

## Sponsors

---

### *We thank the following sponsors:*

#### **Platinum**

Bayer Animal Health  
National Pork Board  
Pfizer Animal Health

#### **Silver**

Boehringer Ingelheim Vetmedica, Inc.

#### **Bronze**

Cargill  
Merck Animal Health  
Novartis Animal Health

#### **Copper**

AgStar Financial Services  
Elanco Animal Health  
IDEXX  
Newport Laboratories  
PIC USA  
PRRS CAP

#### **University of Minnesota Institutional Partners**

College of Veterinary Medicine  
University of Minnesota Extension  
College of Food, Agriculture and Natural Resources Sciences

#### **Formatting**

Tina Smith Graphics  
[www.tinasmithgraphics.com](http://www.tinasmithgraphics.com)

#### **CD-ROM**

David Brown  
[www.davidhbrown.us](http://www.davidhbrown.us)

#### **Logo Design**

Ruth Cronje, and Jan Swanson;  
based on the original design by Dr. Robert Dunlop

The University of Minnesota is committed to the policy that all persons shall have equal access to its programs, facilities, and employment without regard to race, color, creed, religion, national origin, sex, age, marital status, disability, public assistance status, or sexual orientation.

# Detection of influenza virus in aerosols from swine farms

C.A. Corzo<sup>1</sup>; M. Torremorell<sup>1</sup>; S. Dee<sup>2</sup>; M. Gramer<sup>1</sup>; R. Morrison<sup>1</sup>

<sup>1</sup>College of Veterinary Medicine, University of Minnesota, St. Paul, Minnesota

<sup>2</sup>Pipestone Veterinary Clinic, Pipestone, Minnesota

## Introduction

Influenza A virus transmission in swine occurs mainly by direct nose-to-nose contact and aerosol. It has been documented that pigs can shed influenza virus in their nasal secretions for approximately 7 days after infection.<sup>1</sup> Transmission between farms is thought to occur through movement of infected pigs.<sup>2</sup> However, there are reports in which outbreaks of respiratory disease in pig farms have occurred suddenly and are not associated to the introduction of animals. In addition, farm density has been associated to increase in the risk of influenza infections<sup>3</sup> which suggest that airborne transmission of influenza between farms may be possible. Information on transmission of influenza A in pigs is scarce, therefore, the objective of this study is to detect airborne influenza A virus from acutely infected pig populations in the field.

## Materials and methods

Two acutely infected pig populations raised under commercial conditions were identified through constant communication with local veterinarians. Upon diagnostic confirmation of influenza infection by RRT-PCR in pigs affected with respiratory signs, a total of fifteen 30-minute air samples were collected inside and outside the barn using a cyclonic collector. Additionally, 15 oral fluid samples were collected from pigs inside the barn. If a population tested positive in the first visit, a second visit was scheduled seven days after the first visit and testing repeated. Air and oral fluid samples were tested for influenza RNA by RRT-PCR.<sup>4</sup> Further diagnostics included virus isolation, titration, strain subtyping and sequencing.

## Results

In farm 1 (nursery), an influenza A H1N2 virus was detected in pigs with acute respiratory signs suggestive of influenza infection. Based on its clinical history, the herd was going through a clinical episode of respiratory disease. All air samples (inside and outside) tested positive for influenza A. Virus was isolated from 8 (7 inside and 1 outside) air samples. Virus was also isolated from 11 of 15 oral fluid samples. During the second farm visit, out of 15 air samples

collected inside the barn, 6 were classified as suspect. Out of 15 air samples collected outside, 2 tested positive and 1 was classified as suspect. All 15 oral fluid samples tested positive. No virus was isolated from these samples.

On farm 2, a wean-to-finish barn, with influenza clinical signs in the later stage of the disease, four (2 inside and 2 outside) air samples were classified as suspect. All 15 oral fluids tested positive for influenza A. No virus was isolated from these samples.

## Discussion

Our results confirm that acutely infected pig populations do generate airborne influenza A virus viable particles capable of being exhausted from pig barns and likely disseminated to other farms in the vicinity. Additionally, acutely infected populations can generate viable particles for at least a week after infection. Detection of influenza A virus in the field will depend on the course of disease, as it was previously demonstrated.<sup>5</sup> More studies are needed to further understand regional airborne transmission of influenza A virus.

## Acknowledgements

This study was partially funded by the Rapid Agricultural Response Fund, Minnesota Agricultural Experiment Station and the National Institute of Allergy and Infectious Diseases, National Institutes of Health, Department of Health and Human Services, under Contract No. HH-SN266200700007C. Its contents are solely the responsibility of the authors and do not necessarily represent the official views of the NIH.

## References

1. Brown, I. H. 2000. The epidemiology and evolution of influenza viruses in pigs. *Vet. Microbiol.* 74:29–46.
2. Mohan, R., Saif, Y.M., Erickson, G.A., Easterday, B.C. 1980. Serologic and epidemiologic evidence of infection in turkeys with an agent related to the swine influenza virus. *Avian Dis.* 25:11–16.
3. Poljak, Z., C. E. Dewey, S. W. Martin, J. Christensen, S. Carman, and R. M. Friendship. 2008. Prevalence of and risk factors for influenza in southern Ontario swine herds in 2001 and 2003. *Can. J. Vet. Res.* 72:7–17.

**C.A. Corzo; M. Torremorell; S. Dee; M. Gramer; R. Morrison**

4. Spackman, E., D. A. Senne, T. J. Myers, L. L. Bulaga, L. P. Garber, M. L. Perdue, K. Lohman, L. T. Daum, and D. L. Suarez. 2002. Development of a real-time reverse transcriptase PCR assay for type A influenza virus and the avian H5 and H7 hemagglutinin subtypes. *J. Clin. Microbiol.* 40:3256–3260.
5. Corzo, C., Romagosa, A., Gramer, G., Dee, S., Morrison, R., Torremorell, M. 2011. Detection of influenza virus in air from artificially generated aerosols and experimentally infected pigs. *AASV*. 43–44.

