive and respiratory syndrome virus (PRRSV) in Thailand. Vet Microbiol 101: 9-21.

6. Boonsoongnern A, Jirawattanapong P, Lertwatcharasarakul P, Phatthanakunanan S, Poolperm P, et al. (2012) The Prevalence of Mycoplasma hyopneumoniae in Commercial Suckling Pigs in Thailand. World J Vaccine 2: 161-163.

7. Otake S, Dee S, Corzo C, Oliveira S, Deen J (2010) Long-distance airborne transport of infectious PRRSV and Mycoplasma hyopneumoniae from a swine population infected with multiple viral variants. Vet Microbiol 145: 198-208.

8. Wiratsudakul A, Sariya L, Prompiram P, Tantawet S, Suraruangchai D, et al. (2012) Detection and phylogenetic characterization of hepatitis E virus genotype 3 in a captive wild boar in Thailand. J Zoo Wildl Med 43: 640-644.

9. Tun HM, Shi M, Wong CL, Ayudhya SN, Amonsin A, et al. (2011) Genetic diversity and multiple introductions of porcine reproductive and respiratory syndrome viruses in Thailand. Virol J 8: 164.

10. Thacker EL, Halbur PG, Ross RF, Thanawongnuwech R, Thacker BJ (1999) Mycoplasma hyopneumoniae potentiation of porcine reproductive and respiratory syndrome virus-induced pneumonia. J Clin Microbiol 37: 620-627.

11. Vengust G, Valencak Z, Bidovec A (2006) A serological survey of selected pathogens in wild boar in Slovenia. J Vet Med B Infect Dis Vet Public Health 53: 24-27.

12. Erlandson KR, Evans RB, Thacker BJ, Wegner MW, Thacker EL (2005) Evaluation of three serum antibody enzyme-linked immunosorbent assays for Mycoplasma hyopneumoniae. J Swine Health Prod 13: 198-203.

13. Olarnratmanee EO, Wangnaltham S, Thanawongnuwech R, Kunavongkrit A, Tummaruk P (2011) Prevalence of porcine reproductive and respiratory syndrome virus (PRRSV) antigen-positive uterine tissues in gilts culled due to reproductive disturbance in Thailand. Trop Anim Health Prod 43: 451-457.

14. Pieters M, Pijuan C, Fano E, Dee S (2009) An assessment of the duration of Mycoplasma hyopneumoniae infection in an experimentally infected population of pigs. Vet Microbiol 134: 261-266.

15. Desrosiers R, Boutin M (2002) An attempt to eradicate porcine reproductive and respiratory syndrome virus (PRRSV) after an outbreak in a breeding herd: eradication strategy and persistence of antibody titers in sows. J Swine Health Prod 10: 23-25.

16. Große Beilage E, Rohde N, Krüter J (2009) Seroprevalence and risk factors associated with seropositivity in sows from 67 herds in north-west Germany infected with Mycoplasma hyopneumoniae. Prev Vet Med 88: 255-263.