



**North American Spray-Dried Blood and Plasma Producers (NASDBPP)
and
European Animal Protein Association (EAPA)**

Operating production facilities within China

Statement concerning African Swine Fever - Technical Version

Safety of Spray-Dried Porcine Blood Products

The Chinese Center for Disease Control and Prevention recently reported several cases of African Swine Fever (ASF). In each case, the herd animals were slaughtered and movement of pigs in and out of the affected areas was banned along with the feeding of untreated food waste (SHIC August 23, 2018). **African Swine Fever does not infect or harm humans (FAO, 2017).**

African swine fever virus (ASFV) is an enveloped dsDNA virus that can cause high mortality (up to 100%) in all ages of swine. The ASFV is primarily transmitted by oral-nasal pig to pig contact, through excretions from infected pigs, or from ingestion of pork or other contaminated products (swill, waste, carcasses, etc.) containing the virus (FAO, 2017). Other transmission pathways are indirect contact through fomites or vector-borne insects.

NASDBPP and EAPA spray-dried blood products are safe ingredients for use in animal feed.

Spray-dried porcine plasma (SDPP) is used globally in feed to enhance weaned pig performance (Torrallardona, 2010) and can effectively reduce morbidity and mortality of growing pigs during outbreaks of diseases like porcine reproductive and respiratory syndrome (PRRS), porcine circovirus 2 (PCV-2)-systemic disease and swine influenza (Messier et al., 2007). A research review (Pérez-Bosque et al., 2016) reported that spray-drying inactivates several swine viruses including PRRS virus, pseudorabies virus (PRV), swine vesicular disease virus (SVDV) and porcine epidemic diarrhea virus (PEDV).

Heating at 56°C for 70 minutes or 60°C for 20 minutes inactivates the ASF virus (OIE: ASF Technical Swine Disease card). Also, heat treatment of **80°C throughout substance** of meat is recognized to **inactivate many viruses** including Foot and Mouth Disease, Classical Swine Fever, Swine Vesicular Disease, **African Swine Fever**, Avian Influenza, Newcastle Disease, Rinderpest, and Sheep and Goat Plague (EU Directive 2002/99/EC, Annex III).

The **NASDBPP and EAPA** spray-dried blood products for use in animal feed are heated to **80°C throughout substance** during the manufacturing process.

Research has shown that spray-drying inactivates several swine viruses including African Swine Fever. In the table below tested viruses were grown in culture, inoculated in liquid plasma, and then spray-dried. Specific virus detection procedures for each of the tested virus were completed for the inoculated liquid and spray-dried plasma, and no viable virus was detected for PRRSV, PRV, PEDV or SVDV, indicating that these viruses were completely inactivated by spray-drying.

| Virus | Envelope | Thermal Stability | Inactivation Logarithm | Reference |
|-------|----------|-------------------|------------------------|------------------------------|
| PRRSV | Yes | Low | > 4.0 | Polo et al., 2005 |
| PRV | Yes | Medium | > 5.0 | Polo et al., 2005 |
| PEDV | Yes | Low | > 5.2 | Pujols & Segales, 2014 |
| PEDV | Yes | Low | > 3.6 | Gerber et al., 2014 |
| SVDV | No | Medium | > 6.0 | Pujols et al., 2007 |
| ASF | Yes | High | 4.1 | CRISA, 2018 unpublished data |

The **World Health Organization** (WHO, 2004) recognizes that a reduction of 4 logs (logarithms) of virus assures safety of human plasma products used for transfusions. Inactivation means that the virus is killed, removed or not capable of replicating. A 4-logarithm reduction of virus is considered equivalent to inactivation of 99.99% of the virus.

A recently completed study (CRISA, 2018) was conducted to determine the effects of commercial spray-drying conditions on ASFV. Liquid concentrated porcine plasma (28% solids) was inoculated with ASFV (Strain Badajoz 1971 – BA71) to a final median tissue culture infective dose (TCID₅₀) of 10^{5.77} per mL of liquid concentrated plasma. Triplicate 0.5 kg samples of inoculated plasma were spray-dried using a laboratory spray-dryer (Büchi 290 Mini Spray Dryer) at an inlet temperature of 200°C and at an outlet temperature of 80°C. Both liquid and spray-dried samples were analyzed for ASFV infectivity in VERO cell monolayers using a titer assay procedure on 25cm² flasks and subjected to 2 consecutive serial passages to new VERO cell culture. After the second passage the cells were analyzed by immune peroxidase monolayer assays (IPMA) against ASFV antigens to determine the amount of virus inactivation. **Titration results determined that spray-drying under commercial conditions inactivated 4.11 ± 0.20 log₁₀ TCID₅₀/mL of the inoculated ASFV.**

In conclusion, the spray-drying process typical of commercial conditions (80°C throughout its substance) for spray-dried porcine blood products was able to inactivate 4 logarithms (99.99%) of the inoculated ASFV. This result aligns with the EU Directive 2002/99/EC, Annex III, which recognizes that heat treatment of 80°C throughout substance of meat inactivates many viruses, including ASFV. In fact, inactivation of 4 logs of a pathogen is considered an effective safety assurance process for reducing risk of transmission in food or feed products according to guidelines of the WHO (2004).

The scientific evidence indicates that spray-dried porcine blood products are safe and that the spray-drying process effectively inactivates tested swine viruses, including African Swine Fever.

Additional Safety Features:

Members of NASDBPP and EAPA, and their respective plants in China, manufacture spray-dried blood products with several independent features that contribute to the safety of blood products.

1. Blood is **ONLY** collected from healthy animals to be slaughtered for human consumption.
2. Collected blood is pooled from multiple animals which contributes to a dilution effect. **Pooling is a recognized biosafety feature for human plasma products used for transfusions.**
3. The spray-drying process at high temperatures (**80°C throughout its substance**) has been shown and is accepted as effective in inactivating heat resistant viruses, including non-enveloped or enveloped viruses (**EU Directive 2002/99/EC, Annex III**).

Therefore, the spray-drying processes used by the NASDBPP & EAPA members globally and within China, for spray-drying blood products are aligned with the WHO guidelines providing several independent safety features that assure that the final products are safe from pathogens of concern for the swine industry including ASF virus.

References:

SHIC – Swine Health Information Center (<https://www.swinehealth.org>).

FAO. Beltrán-Alcrudo, D., Arias, M., Gallardo, C., Kramer, S. & Penrith, M.L. 2017. *African swine fever: detection and diagnosis – A manual for veterinarians*. FAO Animal Production and Health Manual No. 19. Rome. Food and Agriculture Organization of the United Nations (FAO). 88 pages.

Torrallardona, D. 2010. Spray dried animal plasma as an alternative to antibiotics in weanling pigs – A review. *Asian-Aust. J. Anim. Sci.* 23(1):131-148.

Messier, S., C. Gagne-Fortin, and J. Crenshaw. 2007. Dietary spray-dried porcine plasma reduces mortality attributed to porcine circovirus associated disease syndrome. *Proc. Amer. Assoc. Swine Vet.* p 147-150.

Pérez-Bosque, A., J. Polo, and D. Torrallardona. 2016. Spray dried plasma as an alternative to antibiotic in piglet feeds, mode of action and biosafety. *Porcine Health Management.* 2:16. doi:10.1186/s40813-016-0034-1.

OIE Swine Disease Card:

https://www.oie.int/fileadmin/Home/eng/Animal_Health_in_the_World/docs/pdf/Disease_cards/AFRICA_N_SWINE_FEVER.pdf.

EU - The Council of the European Union. Council Directive 2002/99/EC. Annex III. 2002;http://www.vet.gov.ba/pdf/files/eu_leg/anheu02.pdf.

Polo, J., J. D. Quigley, L. E. Russell, J. M. Campbell, J. Pujols, and P. D. Lukert. 2005. Efficacy of spray drying to reduce infectivity of Pseudorabies and PRRS viruses and seroconversion in pigs fed diets containing spray-dried animal plasma. *J. Anim. Sci.* 83:1933-1938.

Pujols J and J. Segalés. 2014. Survivability of porcine epidemic diarrhea virus (PEDV) in bovine plasma submitted to spray drying processing and held at different time by temperature storage conditions. *Vet Microbiol* 174:427-432.

Gerber, P. F., C. T. Xiao, Q. Chen, J. Zhang, P. G. Halbur, and T. Opriessnig. 2014. The spray-drying process is sufficient to inactivate infectious porcine epidemic diarrhea virus in plasma. *Vet. Microbiol.* 174:86-92.

Pujols, J. R. Rosell, L. Russell, J. Campbell, J. Crenshaw, E. Weaver, C. Rodriguez, J. Rodenas, and J. Polo. 2007. Inactivation of swine vesicular disease virus in porcine plasma by spray drying. *Proc. Amer. Assoc. Swine Vet.*, p 281-283.

WHO - World Health Organization 2004. WHO Technical Report Series No. 924, 2004. Annex 4. Guidelines on viral inactivation and removal procedures intended to assure the viral safety of human blood plasma products.