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Meta-analysis of feed intake and growth responses of growing pigs after a sanitary challenge

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Sanitary challenges negatively affect feed intake and growth, leading to a negative impact on animal well-being and economic losses. The aim of this study was to carry out a meta-analysis to quantify the dynamic feed intake and growth responses of growing pigs after a sanitary challenge. A database was constructed using 122 published experiments reporting the average daily feed intake (ADFI) and the average daily gain (ADG) of pigs subjected to one of six sanitary challenges: digestive bacterial infections, poor housing conditions, lipopolysaccharide (LPS) challenges, mycotoxicoses, parasitic infections and respiratory diseases. The responses to experimental challenges were calculated relative to that of a control group. Statistical analyses were carried out for each challenge to quantify the mean and the dynamic responses in feed intake and growth and to identify the basis of the reduction in growth (i.e. reduction in feed intake or reduction in feed efficiency related to changes in maintenance requirements). All challenges resulted in a reduction in ADFI and ADG, with the strongest responses for mycotoxicoses, respiratory diseases and digestive bacterial infections (8% to 23% reduction in ADFI and 16% to 29% reduction in ADG). The reduction in ADG was linearly related to the reduction in ADFI for digestive bacterial infections, LPS challenge, parasitic infections and respiratory diseases. For poor housing conditions and mycotoxicoses, the relationship was curvilinear. A 10% reduction in ADFI resulted in a reduction in ADG varying from 10% for mycotoxicoses to 43% for digestive bacterial infections. More than 70% of the reduction in ADG could be explained by the reduction in ADFI for digestive bacterial infections, LPS challenge, parasitic infections and respiratory diseases. For poor housing conditions and mycotoxicoses, the relationship was curvilinear. A 10% reduction in ADFI resulted in a reduction in ADG varying from 10% for mycotoxicoses to 43% for digestive bacterial infections. More than 70% of the reduction in ADG could be explained by the reduction in ADFI for mycotoxicoses, LPS challenge and respiratory diseases. For challenges associated with the gastrointestinal tract, a large part of the reduction in ADG was due to an increase in maintenance requirements, suggesting digestive and metabolic changes. A dynamic pattern in the reduction in feed intake and growth rate could be identified for digestive bacterial infections, mycotoxicoses and respiratory diseases. For digestive bacterial infections and mycotoxicoses, pigs did not fully recover from the challenge during the experimental period. The results of this study can be used to quantify the effects of a sanitary challenge in growth models of pigs.

Keywords: meta-analysis, pigs, feed intake, growth, sanitary challenge

Implications
Diseases negatively affect feed intake and growth in pigs. This often results in a decrease in feed efficiency, an increase in production costs (e.g. feed and veterinary costs) and an increase in nutrient excretion and in environmental impact. Medication and/or feed additives are often used to limit the negative consequences of diseases. However, the development of sustainable pig production systems requires reducing the use of medication and feed additives. The ability to predict the performance of pigs under a wide range of sanitary challenges allows using management and feeding strategies to minimize the negative consequences.

Introduction
The health status of pigs is often challenged in commercial farms, leading to a lower performance compared with what is potentially possible under good conditions. Several environmental and housing factors are responsible for the reduced performance (Tillon and Madec, 1985). A wide range of pathogenic agents (Kyrizakis and Houdijk, 2007) such as viruses, bacteria, parasites and fungi, as well as social and climatic conditions (Wellock et al., 2003) and the degree of hygiene (Williams et al., 1997) can reduce feed intake, growth and feed efficiency. The deleterious effects on the performance of these factors, defined hereafter as sanitary challenges, are associated with the stimulation of the immune system, which triggers a series of responses of the animal, including a reduction in feed intake and increases in

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energy expenditure, body protein synthesis and catabolism and body temperature (Black, 2009). These responses are mediated through hormones, such as glucagon and corticosterone, and cytokines that regulate nutrient metabolism, immune function and growth (Klasing et al., 1991). Following a sanitary challenge, nutrient partitioning can be altered away from growth (and in particular from tissue protein accretion) toward metabolic responses in support of immune function (Klasing and Johnstone, 1991; Johnson, 1997; Spurlock, 1997). The extent and the duration of the pathophysiological responses depend on the type and the intensity of pathogen exposure and on the host’s ability to stimulate its immune response (Sandberg et al., 2007; Kyriazakis and Doeschl-Wilson, 2009). Different concepts and approaches have been developed to represent the effect of disease on feed intake and growth in pigs (Black et al., 1999; Sandberg et al., 2006; Kyriazakis and Doeschl-Wilson, 2009). However, predicting the effects for each of the major diseases in a production environment requires quantifying how feed intake and growth are affected by different sanitary challenges (Pomar et al., 1991; Lovatto and Sauvant, 2003; van Milgen et al., 2008). This is the objective of this study using a meta-analysis of published results.

Material and methods

Data entry

A database (not available but the list of references used in the meta-analysis is indicated in supplementary materials) was constructed using 122 experiments published between 1968 and 2009 reporting the effects of different sanitary challenges on feed intake and growth in pigs. The selection of candidate publications was based on three criteria: (1) experiments should deal with at least one of the six sanitary challenges studied; (2) experiments should report the average daily feed intake (ADFI) and the average daily gain (ADG); (3) experiments should include a control treatment of pigs that were not challenged. For each experiment, information describing the animals (e.g. sex, genetic origin, number of pigs per treatment, weaning age, BW), experimental conditions (e.g. animal housing, diet composition, duration of the experimental periods), ADFI and ADG was included in the database.

A total of six sanitary challenges were considered (Table 1). Digestive bacterial infections \( (n = 29) \) corresponded to experimental infections of pigs, mainly with *Escherichia coli* (E. coli). Pigs were infected orally or intragastrically. Most of the experiments consisted of a single inoculation at the beginning of the experiment, although multiple inoculations were also used (nine experiments used two to eight inoculations). Inoculation doses varied between \( 10^8 \) and \( 10^{12} \) cfu. These doses were considered (by the authors) to be optimal to induce infection, but the dose effect was not tested. Poor housing conditions \( (n = 20) \) corresponded to pigs raised under suboptimal conditions including poor hygiene conditions \( (n = 13) \), exposure to extreme temperature (heat or cold stress; \( n = 5 \)) and limiting space allowance \( (n = 2) \). Poor hygiene conditions consisted of keeping pigs in pens that were not cleaned or disinfected after a previous occupation by other pigs, resulting in a moderate inflammation model (Williams et al., 1997; Le Floc’h et al., 2006). Lipopolysaccharide (LPS) challenges \( (n = 12) \) corresponded to experimental inflammation of pigs with LPS from *E. coli*. Pigs received intramuscularly approximately 200 \( \mu \)g LPS/kg BW through one or more infections. Mycotoxicoses corresponded to dietary intoxications with a single mycotoxin \( (n = 31) \) or combinations of two or three mycotoxins \( (n = 8) \). The main mycotoxins used were aflatoxin, deoxynivalenol, fumonisin and zearalenon. The toxin contents varied between experiments but also within experiments because half of the experiments tested a dose effect. Parasitic infections \( (n = 12) \) corresponded mainly to single experimental infections with different species of digestive parasites \( (n = 7) \) or with either blood \( (n = 1) \), kidney \( (n = 1) \) or skin parasites \( (n = 3) \). Except for experiments with skin parasites, pigs were infected orally with varying doses within an experiment to test a dose effect. Respiratory diseases \( (n = 10) \) corresponded either to single experimental infections with porcine reproductive and respiratory syndrome virus (PRRSV; \( n = 5 \)), single experimental infections with respiratory bacteria (e.g. *Mycoplasma hyopneumoniae* or *Pasteurella multocida*; \( n = 3 \)) or both \( (n = 2) \). Pigs were infected once intranasally at the beginning of the experiment. Infectious doses were reported only for PRRSV (more than \( 10^8 \) Median Tissue Culture Infective Dose, which was considered to be the optimal infectious dose by the authors).

Management and analysis of data

Each record of the database corresponded to a group of animals of the same age subjected to the same treatment during a given period in an experiment. If an experiment reported data for different periods, the ADFI and ADG across periods were also included in the database. In some experiments, performance was also recorded during successive measurement periods, and this information was also retained.

Table 1 Number of experiments and treatments for each sanitary challenge in the meta-analysis

<table>
<thead>
<tr>
<th>Sanitary challenge</th>
<th>Experiments</th>
<th>Treatments</th>
<th>Control treatments</th>
<th>Dose</th>
<th>Experiment</th>
<th>ADFI</th>
<th>ADG</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>Digestive bacterial infections</td>
<td>29</td>
<td>96</td>
<td>37</td>
<td>12</td>
<td>20</td>
<td>20</td>
<td>48</td>
<td>122</td>
</tr>
<tr>
<td>Poor housing conditions</td>
<td>20</td>
<td>102</td>
<td>48</td>
<td>54</td>
<td>8</td>
<td>102</td>
<td>24</td>
<td>521</td>
</tr>
<tr>
<td>LPS challenge</td>
<td>12</td>
<td>54</td>
<td>24</td>
<td>61</td>
<td>15</td>
<td>54</td>
<td>15</td>
<td>200</td>
</tr>
<tr>
<td>Mycotoxicoses</td>
<td>39</td>
<td>190</td>
<td>61</td>
<td>12</td>
<td>15</td>
<td>190</td>
<td>15</td>
<td>521</td>
</tr>
<tr>
<td>Parasitic infections</td>
<td>12</td>
<td>47</td>
<td>15</td>
<td>10</td>
<td>15</td>
<td>47</td>
<td>15</td>
<td>200</td>
</tr>
<tr>
<td>Respiratory diseases</td>
<td>10</td>
<td>32</td>
<td>15</td>
<td>2</td>
<td>2</td>
<td>32</td>
<td>2</td>
<td>6</td>
</tr>
<tr>
<td>Total</td>
<td>122</td>
<td>521</td>
<td>200</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

LPS = lipopolysaccharide.
The six challenges were analyzed separately and several coding variables were used to identify experiments, experimental factors and repeated measurements within an experiment (Sauvant et al., 2008). The response of the animals to an experimental challenge was calculated relative to that of the control group and expressed as the percentage (i.e. \( \Delta \text{ADFI} \) and \( \Delta \text{ADG} \), for, respectively, the relative change in ADFI and ADG). This allowed accounting for a large part of the variation between experiments. Descriptive statistics were performed on the main characteristics for each challenge. Graphical analyses were carried out throughout the meta-analytical process to detect outliers and to maintain a global view on the heterogeneity of the data, the nature of the relationships between and within experiments and the relationship between the variables to select statistical models (Sauvant et al., 2008).

In all, three analyses were carried out to study the feed intake and growth responses. The first analysis focused on the mean effect of each challenge on \( \Delta \text{ADFI} \) and \( \Delta \text{ADG} \) measured during the overall experimental period, which was tested using a t-test. The second analysis studied the relationship between \( \Delta \text{ADG} \) and \( \Delta \text{ADFI} \) using linear (1) and quadratic regressions (2):

\[
\Delta \text{ADG} = \alpha + \beta \times \Delta \text{ADFI} \quad (1)
\]

\[
\Delta \text{ADG} = \alpha + \beta_1 \times \Delta \text{ADFI} + \beta_2 \times \Delta \text{ADFI}^2 \quad (2)
\]

The intercept (\( \alpha \)) reflects the reduction in ADG not related to the reduction in ADFI, which can be interpreted as an indicator for maintenance. The slope (\( \beta \)) reflects the extent of the change in ADG associated with the reduction in ADFI between challenged and control pigs, and is an indicator of the feed efficiency. Variation between experiments was not explicitly included in the models because the results were expressed relative to the control group. The age of the pig at the beginning of the challenge (i.e. initial age) and the duration of the experimental periods were considered as covariates in the regression analysis. Because the reduction in the growth rate may be age or BW dependent (Coop and Kyriazakis, 1999; Kyriazakis and Houdijk, 2007), and the duration of the experimental periods varied between studies, the magnitude of the response may also vary. Although the pathogen dose may influence the response of an animal, it was not taken into account, because a dose effect was considered only in the experiments with mycotoxins and parasitic infections. In addition, the types of mycotoxin and parasite differed between experiments. The regression parameters and correlations were considered significant when \( P < 0.05 \), whereas \( P < 0.15 \) indicated a trend. The covariates were maintained in the regression analysis when the effect was significant. The normality of residuals was tested using the Shapiro–Wilk test and outliers were identified on the basis of residuals, HI leverage and Cook’s distance (Sauvant et al., 2008). These analyses were carried out using the Minitab® Statistical Software, version 15 (2007).

The time necessary to observe a response or first clinical signs may depend on the pathogen and the host’s ability to cope with a challenge. In a third analysis, the effect of the duration of the overall experimental period on ADFI and ADG was assessed using linear-plateau (LP) (3) and curvilinear-plateau (CLP) (4) regression models:

\[
\Delta \text{ADG} \text{ and } \Delta \text{ADFI} = \left( \text{duration} > T_1 \right) \times L + \left( \text{duration} \leq T_1 \right) \times \left( -a/T_1 \right) \times \left( \text{duration} - T_1 \right) + L \quad (3)
\]

\[
\left( \text{duration} > T_1 \right) \times L + \left( \text{duration} \leq T_1 \right) \times \left( a + (-2 \times a/T_1) \times \text{duration} + (a/T_1^2) \times \text{duration}^2 + L \right) \quad (4)
\]

The parameters \( L, a \) and \( T_1 \) correspond, respectively, to the plateau value, the difference between intercept and plateau value and the time necessary to reach the plateau. The intercept was defined as the immediate reduction in ADFI and ADG following the challenge (i.e. at time = 0). The parameter \( T_1 \) corresponds to the time of recovery (i.e. the time required beyond which no further improvement in \( \Delta \text{ADFI} \) and \( \Delta \text{ADG} \) is observed), whereas parameter \( L \) indicates the extent of the recovery. A zero value for \( L \) indicates that pigs fully recovered from the challenge. The hypothesis that \( L \) differed from zero was tested using an F-test. This analysis was assessed using the PROC NLIN procedure of SAS, version 8.1 (2000).

**Results**

*Experimental data in the database*

Pigs used in the experiments were weaned between 3 and 4 weeks of age (Table 2) and were mainly of mixed sex and of commercial crossbreds ((Large White \times Landrace) \times Pietrain or Duroc). Most experimental challenges were initiated 1 to 2 weeks after weaning, except for parasitic infections that started 6 weeks after weaning. The post-weaning period was the most predominant physiological stage, with an average initial BW of 10 kg and an overall experimental duration of 3 to 5 weeks. Most experiments with parasitic infections and approximately 25% of the experiments with mycotoxins were carried out with growing–finishing pigs and the experimental period ranged from 6 to 11 weeks. The ingredient composition of the diets was rarely reported. Some information about the nutrient composition was indicated in experiments testing poor housing conditions or LPS challenge and the CP content was the most frequently reported nutrient (reported in more than 60% of treatments). According to the authors, the nutrient characteristics met or exceeded the nutrient requirements of the animals.

*Consequences of sanitary challenges on feed intake and growth rate*

All sanitary challenges resulted in a significant decrease in ADFI and ADG compared with the control groups, except for the LPS challenge, where a trend (\( P = 0.14 \)) for a reduction in ADG was observed (Table 2). The most important effects were observed for mycotoxicoses (\( -23\% \text{ ADFI}, -30\% \text{ ADG} \)) and respiratory diseases (\( -16\% \text{ ADFI}, -16\% \text{ ADG} \)). The smallest effects were observed for parasitic infections.
Table 2 Characteristics of experiments used in the meta-analysis to quantify the effect of a sanitary challenge on feed intake and growth in pigs

<table>
<thead>
<tr>
<th>Digestive bacterial infections</th>
<th>Poor housing conditions</th>
<th>LPS challenge</th>
<th>Mycotoxoses</th>
<th>Parasitic infections</th>
<th>Respiratory diseases</th>
</tr>
</thead>
<tbody>
<tr>
<td>96</td>
<td>13 ± 12</td>
<td>98</td>
<td>50 ± 55</td>
<td>54</td>
<td>18 ± 11</td>
</tr>
<tr>
<td>93</td>
<td>29 ± 11</td>
<td>94</td>
<td>26 ± 4</td>
<td>47</td>
<td>21 ± 5</td>
</tr>
<tr>
<td>93</td>
<td>32 ± 11</td>
<td>94</td>
<td>30 ± 8</td>
<td>47</td>
<td>31 ± 7</td>
</tr>
<tr>
<td>78</td>
<td>9.8 ± 6.3</td>
<td>96</td>
<td>11.0 ± 7.5</td>
<td>54</td>
<td>12.9 ± 13.7</td>
</tr>
<tr>
<td>96</td>
<td>17 ± 9</td>
<td>102</td>
<td>31 ± 19</td>
<td>54</td>
<td>21 ± 8</td>
</tr>
</tbody>
</table>

Experimental protocols

| Animal number/Treatment | 96 | 13 ± 12 | 98 | 50 ± 55 | 54 | 18 ± 11 | 190 | 10 ± 9 | 47 | 13 ± 6 | 32 | 30 ± 30 |
| Weaning age (days) | 93 | 29 ± 11 | 94 | 26 ± 4 | 47 | 21 ± 5 | 159 | 29 ± 4 | 31 | 30 ± 3 | 32 | 19 ± 7 |
| Initial age* (days) | 93 | 32 ± 11 | 94 | 30 ± 8 | 47 | 31 ± 7 | 159 | 38 ± 12 | 31 | 71 ± 15 | 29 | 38 ± 15 |
| Initial BW (kg) | 78 | 9.8 ± 6.3 | 96 | 11.0 ± 7.5 | 54 | 12.9 ± 13.7 | 190 | 13.6 ± 10.0 | 45 | 24.8 ± 4.6 | 32 | 13.1 ± 8.7 |
| Duration of experiments (days) | 96 | 17 ± 9 | 102 | 31 ± 19 | 54 | 21 ± 8 | 190 | 42 ± 27 | 47 | 79 ± 13 | 32 | 35 ± 29 |

Responses

| Feed intake reduction (%) | 49 | 8.1 ± 12.1*** | 54 | 3.9 ± 10.3** | 24 | 9.8 ± 13.3* | 92 | 23.1 ± 27.7*** | 32 | 2.9 ± 8.7* | 15 | 16.3 ± 14.6*** |
| Growth rate reduction (%) | 16.5 ± 23.1*** | 9.6 ± 9.6*** | 24.5 | 39.5 | 29.7 | -38.2*** | 8.4 | 11.0*** | 16.2 ± 16.0* |

Difference between feed intake and growth reduction

| LPS | -5.4% ± 1.2% | Poor housing | -7.3% ± 2.7% | LPS | -7.3% ± 2.7% | Mycotoxoses | -7.3% ± 2.7% | Parasitic | -5.4% ± 1.2% | Respiratory | -7.3% ± 2.7% |

Note: LPS = lipopolysaccharide.

*aNumber of treatments used for calculating the mean.

*bAt the beginning of the experimental challenge.

**The response was calculated as the difference between challenged pigs and control pigs and expressed as a percentage of the control.

***P < 0.001; **P < 0.01; *P < 0.05; -P < 0.15; ns = non-significant.
occurred independent of ΔADFI (Figure 2). Although the numerical value of the intercept in parasitic infections was low, it represented almost 70% of ΔADG. For the other three challenges, the intercept explained less than 30% of ΔADG.

Temporal changes in both responses
The LP and the CLP models could be used to describe the ΔADFI and ΔADG as a function of the experimental duration for digestive bacterial infections, mycotoxoses and respiratory diseases, but not for the other three challenges (Figure 3). For all challenges and both responses, the models fitted the data with similar accuracy, although parameter estimates differed (Table 3). The estimates of the immediate reduction in ADFI or ADG (i.e. intercept) were always lower for model LP than for model CLP. The same was true for the time necessary to reach the plateau value. Estimates for the plateau values were slightly higher for model LP than for model CLP. The highest immediate reduction in ADFI and ADG was observed for mycotoxoses (−98% ± 12.1% to −108% ± 14.8% for ΔADFI and −149% ± 13.7% to −176% ± 16.7% for ΔADG in models LP and CLP, respectively). The immediate reduction in ADG was fivefold greater than the reduction in ADFI for digestive bacterial infections. This ratio did not exceed 1.5 for the other two challenges. The time necessary to reach the plateau value was

Figure 1 Relationship between the change in growth (ΔADG) and feed intake (ΔADFI) of pigs challenged with digestive bacterial infections (a), poor housing conditions (b), LPS challenge (c), mycotoxoses (d), parasitic infections (e) and respiratory diseases (f). Responses are expressed as results of the challenged pigs relative to that of a control group. The lines represent the linear or the quadratic model adjustments. Estimated parameters differed from zero (P < 0.05), except for the intercept in LPS challenge.
Discussion

Methodological considerations

The experiments retained in the meta-analysis consisted of experimental models of infection, stress or inflammation, considered as sanitary challenges, where the response of challenged animals could be expressed relative to that of a control group. We also assumed that the sanitary challenges were validated (i.e. potentially sufficient to induce a response). A disadvantage of our choice to include only experiments with challenged and control pigs is that the number of experiments retained was relatively small. Publications reporting data of health disorders in pigs in association with feeding factors but without a specific experimental challenge were not included in the database.

Approximately half of the experiments in the database tested the effect of the challenge in combination with other experimental factors (e.g. in combination with feeding). However, it was difficult to exploit this information further because the experiments tested different nutritional factors. This probably accounted for a large part of the variation between experiments (Kyriazakis, 2010).

Figure 2  Partitioning of the reduction in the average growth rate following a sanitary challenge between the fraction due to the change in maintenance requirement (i.e. not associated with a reduction in feed intake) or due to the change in feed efficiency (i.e. associated with a reduction in feed intake).

Similar for ∆ADFI and ∆ADG for mycotoxoses (30 to 40 days) and for respiratory diseases (30 to 50 days). For respiratory diseases, the plateau value did not differ from zero, indicating that pigs totally recovered. For digestive bacterial infections, growth recovery was incomplete, with an average plateau value of −8% ± 4% during the experimental period, although feed intake recovery was complete (i.e. plateau value not different from zero). The time necessary for total recovery was almost threefold greater for ∆ADFI (38 ± 12 and 60 ± 27 days for models LP and CLP, respectively) than for ∆ADG (14 ± 3 and 21 ± 5 days for models LP and CLP, respectively). For mycotoxoses, recovery was incomplete during the experimental period and ADFI and ADG remained on average 13% lower in challenged pigs compared with control pigs for both models.

Consequences of immune challenges on feed intake

The amplitude and duration of the reduction in ADFI differed considerably between sanitary challenges, and confirm the results reported by others (Kyriazakis et al., 1998; Sandberg et al., 2006). A reduction in appetite and ADFI is frequently observed following exposure to a sanitary challenge, but the mechanisms implied may differ between challenges. The reduction in ADFI observed for sanitary challenges associated with pathogens (bacteria, virus, parasites) or antigens (LPS) probably results from the activation of the innate immune response through the development of an acute-phase response mediated by cytokines (Klasing, 1998a). The poor housing conditions increase microbial pressure, resulting in a continuous activation of the immune system (Klasing et al., 1991). Toxins such as deoxynivalenol act directly on brain transmitters implied in appetite regulation (Etienne, 2007).

For a large range of pathogen types and doses that lead to subclinical diseases in a naïve host, reductions in feed intake of 25% have been reported (Kyriazakis et al., 1998; Sandberg et al., 2006). When pigs develop clinical signs after pathogen exposure, a more significant decrease in feed intake and even complete cessation of eating can occur (Kyriazakis and Houdijk, 2007). The intensity and duration of feed intake reduction may differ between challenges and is related to the type and dose of pathogen (Kyriazakis and Houdijk, 2007), but is also influenced by the ability of the pig to cope with challenges (Sandberg et al., 2007; Kyriazakis and Doeschl-Wilson, 2009).

The immediate reduction in ADFI estimated in this study for pigs challenged with digestive bacterial infections (−15%) is slightly lower than that estimated by Kyriazakis et al. (1998). This difference may be due to the limited number of experiments where the experimental period lasted less than 5 days in our study. A stronger reduction in ADFI (−55%) and even a complete cessation of feeding (−100%) occurred for respiratory diseases and mycotoxoses. Respiratory diseases were mainly caused by PRRSV, which is...
a virulent pathogen (Zimmerman et al., 1997) often leading to anorexia for several days (Done et al., 1996). Mycotoxicoses correspond mainly to diet contamination with aflatoxin or deoxynivalenol, which are known to affect feed intake and can even lead to anorexia for higher doses of mycotoxins (Boudergue et al., 2009).

The time of the recovery period, corresponding to the time between the immediate reduction and the constant reduction in ADFI (i.e. the plateau value), was similar for pigs challenged with mycotoxins and respiratory diseases (approximately 30 days) and longer for digestive bacterial infections (at least 40 days). For respiratory and digestive diseases that are associated with pathogens, the duration of the reduction in ADFI depends on the development of the pathogen and reflects the time required for the acquisition and expression of immunity (Coop and Kyriazakis, 1999). The longer recovery time observed for digestive bacterial infections might be due to the age of challenged pigs, which, in our study, mostly concerned piglets immediately after weaning. Weaning may slow down the maturing of the immune system (Lalles et al., 2007), leading to longer recovery times than in older pigs. Also, digestive bacteria may impair the integrity of the gastrointestinal tract, thereby affecting the digestive capacity of the piglets and thus increasing the time required to recover. In addition, because of abdominal pain caused by digestive infections, the host can develop a feed aversion through a learned association between the sensory properties of the feed and post-ingestive

Figure 3 Relationships between the changes in feed intake (ΔADFI; 1) or growth (ΔADG; 2) and the duration of overall experimental periods in pigs challenged with digestive bacterial infections (A), mycotoxicoses (D) and respiratory diseases (F). Responses are expressed as results of the challenged pigs relative to that of a control group. The lines represent the adjustment with linear-plateau (— LP) and curvilinear-plateau (— - - CLP) models.
Table 3 Parameter estimates obtained from the LP and CLP models describing the changes in feed intake ($\Delta\text{ADFI}$) and growth ($\Delta\text{ADG}$) of pigs challenged with digestive bacterial infections, mycotoxicoses and respiratory diseases relative to the control group as a function of the duration of the overall experimental periods.

<table>
<thead>
<tr>
<th>Digestive bacterial infections ($n = 49$)</th>
<th>Mycotoxicoses ($n = 92$)</th>
<th>Respiratory diseases ($n = 15$)</th>
</tr>
</thead>
<tbody>
<tr>
<td>$\Delta\text{ADFI}$</td>
<td>$\Delta\text{ADG}$</td>
<td>$\Delta\text{ADFI}$</td>
</tr>
<tr>
<td>LP</td>
<td>CLP</td>
<td>LP</td>
</tr>
<tr>
<td>$\delta$</td>
<td>$\delta$</td>
<td>$\delta$</td>
</tr>
<tr>
<td>-15.2 ± 4.2</td>
<td>-15.9 ± 5.2</td>
<td>-41.9 ± 6.2</td>
</tr>
<tr>
<td>$\eta$</td>
<td>$\eta$</td>
<td>$\eta$</td>
</tr>
<tr>
<td>37.9 ± 12.2</td>
<td>59.9 ± 27.1</td>
<td>53.9 ± 15.1</td>
</tr>
<tr>
<td>$L^*$</td>
<td>$L^*$</td>
<td>$L^*$</td>
</tr>
<tr>
<td>0</td>
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**Note:**
- $\Delta\text{ADFI}$: Average daily feed intake; $\Delta\text{ADG}$: Average daily gain; r.s.d.: Residual standard deviation.
- The response was calculated as the difference between challenged pigs and control pigs expressed as a percentage of the control.
- $\delta$: Estimate and standard error of the difference between intercept and plateau value.
- $\eta$: Estimate and standard error of the time (in days) necessary to reach the plateau.
- $L^*$: Estimate and standard error of the plateau value.
- *** $P < 0.001$; ** $P < 0.01$; * $P < 0.05$, indicate a plateau value significantly different from zero.
- Calculated from the sum of $a$ and $L^*$. 

Consequences of sanitary challenges on growth. It has been suggested that the growth after a sanitary challenge is not only due to the reduction in ADFI but also due to an increase in requirements related to the challenged pigs. The reduction in ADFI is proportional to that of ADG for digestive bacterial infections, parasitic infections, LPS and mycotoxicoses. The reduction in ADG was proportional to that of ADFI for digestive bacterial infections, parasitic infections, LPS and mycotoxicoses. The reduction in ADG may result from the repair of damaged tissues, as well as the increased energy and amino acid requirements associated with the stimulation of the immune response.

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immune system (Sandberg et al., 2007). Energy and amino acids are required for the synthesis of cytokines, antibodies, acute-phase proteins and specific immune cells (Klasing, 1998b). The decrease in ADFI combined with the increase in the nutrient requirements for the immune response may lead to significant changes in metabolism and nutrient fluxes in many organs (Klasing, 1998a). In skeletal muscle, protein synthesis is reduced while protein catabolism is stimulated through the direct action of cytokines (Klasing and Johnstone, 1991). This will contribute to the supply of amino acids to the liver to synthesize acute-phase proteins (Klasing et al., 1991), but also lead to a reduction in ADG. Sandberg et al. (2007) suggested that the cost associated with the repair of damaged tissues is greater than that associated with immune response. This may explain why bacterial, viral or parasitic challenges have a greater effect on ADG than the LPS challenge.

In our study, recovery was interpreted as the situation in which the ADG of challenged pigs was no longer different from that of control pigs. According to the concept of homeorhesis, an animal under non-limiting conditions has a growth trajectory related to its genetic potential. After a sanitary challenge, the animal deviates from this trajectory but homeostatic regulations ensure that the animal eventually returns to its growth trajectory (Lovatto and Sauvant, 2003). Total recovery was observed in pigs challenged with respiratory diseases, which supports the idea of compensatory feed intake and compensatory growth (Kyriazakis and Emmans, 1992; Lovatto et al., 2000). However, recovery was incomplete for pigs challenged with mycotoxins, probably resulting from the incomplete recovery in ADFI. Incomplete recovery was also observed with digestive bacterial infections, despite a greater ADFI per kg of BW for challenged pigs. The compensatory feed intake and nutrient supply might be insufficient to cover the important requirements to sustain compensatory growth (Kyriazakis and Emmans, 1992). Another explanation is that total recovery may require a longer time than the duration of the experimental period. Also, it cannot be excluded that the growth potential of the animal (most of which are weaned piglets) has been changed following a sanitary challenge (Escobar et al., 2004).

Conclusion

This meta-analytic approach allowed combining the results of different published experiments on the effect of a sanitary challenge on performance in pigs. For major health disturbances that may occur in commercial pig production, equations were proposed to quantify the reduction in feed intake and growth following a challenge. The reduction in growth is partly due to a reduction in feed intake but also due to changes in digestion and metabolism. The contribution of the latter was most prominent for sanitary challenges that directly affected the gastrointestinal tract. The dynamic pattern in the reductions in feed intake and growth illustrates the responsiveness of the immune system and underlines the necessity to further exploit studies in which the response of animals are recorded during successive measurement periods. The results of this study can also be used to account for the effect of sanitary challenges in predictive growth models of pigs.

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Supplementary material

The supplementary material referred to in this article is available online at http://www.journals.cambridge.org/anm

References


Response of feed intake and growth to a sanitary challenge


