


A review of strategies to impact swine feed biosecurity

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Review

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Abstract

Global pork production has largely adopted on-farm biosecurity to minimize vectors of disease transmission and protect swine health. Feed and ingredients were not originally thought to be substantial vectors, but recent incidents have demonstrated their ability to harbor disease. The objective of this paper is to review the potential role of swine feed as a disease vector and describe biosecurity measures that have been evaluated as a way of maintaining swine health. Recent research has demonstrated that viruses such as porcine epidemic diarrhea virus and African Swine Fever Virus can survive conditions of transboundary shipment in soybean meal, lysine, and complete feed, and contaminated feed can cause animal illness. Recent research has focused on potential methods of preventing feed-based pathogens from infecting pigs, including prevention of entry to the feed system, mitigation by thermal processing, or decontamination by chemical additives. Strategies have been designed to understand the spread of pathogens throughout the feed manufacturing environment, including potential batch-to-batch carryover, thus reducing transmission risk. In summary, the focus on feed biosecurity in recent years is warranted, but additional research is needed to further understand the risk and identify cost-effective approaches to maintain feed biosecurity as a way of protecting swine health.

Pathogenic bacteria in swine feed and ingredients

Biological hazards that may be pathogenic to swine health include bacteria, such as *Salmonella* spp. and *Escherichia coli*, and viruses, such as PEDV, ASFV, SVA, Classical Swine Fever Virus (CSFV), Pseudorabies Virus (PRV), and Foot and Mouth Disease (FMD). These hazards differ in chemical and molecular structure, and therefore their prevalence may differ in feedstuffs. However, fecal contamination may lead to entry of many of these pathogens into ingredients, and the type of feedstuff and manner of contamination event may impact their survivability in feed and infectivity in swine.

Of the potential biological hazards in feed, *Salmonella* spp. is the most researched and understood. Feed-based transmission of *Salmonella* has been demonstrated to impact swine health, including a feed-based outbreak of *Salmonella enterica* subsp. *enterica* serovar Cubana in Sweden (Österberg *et al.*, 2006). Furthermore, commercial feed was reported to have a high significance as a potential vehicle for *Salmonella* transmission in the United States (Molla *et al.*, 2010). Researchers found 3.6% of feed samples and 17.2% of fecal samples positive for *S. enterica* subsp. *enterica* 36 barns and more than 6500 pigs. Of the *Salmonella* isolates, more than half were genotypically related to similar phenotypes and patterns of antimicrobial resistance. Currently, the United States Food and Drug Administration (FDA) considers *S. enterica* serotype Choleraesuis as an adulterant in swine feed, but adulteration by other serotypes is evaluated on a case-by-case basis (FDA, 2013). While *Salmonella* spp. has been reported by the FDA to be present in approximately 8% of animal feeds, neither *Salmonella* Cubana nor Choleraesuis are in the top 25 most prevalent serotypes found by the agency during routine surveillance (Li *et al.*, 2012).

One of the emerging serotypes of concern for swine feed is *S. enterica* serovar 4,5,12:i:–, a monophasic variant of *S. enterica* serovar Typhimurium. This serotype was responsible for a recall of whole roaster hogs in the United States in 2016, and has been associated with resistance to many common antimicrobials (Moreno Switt *et al.*, 2009; Centers for Disease Control, 2016). In 2012, Li *et al.* reported the serotype was the sixth most prevalent serotype found in animal feeds, and the seventh most common serotype in human infections. In a recent survey of 11 United States feed mills, *S. enterica* serovar 4,5,12:i:– was found in the manufacturing environment of two different mills (Magossi *et al.*, 2019). Contaminated surfaces included the ingredient pit grating, floor dust in the ingredient receiving area, and floor dust in the

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control room (Magossi *et al.*, 2019). Due to its multidrug resistance and links to both pork safety and prevalence in feed mills, *S. enterica* serovar 4,5,12:i:– is likely the key *Salmonella* serotype to control through future feed biosecurity.

The presence of other pathogenic bacteria in swine feed is less established. Tulayakul *et al.* (2012) reported 17 of 24 nursery, finishing, and sow feed samples collected in central Thailand were positive for *E. coli*, but only one sample had >100 colony forming units (CFU) mL⁻¹. Doane *et al.* (2007) reported two of 24 United States swine feed samples contained *E. coli* O157:H7, both of which were obtained from the state of Washington. The recent survey of 11 United States feed mills described above also identified *E. coli* in one sample of finished swine feed (Magossi *et al.*, 2019).

Both *Salmonella* and *E. coli* belong to a family of bacteria called Enterobacteriaceae. Active surveillance of this bacteria family may act as an indicator of biosecurity compliance and even predict future outbreaks. Enterobacteriaceae and *Salmonella* spp. in the 11 feed mills by Magossi *et al.* (2019) are shown in Fig. 1. Most Enterobacteriaceae identified in feed or the manufacturing environment were generally non-pathogenic in nature, such as *Enterobacter* and *Citrobacter*. However, areas with high levels of Enterobacteriaceae also had high levels of *Salmonella* spp. (association $P=0.05$). Analysis of retained samples showed that worker shoes also carried Senecavirus A in one feed mill. When another mill that was part of the surveillance was associated with an outbreak of Porcine Deltacoronavirus, the virus was found in the load-out auger, cooler air intake, ingredient pit grating, all locations of floor dust, broom, and worker shoes. Enterobacteriaceae is commonly used to indicate hygiene and/or biosecurity compliance in human food, rendering, and poultry feed manufacturing facilities (Van Schothorst and Oosterom, 1984; Jones and Richardson, 2004; Nestle, 2014). The proactive monitoring of Enterobacteriaceae should be further evaluated and considered as a method to better identify and control the highest risk points of entry into the swine feed supply chain.

Pathogenic viruses in swine feed and ingredients

Research has demonstrated that viruses, such as PEDV, ASFV, SVA, CSFV, PRV, and FMD, are able to survive in at least some commonly imported feed ingredients (Dee *et al.*, 2018). Modeling performed to simulate the environmental conditions during transport of ingredients from China to the United States has shown that a viable PEDV sample is able to survive in certain ingredients, including soybean meal (both conventional and organic), vitamin D, lysine hydrochloride, and choline chloride (Dee *et al.*, 2016). In addition to PEDV, 11 other pathogens have been subjected to a similar modeling procedure in a variety of different ingredients (Dee *et al.*, 2018). The survivability of a pathogen varied depending on the genetic and physicochemical properties of the virus, and differed between pathogens and the feed ingredients tested. Certain feed ingredients or feed products presented a better matrix for virus survival than the others and select ingredient matrices seemed to enhance the survival of multiple viruses. For example, conventional soybean meal had a higher level of virus survival in comparison with organic soybean meal. The exact reason for this difference in survivability in sources of soybean meal is unknown, but could be attributed to the higher levels of fat present in the organic variety used in the trial, as there has been some evidence that medium chain fatty acid (MCFA) blends have viricidal effects (Cochrane,

2018). It has also been hypothesized that higher protein ingredients have greater capability of retaining viral infectivity, but the mechanism is not yet understood. Overall, laboratory simulations have indicated that certain feed ingredients exhibit a higher risk of transporting viral pathogens (Dee *et al.*, 2018). Additional research is needed to better understand what ingredient attributes are associated with enhanced survivability.

Infectivity of biological hazards in swine feed and ingredients

Once it has been established that biological hazards can survive in feed and ingredients, it is important to understand their infectivity at a dose that may cause infection. Infectivity frequently relies on ensuring the viral capsids or bacteria lipid membranes are intact as they protect the pathogen from deterioration during storage. Sufficient numbers of intact particles are needed to cause infection in otherwise healthy animals, and this is known as the minimum infectious dose. Loynachan and Harris (2005) first published the minimum infectious dose of *S. enterica* serovar Typhimurium in pigs as >10³ CFU g⁻¹ of feed. Cornick and Helgeson (2004) reported the infectious dose of *E. coli* O157:H7 is 6 × 10³ CFU g⁻¹ in 3-month old pigs. As Österberg *et al.* (2006) reported, infectious dose is difficult to determine, especially in bacteria, because challenge doses are strongly associated with fecal shedding, but weakly associated with infection.

Schumacher *et al.* (2016) reported the minimum infectious dose for PEDV-inoculated feed is 5.6 × 10¹ TCID₅₀, equivalent to antiprimer-based quantitative real-time polymerase chain reaction (aqRT-PCR) cycle threshold (CT) of 37.1. Notably, this was above the threshold of many PEDV PCR assays in diagnostic laboratories. This research helped demonstrate why PEDV was so easily spread through a feed matrix, as 1 g of feces from an acutely infected pig could infect 500 ton of feed, with all the feed being infected at a dose capable of causing illness.

Ongoing research focuses on determining the median infectious dose of African Swine Fever Virus in both feed and water (Niederwerder, 2018). Additional research is needed to determine the minimum or median infectious dose for a number of bacteria and viruses, including Enterotoxigenic *E. coli*, SVA, CSF, and PRV. These doses are necessary as they become targets for mitigation measures. While ideally there is no detectable pathogen in feed or ingredients, it must at least be prevented or reduced to levels below an infectious dose to sustain animal health.

Once biological hazards that are considered a risk have been identified, procedures should be created that prevent entry of the hazard into the mill, as well as procedures for mitigation and decontamination in case hazard entry cannot be prevented. Cochrane *et al.* (2016a, 2016b) published an overview of a feed mill biosecurity plan that can easily serve as the foundation for developing a mill-specific biosecurity plan. Some of their recommendations are highlighted below.

Preventing biological hazards in swine feed and ingredients

The most effective component of a feed mill biosecurity plan is prevention of hazard entry. There is incentive to prevent a hazard's entry into a facility altogether because it has been shown that the introduction of a contaminated material into a feed mill can lead to the mill being contaminated for an extended period (EFSA, 2008). Controlling the entry of biological hazards into a facility should begin with evaluation of the ingredient

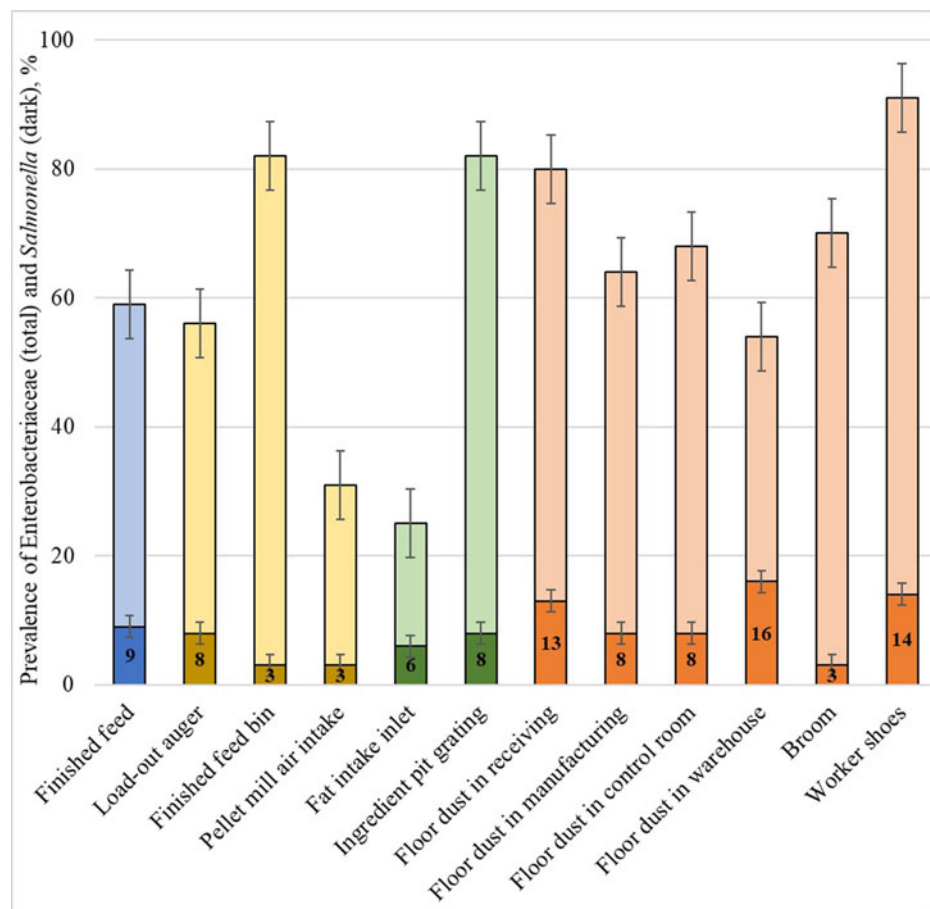


Fig. 1. Presence of Enterobacteriaceae in 11 United States feed mills. The levels of Enterobacteriaceae (total bars) vary across location, but are associated with *Salmonella* spp. (dark portion of bars) presence. High Enterobacteriaceae levels may indicate biosecurity compliance and even predict future outbreaks.

suppliers. The development of a supplier verification program that includes specific requirements for ingredients being purchased, as well as communicating safety expectations to the supplier of an inbound ingredient is an important step in preventing the entry of a biological hazard. This may also include verification of ingredient-supplier protocols and on-site manufacturing facility reviews and assessments. As mentioned in the previous section, some ingredients have the potential to maintain bacteria or virus survivability and infectivity more than others. As a result, the best way to prevent hazard entry into the mill is to eliminate high risk ingredients from diet formulations. Thus, coordinated efforts between nutritionists, formulators, purchasers, and the rest of the integrated feed supply team is essential for maintaining an effective feed mill biosecurity plan.

While having a supplier control program is an important step when controlling the entry of a biological hazard into a facility, routine sampling and analysis of bagged, bulk, or liquid ingredients that are considered high-risk for certain pathogens is a valuable tool. All samples' collection should be performed by using an aseptic method, as cross-contamination of samples during the collection process needs to be prevented. If an ingredient is considered high risk, every lot should be analyzed separately. If it is lower risk, it may be more practical to collect samples and pool them for more intermittent analysis as a way to reduce analytical cost. Determining and setting a schedule for sampling of ingredients that are considered higher risk, as well as defining an inventory holding procedure until analytical results are obtained can help lower the potential of a biological hazard being introduced into the mill. Traceability of ingredients is essential, and

maintaining records that indicate information such as the receiving date, time, lot number during unloading, and prior haul data that is connected to specific batches of finished feed allows for a quick response if a biological hazard is suspected.

Movement of people or vehicles in or out of a facility also has the potential to introduce biological hazards. Employees in the feed mill and visitors, such as guests, truck drivers, and subcontractors have the ability to introduce contaminants into a feed manufacturing facility. People may unknowingly carry fecal, dirt, or dust particles contaminated with undesirable microorganisms on the bottoms of their shoes or on clothing and are at a particularly higher risk if they are coming from another farm or feed mill where the hazard is present. The risk of people introducing biological hazards is easily illustrated in Fig. 1 (Magossi *et al.*, 2019), as 91% of samples collected from worker boots were contaminated with Enterobacteriaceae. Controlling and minimizing foot traffic across receiving pit grates or around hand-add port grates is a logical, low-cost method to reduce the risk of a biological pathogen being introduced into the manufacturing system, and can easily be accomplished by covering the grates when not in use. No-walk zones or even hygienic zoning may be appropriate to include in biosecurity plans in feed mills that have a higher risk of biological pathogen introduction. Procedures requiring that all visitors must be accompanied at all times by a trained employee can to help prevent biosecurity breaches. Visitors should be provided clean footwear, plastic boots, or boot covers to limit the entry of outside hazards. This includes the drivers of inbound trucks. Ideally, drivers should stay inside their trucks at all times to minimize foot traffic, especially over the receiving

grates. If the driver must exit the vehicle, wearing disposable plastic boots or covers will limit the potential of hazards being introduced from their shoes. Trucks entering the feed mill should have mud and sludge removed from the trailer opening before the vehicle reaches the receiving pit, and the pit should remain covered until the truck is ready to unload. Ingredients may be contaminated prior to unloading, but they may also be contaminated during the unloading process due to mud or floor sweepings intermingling with ingredients at the point of entry. Ensuring the receiving pit remains covered while trucks are being moved reduces the risk of contamination during unloading, which is important considering the impracticality of thoroughly cleaning conveying equipment such as the central pit or, bucket elevators. Use of cones and funneling devices can also be used to limit the quantity of material that spills during unloading and prevents mill employees from sweeping spilled ingredients into the pit.

Floor sweepings, including those from the unloading process, should be disposed of and not swept into the pit. In addition, many feed manufacturing facilities have grain cleaners and dust collection equipment in place, and it has been well established that dust and other screened particles can act as a carrier for biological hazards including PEDV (Gebhardt *et al.*, 2016) or mycotoxins (Yoder *et al.*, 2018), among others. Many feed manufacturers have the mentality that adding back the dust or screened material to the finished feed is acceptable because it will reduce ingredient shrink. However, the cost associated with reduced animal performance and/or increased mortality is much greater than the loss of mill efficiency, and therefore all dust and screened materials should always be disposed of compared to being added back into the feed.

Reducing biological hazards in swine feed and ingredients

Once a biological hazard is introduced into a facility, it can be almost impossible to control because most feed manufacturing facilities were not hygienically designed. Furthermore, mitigation strategies that may be possible in some systems may not work in others because of differences in facility design and equipment, manufacturing operations, and other associated risk factors among feed mills. For instance, Muckey (2016) reported that the surface type (concrete, plastic, rubber, stainless steel, etc.) impacts pathogen survivability in the presence of different decontamination procedures. Stainless steel and smooth plastic surfaces, while easier to clean than tires, rubber belts, or polyethylene totes, are more difficult to sanitize due to the formation of biofilms that protect the bacteria or virus from a chemical sanitizer. Therefore, both cleaning and sanitizing is often necessary, and nearly impossible based on current equipment design constraints.

Physical prevention of hazard spread via cross-contamination is especially difficult due to the highly infective nature of contaminated dust and the impracticality of physical clean-out in most mills (Fig. 2). In Schumacher *et al.* (2017), the role of PEDV cross-contamination in feed mills was evaluated. Initially, a PEDV-negative corn- and soybean meal-based nursery pig diet was mixed, conveyed, and discharged using pilot scale feed manufacturing equipment. Next, a diet was manufactured, including an ingredient that had been spiked with infectious PEDV. Subsequently, four separate PEDV-negative diets were mixed, conveyed, discharged to test how many negative diets were necessary to 'flush' contamination from the manufacturing surface. Environmental swabs were collected prior to and after each

batch of feed by swabbing direct feed contact surfaces, adjacent surfaces located within 1 m of manufacturing equipment, and other surfaces located at least 1 m away from manufacturing equipment. The presence of PEDV RNA was reported in CT of qRT-PCR using PROC GLIMMIX (SAS Institute, Inc., Cary, NC). The statistical model evaluated the effect of manufacturing sequence (negative, positive, flush 1, flush 2, flush 3, and flush 4) and location (direct feed contact, adjacent, or other surface) and the associated interaction. The LSMEANS procedure compared surface type among treatments within animal food-contact surfaces by pairwise comparison. Both main effects and the interaction were significant ($P < 0.05$). Subsequently, Gebhardt *et al.* (2018) demonstrated that dust collected from feed manufacturing surfaces can cause infectivity in a swine bioassay. Therefore, limiting and controlling dust created during manufacturing should be a priority, as it can serve as a vector in viral disease transmission such as PEDV. Sequencing procedures in order to minimize risk to the most sensitive phases of production should be utilized. Furthermore, flushing protocols should be established to help minimize cross contamination risk. Gebhardt *et al.* (2016) showed in a PEDV model that rice hull flushes can be a cost-effective strategy to reduce cross-contamination risk.

For RNA viruses in particular mitigation techniques depend on disrupting the viral capsid which removes the protective shell around the virus (Cliver, 2009). Three main categories of mitigation strategies have been identified and include biological, physical, and chemical. Deng and Cliver (1995) reported that biological inactivation typically occurs with the use of specific enzymes or other products of microbial origin that attack viruses or bacteria, but research is lacking to determine if this is a feasible mitigation strategy for the feed manufacturing industry. Physical inactivation in feed manufacturing is most commonly achieved thermally, but should be considered a point-in-time mitigation strategy, because it would not prevent post-processing contamination risk. The use of chemical agents, such as formaldehyde or medium-chain fatty acids as feed additives has been shown to have excellent potential to inhibit virus and bacterial hazards in feed. The benefit of these chemical agents is that they have the potential to have immediate as well as residual efficacy which could help with mitigation from the point of application until the time the feed is consumed. Specific research identifying mitigation strategies that can be used in the feed manufacturing process are reviewed below.

Thermal Processing: In a benchtop model, Goyal (2013) confirmed that PEDV is a heat-sensitive virus and that a temperature \times time relationship could be used as a guide for PEDV inactivation. Based on this information, two studies were conducted to determine if passing feed through a pellet mill would be sufficient to apply thermal insult to a great enough extent to prevent PEDV infectivity. Cochrane *et al.* (2017) showed in the first trial that when a low or high dose of PEDV was used to inoculate feed, with the resulting feed subsequently processed at one of nine combinations of conditioning temperature (68, 79, or 90 °C) or conditioner retention time (45, 90, or 180 s) all processed batches of feed were unable to generate infectivity in a pig bioassay model, even though the unprocessed feed did lead to PEDV infectivity. In a subsequent trial, the same researchers processed feed through a conditioner utilizing a 30 s retention time and one of five condition temperatures (38, 46, 54, 63, or 71 °C) and observed that feed processed at or above 54 °C was able to prevent PEDV infectivity, while feed that was processed at the two low temperatures did lead to PEDV infection when fed to pigs. This

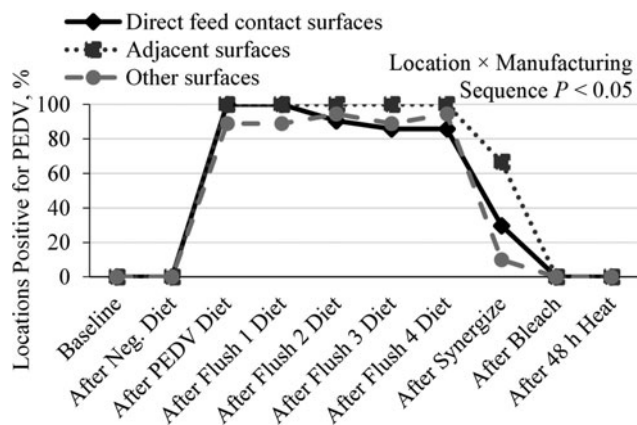


Fig. 2. Schumacher *et al.* (2017) reported the role of PEDV cross-contamination in feed mills. Initially, a PEDV-negative swine diet was manufactured, followed by a diet that included an ingredient that had been spiked with infectious PEDV, then four subsequent PEDV-negative diets in an attempt to ‘flush’ contamination from the manufacturing surfaces. Environmental swabs were collected prior to and after each batch of feed by swabbing direct feed contact surfaces, adjacent surfaces located within 1 m of manufacturing equipment, and other surfaces located at least 1 m away from manufacturing equipment. Both main effects and the interaction were significant ($P < 0.05$). Subsequently, Gebhardt *et al.* (2018) demonstrated that dust collected from feed manufacturing surfaces can cause infectivity in a swine bioassay. Therefore, limiting and controlling dust created during manufacturing should be a priority, as it can serve as a vector in viral disease transmission such as PEDV. Sequencing procedures in order to minimize risk to the most sensitive phases of production should be utilized.

series of trials demonstrated that thermal mitigation is a possible means of minimizing PEDV-associated risk, and more importantly demonstrated that equipment commonly found in commercial feed mills was effective at applying the thermal stress. However, it is important to remember that even though the feed mill may target a specific processing temperature adequate to inactivate PEDV, there are times during the feed manufacturing process (such as at equipment startup, or if steam flow is turned off to ameliorate a plugged die) that the feed may not be processed at a high enough temperature to effectively eliminate all virus transmission risk. Furthermore, the research demonstrates that the pellet mill is an effective point-in-time mitigation strategy, but it cannot prevent post-processing recontamination risk.

Residual control measures

The use of chemical feed additives as strategies to reduce biological hazards in feed is appealing because they allow for efficacy throughout the remainder of the feed supply chain, with the potential to also influence animal performance once consumed. As a result, a number of different products have been tested as chemical-based feed hazard mitigants. Some compounds that have shown mixed efficacy at reducing or eliminating virus or bacterial risk include organic acids (Eklund 1985), essential oils (Orhan *et al.*, 2012), sodium bisulfate (Knueven, 1998), or sodium chlorate (Smith *et al.*, 2012); however, the cumulative data suggest that the effectiveness of any chemical-based feed mitigant is not only target specific but also feed ingredient/matrix specific (Cochrane, 2018). Of all the potential chemical mitigants available, the two that have garnered the most commercial interest are formaldehyde and MCFAs.

Formaldehyde has been shown to be effective at preventing risk associated with PEDV (Dee *et al.*, 2014, 2015; Cochrane,

2018) as well as *Salmonella* (Cochrane *et al.*, 2016a, 2016b). However, regulatory restrictions can limit some applications as the product is only approved for use to prevent contamination with *Salmonella*. Additionally, specialized equipment must be used for accurate application, and there are worker health concerns as well as negative perception by some consumers, which can lead to formaldehyde being limited in its commercial application. Furthermore, the use of formaldehyde in feed may lead to detrimental bacterial shifts in the pig gut (Williams *et al.*, 2018).

The use of MCFAs as chemical-based feed mitigants was reviewed by Cochrane *et al.* (2018). They observed that MCFA are effective at preventing risk associated with feed contaminated with PEDV in addition to their effectiveness against *Salmonella* (Cochrane *et al.*, 2016a, 2016b). Through a series of trials this group of researchers has shown that combinations of caproic, caprylic, and capric acid are the most effective with little efficacy of lauric acid against PEDV. Interestingly, the same group of researchers also showed that increasing concentrations of a 1:1:1 blend of caproic, caprylic, and capric acid also resulted in a linear increase in growth performance with a 1.50% inclusion resulting in an almost 2 kg BW advantage compared to a diet with no MCFA after feeding nursery pigs for 35 days (Thomson *et al.*, 2018). Furthermore, Gebhardt *et al.* (2018) showed that feed used in this trial that was collected 40 days after MCFA application was still successful at reducing PEDV risk which demonstrates the residual mitigation potential of MCFA.

Addressing feed mills contaminated with biological hazards

Due to the high quantity of airborne particulates in animal food manufacturing facilities, dust contamination is a widespread mechanism for both viral and bacterial hazard transmission (Fig. 2). This can be specifically challenging because of the difficulties associated with physical cleaning (Muckey, 2016). Highly aggressive procedures, such as the use of liquid chemical sanitizers and heat have been shown to be necessary when reducing bacteria on environmental surfaces to completely decontaminate manufacturing surfaces (Fig. 2; Huss *et al.*, 2017; Schumacher *et al.*, 2017). Effective cleaning, which may require both physical cleaning and the use of cleaning solutions, removes biofilm formations that will allow for subsequent penetration and removal of vegetative bacteria by a sanitizer. Both steps are necessary, but can prove to be difficult in many feed manufacturing systems due to a lack of access or ability to thoroughly clean out or safely sanitize dry bulk manufacturing systems. Cleaning of non-animal food-contact surfaces should not be overlooked as biological hazards can efficiently spread throughout a facility through dust and other airborne particulates. This contamination is not mitigated during flushing procedures, and can contaminate subsequent feed batches (Schumacher *et al.*, 2017).

Because complete physical clean-out of feed manufacturing systems can prove to be difficult, flushing procedures including the use of added substances such as formaldehyde, MCFA, and dry essential oil blends may be used to help reduce the presence of biological hazards on feed-contact surfaces. Data suggest that biological hazard risk can be reduced after a third flush, or after the use of a chemically enhanced flush (Gebhardt *et al.*, 2016; Muckey, 2016; Schumacher *et al.*, 2017). Formaldehyde-based products and an MCFA blend have been shown to reduce the presence of PEDV on these surfaces when used in conjunction with a rice hull flush. Similarly, MCFA blends have been found to be effective at reducing *S. enterica* serovar Typhimurium on

stainless steel surfaces, in addition to reducing the quantity of post-processing *S. enterica* serovar Typhimurium contamination if 2% is applied to swine feed prior to its inoculation with bacteria (Cochrane, 2016a).

Future directions for swine feed safety

Clearly, additional research is necessary to better understand both the risk and prevention of biological hazards in swine feed and ingredients. Our knowledge of survivability, infectivity, mitigation, and decontamination strategies all must be improved to maintain the safety of swine feed in the future. Additional research is warranted to evaluate the role of beneficial bacteria to competitively exclude pathogens in feed manufacturing environments, such as those described by Zhao *et al.* (2013) for controlling *Listeria monocytogenes* and *Listeria* in poultry processing plants. The concept of competitive bacteria for inhibitory exclusion is being tested for controlling *Salmonella* and other pathogenic bacteria in meat processing and rendering facilities, and may be viable to consider for feedstuff production facilities. Furthermore, it will be important to understand hygienic design for retrofits and new construction of feed mills in the future.

One of the items that limits the ability to make faster progress on feed safety is that few molecular diagnostics methods have been appropriately validated for feedstuffs. Our team has consistently witnessed lower recovery rates of viral nucleic acids when moving from inoculant (virus stock) into dry feed or ingredients. We reported this challenge in Schumacher *et al.* (2016), where we established the infectious dose of PEDV via feed. Increasing levels of virus were associated with lower qRT-PCR CT in the inoculum. However, the CT value of the same virus dose in feed was, on average, 9.5 CT higher than that in the inoculum after correcting for the dilution. The loss in viral RNA and diagnostic sensitivity did not appear to be dose-related, since a similar CT was detected despite different amounts of PEDV were spiked into the feed. This was later confirmed by Cochrane *et al.* (2017), where we reported a consistent loss in PEDV detectability by RT-PCR when feed was inoculated with a low (10^3 TCID₅₀ g⁻¹) or high (10^5 TCID₅₀ g⁻¹) dose of PEDV. Tissue culture medium inoculum was 20 or 13 CT, while the detectable PEDV in feed was 30.7 or 23.9 CT, leading to a 10.7 or 10.9 CT loss in sensitivity for both the low and high PEDV doses, respectively. This loss in viral RNA and diagnostic sensitivity is not isolated to our laboratories. A similar loss in sensitivity was found by Iowa State University, when our samples were tested in their diagnostic laboratory. In our collaboration with Pipestone Systems and SDSU, we (Dee *et al.*, 2016) reported an up to 11.4 CT loss when moving from stock virus to feed or ingredients using diagnostic assays developed at the SDSU Veterinary Diagnostic Laboratory. The challenge in loss of sensitivity is not confined to PEDV. We recently observed a similar CT sensitivity loss between liquid inoculum and dry feed with Senecavirus A (SVA) (Sardella *et al.*, 2019). Furthermore, Dee *et al.* (2018) reported a loss in sensitivity between stock virus and various ingredients or complete feed for SVA, BVDV (surrogate for CSFV), PRRSV, and ASFV. Finally, the loss in diagnostic sensitivity appears to vary from one feed or ingredient matrix to another. Cochrane *et al.* (2016a, 2016b) demonstrated that complete swine diet, blood meal, meat and bone meal, and spray-dried porcine plasma inoculated with the same quantity of PEDV inoculum resulted in CTs ranging from 26 to 31 using identical sample preparation, extraction and RT-PCR conditions. Furthermore, Dee *et al.* (2016) reported a loss of sensitivity

ranging from 1.4 to 11.4 CT for PEDV, depending upon the ingredient matrix.

We initially accepted this loss in sensitivity was inherently part of the assay, and that it posed a problem for veterinary diagnostic laboratories, but had limited biological relevance. However, we have recognized that the poor recovery of nucleic acids using the current methods in ingredients has substantial ramifications: it leads to false negative results via a type-II error. Natural fecal contamination in ingredients or from cross-contamination due to poor biosecurity is likely to have low levels of virus, and a 10-CT reduction in sensitivity may lead to the determination that a contaminated product is actually safe. For example, we reported in Schumacher *et al.* (2018) that a pig gavaged with a PEDV-spiked feed sample that was qRT-PCR-negative (>40 CT) became infected and symptomatic, i.e. presence of infectious PEDV was confirmed in the bioassay.

Presently, the current methods for sample preparation, extraction, and detection of nucleic acids in feedstuffs are too variable and pose too high of a risk for a false negative, which negates the value of the test. Urgent research is needed to validate molecular detection methods in feedstuffs, which can then be used to create an appropriate sampling method and point-of-use diagnostic devices. Until then, environmental monitoring and product testing is not a viable option for ensuring feed safety; instead efforts must be more preventative than reactive. The swine feed industry must embrace feed biosecurity as regulators and consumers shift their thinking of our product as swine feed to swine food.

Recommendations to maximize swine feed biosecurity

In conclusion, biosecurity is a well-known topic at the farm level, but only recently has begun to gain importance in the feed manufacturing process. Evidence demonstrating the ability of feed and feed ingredients to support virus infectivity and bacterial survivability has been collected which points to the fact that feed and ingredients can be a vector for biological hazard transmission. Consequently, a series of steps should be taken to help maximize feed biosecurity:

- (1) Assess biological hazard risk: feed manufacturing facilities must take a proactive approach to understanding biological hazards for their own operations and the security of their customers. The biosecurity procedures employed by a specific mill may not be the same as other mills depending on the customers they serve and the associated risk tolerance versus price for mitigation strategies that are employed.
- (2) Define protocols to prevent entry of hazard into the mill: the most important part of a feed mill biosecurity plan is to prevent hazards from entering the mill. Identifying and eliminating high risk ingredients, minimizing entry via people and equipment, covering all open points of entry when not being used, and other strategies can be used to prevent hazard entry into the mill.
- (3) Utilize mitigation strategies to prevent risk: not all hazards can be prevented from entering the mill and consequently mitigation strategies should be utilized. The best option is to identify the mitigation strategies that are effective against the specific hazards of concern and utilize a combination of point-in-time mitigants as well as those that have residual effectiveness for continue protection through the remainder of the feed supply chain. Some mitigation strategies have multiple benefits. As an example, dust collection and elimination

not only create a safer and better environment for the workers, but also can eliminate a major point of contamination.

- (4) Feed mill decontamination: while it is extremely difficult to completely accomplish, a feed mill decontamination strategy must be developed and should include a combination of physical cleaning, chemical cleaning, and if applicable the use of high heat as the final step.

References

- Centers for Disease Control** (2016) *Multistate Outbreak of Multidrug-Resistant Salmonella I 4,[5],12:i:- and Salmonella Infantis Infections Linked to Pork*. Final update report. <https://www.cdc.gov/salmonella/pork-08-15/index.html>.
- Cliver DO** (2009) Capsid and infectivity in virus detection. *Food and Environmental Virology* **1**, 123–128.
- Cochrane RA** (2018) *Interventional Strategies to Reduce Biological Hazards in Animal Feeds* (PhD Dissertation). Kansas State University, Manhattan.
- Cochrane RA, Huss AR, Aldrich GC, Stark CR and Jones CK** (2016a) Evaluating chemical mitigation of *Salmonella* Typhimurium ATCC 14028 in animal feed ingredients. *Journal of Food Protection* **79**, 672–676.
- Cochrane RA, Dritz SS, Woodworth JC, Stark CR, Huss AR and Jones CK** (2016b) Feed mill biosecurity plans: a systematic approach to prevent biological pathogens in swine feed. *Journal of Swine Health & Production* **26**, 154–164.
- Cochrane RA, Schumacher LL, Dritz SS, Woodworth JC, Huss AR, Stark CR, DeRouchey JM, Tokach MD, Goodband RD, Bai J, Chen Q, Zhang J, Gauger PC, Derscheid RJ, Magstadt DR, Main RG and Jones CK** (2017) Effect of pelleting on survival of Porcine Epidemic Diarrhea Virus (PEDV)-contaminated feed. *Journal of Animal Science* **95**, 1170–1178.
- Cochrane RA, Amachawadi RG, Remfry S, Lerner AB, Gebhardt JT, Nagaraha TG, Pluske JR, Niederwerder MC, Woodworth JC, Dritz SS and Jones CK** (2018) A review of medium chain fatty acids and their recent role in feed safety. *Journal of Animal Science* **96**, 55.
- Cornick NA and Helgerson AF** (2004) Transmission and infectious dose of *Escherichia coli* O157:H7 in swine. *Applied and Environmental Microbiology* **70**, 5331–5335.
- Dee S, Neil C, Clement T, Christopher-Hennings J and Nelson E** (2014) An evaluation of a liquid antimicrobial (Sal CURB™) for reducing the risk of porcine epidemic diarrhea virus infection of naïve pigs during consumption of contaminated feed. *BMC Veterinary Research* **10**, 220.
- Dee S, Neill C, Clement T, Singrey A, Christopher-Hennings J and Nelson E** (2015) An evaluation of porcine epidemic diarrhea virus survival in individual feed ingredients in the presence or absence of a liquid antimicrobial. *Porcine Health Management* **1**, 9.
- Dee S, Neill C, Clement T, Nelson E, Singrey A, Christopher-Hennings J, Jones CK, Cochrane RA, Patterson G and Spronk G** (2016) Modeling the transboundary risk of feed ingredients. Contaminated with porcine epidemic diarrhea virus. *BMC Veterinary Research* **12**, 51–63.
- Dee S, Bauermann R, Niederwerder ME, Singrey A, Clement T, DeLima M, Long C, Patterson G, Shehan M, Stoian A, Petrovan V, Jones CK, De Jong J, Ji J, Spronk G, Hennings J, Zimmerman J, Rowland B, Nelson E, Sundberg P, Diel D and Minion L** (2018) Survival of viral pathogens in animal feed ingredients under transboundary shipping models. *PLoS One* **13**, e0194509.
- Deng MY and Cliver DO** (1995) Antiviral effects of bacteria isolated from manure. *Microbial Ecology* **30**, 43–45.
- Doane CA, Pangloli P, Richard HA, Mount JR, Golden DA and Draughon FA** (2007) Occurrence of *Escherichia coli* O157:H7 in diverse farm environments. *Journal of Food Protection* **70**, 6–10.
- Eklund T** (1985) Inhibition of microbial growth at different pH levels by bezoic and propionic acids and esters of *p*-hydroxybenzoic acid. *International Journal of Food Microbiology* **2**, 159–167.
- European Food Safety Authority** (2008) Scientific opinion of the panel on biological hazards on a request from the health and consumer protection, directorate general, European commission on microbiological risk assessment in feeding stuffs for food producing animals. *EFSA Journal* **720**, 1–84.
- FDA** (2013) *Compliance Policy Guide Sec. 690.800 Salmonella in Food for Animals*. United States Department of Health and Human Services Center for Veterinary Medicine Office of Regulatory Affairs. Rockville, MD.
- Gebhardt JT, Woodworth JC, Jones CK, Gauger PC, Tokach MD, DeRouchey JM, Goodband RD, Muckey MB, Cochrane RA, Niederwerder M, Stark CR, Bai J, Chen Q, Zhang J, Ramirez A, Derscheid R, Main R and Dritz SS** (2016) Evaluation of the effects of flushing feed manufacturing equipment with chemically-treated rice hulls on porcine epidemic diarrhea virus cross contamination during feed manufacturing. *Kansas Agricultural Experiment Station Research Reports* **2**, 12–82.
- Gebhardt JT, Woodworth JC, Tokach MD, DeRouchey JM, Goodband RD, Jones CK and Dritz SS** (2018) Medium chain fatty acid mitigation activity against porcine epidemic diarrhea virus (PEDV) in nursery pig diets after 40 d of storage. *Journal of Animal Science* **96**, 153.
- Goyal S** (2013) Environmental stability of PEDV (porcine epidemic diarrhea virus). *Allen D. Leman Swine Conference*, St. Paul, MN, 14 September 2013.
- Huss AR, Schumacher LL, Cochrane RA, Poulsen E, Bai J, Woodworth JC, Dritz SS, Stark CR and Jones CK** (2017) Elimination of porcine epidemic diarrhea virus in an animal feed manufacturing facility. *PLoS One* **12**, e0169612.
- Jones FT and Richardson KE** (2004) *Salmonella* in commercially manufactured feeds. *Poultry Science* **83**, 384–391.
- Knueven C** (1998) U.S. Patent No. 5773063. Washington, DC: U.S. Patent and Trademark Office.
- Li X, Bethune LA, Jia Y, Lovell RA, Proescholdt TA, Benz SA, Schell TC, Kaplan G and McChesney DG** (2012) Surveillance of *Salmonella* prevalence in animal feeds and characterization of the *Salmonella* isolates by serotyping and antimicrobial susceptibility. *Foodborne Pathogens and Disease* **9**, 692–698.
- Loynachan AT and Harris DL** (2005) Dose determination for acute *Salmonella* infection in pigs. *Applied Environmental Microbiology* **71**, 2753–2755.
- Magossi G, Cernicchiaro N, Dritz S, Houser T, Woodworth J, Jones C and Trinetta V** (2019) Evaluation of *Salmonella* presence in selected United States feed mills. *MicrobiologyOpen* **8**, e00711.
- Molla B, Sterman A, Mathews J, Artuso-Ponte V, Abley M, Farmer W, Rajala-Schultz P, Morrow WEM and Gebreyes WA** (2010) *Salmonella enterica* in commercial swine feed and subsequent isolation of phenotypically and genotypically related strains from fecal samples. *Applied Environmental Microbiology* **76**, 7188–7193.
- Moreno Switt AI, Soyer Y, Warnick LD and Wiedmann M** (2009) Emergence, distribution, and molecular and phenotypic characteristics of *Salmonella enterica* serotype 4,5,12:i:-. *Foodborne Pathogens and Disease* **6**, 407–415.
- Muckey MB** (2016) *Evaluation of Surface Sanitation to Prevent Biological Hazards in Animal Food Manufacturing* (MS Thesis). Kansas State University, Manhattan.
- Nestle** (2014) *Microbiological Specifications*. St. Louis, MO. Switzerland: Nestec Ltd., Co. <https://www.nestle.com/asset-library/documents/library/documents/suppliers/microbiological-specifications-nestle.pdf>.
- Niederwerder MC** (2018) Median infectious dose of ASFV in swine feed. *American Association of Swine Veterinarians Committee on Emerging and Transboundary Diseases Meeting*, 3 March 2018, Manchester Grand Hyatt Hotel, San Diego, CA. Presentation.
- Orhan IE, Özçelik B, Kartal M and Kan Y** (2012) Antimicrobial and antiviral effect of essential oils from selected Umbelliferae and Labiatae plants and individual essential oil components. *Turkish Journal of Biology* **36**, 239–246.
- Österberg J, Vågsholm I, Boqvist S and Sternberg Lewerin S** (2006) Feed-borne outbreak of *Salmonella* Cubana in Swedish pig farms: risk factors and factors affecting the restriction period in infected farms. *Acta Veterinaria Scandinavica* **47**, 13–21.
- Sardella CA, Petrovan V, Davis SK, Stewart SS, Niederwerder MC, Woodworth JC, Dritz SS, Rowland R and Jones CK** (2019) Validation of environmental swabbing to detect Senecavirus A in feed. *Journal of Animal Science* **97**, 245–246.
- Schumacher LL, Woodworth JC, Jones CK, Chen Q, Zhang J, Gauger PC, Stark CR, Main RG, Hesse RA, Tokach MD and Dritz SS** (2016)

- Evaluation of the minimum infectious dose of porcine epidemic diarrhea virus in a feed matrix. *American Journal of Veterinary Research* 77, 1108–1113.
- Schumacher LL, Cochrane RA, Huss AR, Stark CR, Woodworth JC, Bai JF, Poulsen EL, Chen Q, Main RG, Zhang JQ, Gauger PC, Derscheid RJ, Magstadt DR, Dritz SS and Jones CK** (2017) Characterizing the rapid spread of porcine epidemic diarrhea virus (PEDV) through an animal food manufacturing facility. *PLoS One* 12, e0187309.
- Schumacher LL, Cochrane RA, Huss AR, Gebhardt JT, Woodworth JC, Stark CR, Jones CK, Bai J, Main RG, Chen Q, Zhang J, Gauger PC, DeRouchey JM, Goodband RD, Tokach MD and Dritz SS** (2018) Feed batch sequencing to decrease the risk of porcine epidemic diarrhea virus (PEDV) cross-contamination during feed manufacturing. *Journal of Animal Science* 96, 4562–4570.
- Smith DJ, Oliver CE, Taylor JB and Anderson RC** (2012) Efficacy, metabolism, and toxic responses to chlorate salts in food and laboratory animals. *Journal of Animal Science* 90, 4098–4117.
- Thomson KA, Gebhardt JT, Lerner AB, Woodworth JC, Tokach MD, DeRouchey JM, Goodband BD and Dritz SS** (2018) Evaluation of medium chain fatty acids as a dietary additive in nursery pig diets. *Journal of Animal Science* 96, 252–253.
- Tulayakul P, Boonsoongnern A, Kasemsuwan S, Ratanavanichrojn N, Netvichian R and Khaodiar S** (2012) Heavy metal, *Escherichia coli* and *Salmonella* spp. in feeds, reused water, wastewater, and manure from swine farms: a case report. *Natural Science* 46, 882–893.
- Van Schothorst M and Oosterom J** (1984) Enterobacteriaceae as indicators of good manufacturing practices in rendering plants. *Antonie van Leeuwenhoek* 50, 1–6.
- Williams HE, Cochrane RA, Woodworth JC, DeRouchey JM, Dritz SS, Tokach MD, Jones CK, Fernando SC, Burkey TE, Li YS, Goodband RD and Amachawadi R** (2018) Effects of dietary supplementation of formaldehyde and crystalline amino acids on gut microbial composition of nursery pigs. *Nature Scientific Reports* 8, 8164–8171.
- Yoder AD, Stark CR, DeRouchey JM, Tokach MD and Jones CK** (2018) Effect of cleaning corn on mycotoxin concentration. *Journal of Animal Science* 96, 122–123.
- Zhao T, Podtburg TC, Zhao P, Chen D, Baker DA, Cords B and Doyle MP** (2013) Reduction by competitive bacteria of *Listeria monocytogenes* in biofilms and *Listeria* bacteria in floor drains in a ready-to-eat poultry processing plant. *Journal of Foot Protection* 76, 601–607.