INTRODUCTION

The diets for pigs must contain optimal amounts of phosphorus (P) to promote proper performance and avoid excessive excretion resulting in an increased environmental impact of pig production. The amount of total P (PTOT) in most of the ingredients used in diets for pigs is variable, as is the ratio of PTOT for animals (Rostagno et al., 2011). Availability is a way to express the P content in feedstuffs and is defined as the amount of P that is completely absorbed and utilized by the animal. Another measure used to express the P content in ingredients for pig diets is digestibility, which quantifies the digestive P disappearance.

The digestibility can be expressed as apparent, standardized or true. The apparent digestibility is calculated by the difference between the ingested P (PING) and P
recovered in feces. To obtain standardized and true digestibility values of P, it is necessary to have an estimate of endogenous losses of P, which comes from the salivary juice, gastric, biliary, pancreatic secretion and cell desquamation (Fan et al., 2001). In standardized digestibility, only the endogenous basal losses of phosphorus (EBLP) are considered (NRC, 2012), while in true digestibility, both basal and specific endogenous losses of P are considered.

The standardized digestibility is a good alternative to expressing the P content of feedstuffs for pigs, mainly because fecal endogenous losses of P are relatively easy to measure. Several methods are used to determine the endogenous P, with the most common being the P-free diet and the regression method. In the first case, the idea is to supply a P-free diet for pigs and quantify the P recovered in feces (Petersen and Stein, 2006; Almeida and Stein, 2012). On the other hand, in the regression method diets with increasing levels of P are provided to the animals and the EBLP are determined by the intercept of the relationship between absorbed P (PABS) and PING (Fan et al., 2001). In theory, the use of both P-free diet and the method of regression make it possible to obtain estimates of EBLP (Almeida and Stein, 2010).

The use of gelatin (GEL) as a protein source in P-free diets is recommended by authors who have studied this topic (Petersen and Stein, 2006). In Brazil, diets with low P content have been formulated with spray-dried porcine plasma (SDPP) as a protein source (Bünzen et al., 2012). Despite the low content and high digestibility of P present in the SDPP, its use can generate incorrect estimates of EBLP and, consequently, overestimate the standardized total tract digestibility (STTD) of P in pig feedstuff. One of the advantages of using the SDPP is the lower cost compared to GEL and the SDPP may be used as an alternative protein ingredient in P-free diets. However, no comparison of the endogenous losses with the GEL or SDPP as a protein source has been found in the literature.

Therefore, one study was conducted with the purpose of comparing the EBLP in pigs fed diets containing GEL or SDPP as a source of protein and determine the STTD of P in SDPP.

MATERIALS AND METHODS

Experimental design, animals and diets

The experiment was conducted in the Department of Animal Science of the Federal University of Santa Maria (UFSM); twelve castrated pigs from a commercial line with average weight of 55.0±2.5 kg were used. The animals were housed individually in metabolic cages equipped with feeders, drinkers and containers for the collection of feces and urine. The cages had variable widths and heights in order to adjust the area to the weight of the animal. The cages were kept in a room with a temperature range of 22°C±3°C with the aid of an air conditioner.

The experimental period was divided in two consecutive phases with 12 days (seven days of adaptation to experimental conditions and five days for fecal collection). The animals were distributed through a balanced changeover design, with six replications. The treatments were four semi-purified diets, one of which was free of P but had GEL as a protein source; the others contained 10%, 20%, or 30% of SDPP as the protein source (Table 1). The diets with the addition of SDPP were isocaloric and the ratio of calcium (Ca) and P (Ca:P) in all treatments was 1:1.5; the remainder of the minerals and vitamins were included to meet or exceed the recommendations of Rostagno et al. (2011).

The animals received four meals (8:00, 11:30, 14:00, and 17:30 h), in the amount of 75 g/kg BW0.75 per day. Daily feed refusals were collected, dried, weighed and discounted from the total amount provided.

Table 1. Ingredients and chemical composition as feed basis of semi-purified diets with different spray-dried porcine plasma (SDPP) levels and phosphorus-free diet

| Items | P-free diet | Spray-dried porcine plasma (%) |
|-------|-------------|-------------------------------|
|       | 10          | 20                            | 30                            |
| Ingredient (%) |               |                               |                               |
| Gelatin  | 30.00       | -                             | -                             |
| SDPP    | -           | 10.00                         | 20.00                         | 30.00                         |
| Starch  | 37.25       | 61.15                         | 50.51                         | 39.90                         |
| Sugar   | 20.00       | 20.00                         | 20.00                         | 20.00                         |
| Salt    | 0.40        | -                             | -                             | -                             |
| Soy oil | 6.00        | 3.00                          | 3.00                          | 3.00                          |
| Cellulose| 4.20        | 4.20                          | 4.20                          | 4.20                          |
| Limestone| 0.20        | 0.50                          | 1.00                          | 1.50                          |
| L-threonine| 0.30        | 0.03                          | 0.05                          | 0.05                          |
| DL-methionine| 0.45    | 0.10                          | 0.20                          | 0.30                          |
| L-tryptophan| 0.20    | 0.02                          | 0.04                          | 0.05                          |
| Vitamin/mineral premix |         |                               |                               |
| ME (kcal/kg) | 3,215       | 3,504                         | 3,510                         | 3,516                         |
| Crude protein (%) | 28.20     | 7.20                          | 14.50                         | 21.70                         |
| Dry matter (%) | 90.83       | 89.99                         | 90.34                         | 90.59                         |
| Mineral matter (%) | 0.80        | 1.40                          | 2.48                          | 3.48                          |
| NDF (%) | 3.80        | 3.80                          | 3.80                          | 3.80                          |
| Calcium (%) | 0.09        | 0.21                          | 0.43                          | 0.64                          |
| Total phosphorus (%) | 0.01        | 0.14                          | 0.29                          | 0.43                          |
| Ca:P (%) | -           | 1:1.5                         | 1:1.5                         | 1:1.5                         |

ME, metabolizable energy; NDF, neutral detergent fiber; Ca, calcium.

1 Amount/kg of vitamin and mineral premix: vitamin A 1,750,000 IU; vitamin D3 300,000 IU; vitamin E 10,000 mg; vitamin K3 400 mg; vitamin B1 250 mg; vitamin B2 750 mg; vitamin B6 250 mg; vitamin B12 3,000 mg; niacin 5,000 mg; Pantothenic Acid 3,000 mg; choline, 3,000 mg; antioxidant 5,000 mg; Fe, 8,000 mg as iron sulfate; Cu, 1,200 mg as copper sulfate; Mn, 7,000 mg as manganese sulfate; Zn, 10,000 mg as zinc oxide.

2 Analyzed values.

3 Calculated values according Rostagno et al. (2011), except gelatin (NRC, 1998), and amino acids (Sauvant, 2006).
Data and sample: collections and analysis

The feces were collected according to the marker-to-marker approach using ferric oxide (1%) as a marker. Feces were collected twice daily (Adeola, 2001), placed in plastic bags and kept in a freezer at −18°C. At the end of each period, the feces were thawed and homogenized, and an aliquot was removed for drying at 65°C in a forced air oven, before being milled for chemical analysis. The analyses of dry matter (DM, method 930.15), mineral matter (MM, method 942.05) and P (method 946.06) were performed according to AOAC (2005).

P concentration in feed and fecal samples was determined by gravimetric procedures after wet ashing.

The EBLP derived from diets with inclusion of SDPP was obtained by the regression method with the value of the intercept (Y) of the linear relationship between the PABS and PING, while the STTD of P in SDPP was obtained from the slope of the relationship mentioned previously (Fan et al., 2001). For the method of P-free diet, the entire P recovered in the feces of pigs receiving a P-free diet was considered EBLP (Petersen and Stein, 2006).

Statistical analysis

The data were subjected to analysis of variance, using in the model fixed effects of animal, treatment and period. Means comparison was performed using the Tukey test for contrast between protein sources (GEL×SDPP) and within SDPP diets. Endogenous loss estimates were compared by the Student T test and the relations between PING and PABS were studied by linear regression procedures. The experimental unit was a pig. An alpha level below of 0.05 was considered as statistically significant and values of alpha above 0.05 or below then 0.10 were considered tendency. All statistical analyses were performed using the statistical program MINITAB (2013).

RESULTS AND DISCUSSION

The animals remained healthy during the experimental period and feces samples were obtained without any problem. The GEL used to formulate the P-free diet showed 0.03% PTOT and was supplemented with DL-methionine, L-threonine, and L-tryptophan to improve the quality of its protein (Table 1). The SDPP (AP 920) used in this study contained 1.4% total P.

The dry matter intake (DMI) was different between the treatments (p<0.01; Table 2). The animals eating P-free diets, with GEL as a protein source, had DMI that was 18.5% lower (p<0.01) than animals fed diets with the inclusion of SDPP (1,128.6 vs. 1,382.4 g/d, respectively). It is possible that the lower intake observed in animals fed diets with GEL occurred due to the reduced palatability of this ingredient. However, in other experiments determining endogenous P with GEL as a protein source in a P-free diet, no problem with feed intake was observed (Kim et al., 2012; Rojas and Stein, 2012; Baker et al., 2013).

The comparison between treatments with SDPP showed an intake that was 15% higher (p<0.01) in pigs ingesting diets with 20% of the ingredient tested and equality between the levels of 10% and 30%. At present, no explanations exist for the higher intake of pigs fed diets with intermediate levels.
The apparent total tract digestibility (ATTD) of DM was higher in the diet using GEL (p<0.05) compared with SDPP diets (95.5% and 93.2%, respectively). It is believed that these results were due to differences in the digestibility of dry matter between GEL and SDPP. On the other hand, the comparison between diets containing SDPP indicated no differences (p>0.05) between inclusion levels of SDPP.

There were differences (p<0.01) in the MM intake between GEL and SDPP treatments. These results reflect the amount of MM present in GEL and SDPP (1.45% and 7.74%, respectively). However, the inverse occurred with the ATTD of MM, where the GEL diet provided an ATTD of MM that was around 3 percentage points lower (p<0.01) than the diet with SDPP.

The diets with SDPP provided a P intake that was greater than that of the GEL diet, which was expected because GEL is virtually free of P. In the treatments with SDPP, there was a linear increase in PING with a higher SDPP level in the diet. The linearity in the relationship between the intake and excretion of P is one of the assumptions for validating the conclusions when using the regression method (Fan et al., 2001; Stein et al., 2007).

The EBLP measured in pigs receiving the P-free diet was 128.95 mg/kg DMI; this value is about 8% lower than that found by Petersen and Stein (2006) using diets containing 30% GEL and pigs with an average weight of 53.1 kg. Sulabo and Stein (2013) obtained values that were 21% lower in one study including 20% GEL in the diet tested and using pigs with an average weight of 18 kg. When considering the number of considered studies, it was verified that the endogenous losses from P-free diets containing GEL as a source of protein had an average of 166.3 and standard deviation 35.8 mg/kg DMI (Petersen and Stein, 2006; Almeida and Stein, 2010; Almeida and Stein, 2012; Kim et al., 2012; Rojas and Stein, 2012; Baker et al., 2013; Sulabo and Stein, 2013). Thus, we realized that the estimated values in this study approached from average values reported in the literature.

The results of endogenous losses and standardized digestibility of P in diets with SDPP were calculated by the technique of simple linear regression, using the methodology described by Fan et al. (2001). The relationship between PABS and PING indicated an intercept of 153 mg/kg DMI (standard error [SE] = 77.0, p<0.06) as an estimate of the EBLP in feces (Figure 1). For the analysis of data from the literature, it can be seen that EBLP estimated by the regression method are quite variable. In experiments with soybean meal as a source of P, for example, average values of 505 mg/kg DMI from EBLP have been found with a standard deviation of 270 mg/kg DMI (Fan et al., 2001; Ajakaiye et al., 2003; Zou et al., 2007a,b; Zhang et al., 2008; Akinmusire and Adeola, 2009). Differences in experimental conditions such as basal diet, housing, and animal genetics can determine these differences. Obviously, the good estimates of EBLP are important for feed evaluation, such as the maintenance requirements of P (Yang et al., 2007).

The fecal content of P is due to unabsorbed P originating in the diet consumed by the animal and also the endogenous P. Endogenous losses, in turn, can be decomposed into basal or non-specific, and extra or specific, with the EBLP being attributed exclusively to the dry matter intake, and the latter being related to the composition of the test food (Mosenthin et al., 2000). In theory, it is expected that the regression method results estimate the EBLP similar to that obtained with a P-free diet, because the composition of the basal diet was similar for both methods.

The aim of this study was to determine whether the EBLP with the method of regression, using the SDPP as a protein source, would be compatible with the basal losses estimated using P-free diets. The comparison between the endogenous losses shows that there was no difference (p>0.05) when it was obtained by the P-free method, with GEL as a protein source, or by the regression method. We emphasize, however,
that the P-free diet induced a loss that was approximately 18% lower than the regression method (Figure 2).

Therefore, it is suggested that the SDPP can serve as a source of amino acids in diets designed to estimate the EBLP. However, it must be considered that the digestibility of P in SDPP is high, but not 100%, which leads to the need to consider the indigestible fraction in the calculations of endogenous losses.

Pigs with a body weight of 60 kg should ingest 6.4 g of digestible P (PDIG) a day (Rostagno et al., 2011). Considering a daily consumption of 2.3 kg DM and associating that consumption with our EBLP estimates, this implies that EBLP lost in the feces represent 7% of PDIG required by pigs (Rostagno et al., 2011). This value is 25% higher than that reported by Shen et al. (2002), which may be justified because the authors have obtained EBLP that are 4 times higher than the average value obtained in this experiment.

The comparison between the apparent digestibility values of P, with the standardized digestibility values of P (97.4%) obtained from the simple regression technique, is shown in Figure 3. The apparent digestibility of P was 87.9%, 94.2%, and 92.9% for treatments with 10%, 20%, and 30% of SDPP inclusion, respectively. Apparent digestibility values increase in response to the higher consumption of P because the endogenous losses, when expressed as a percentage of total P recovered in the feces, decrease proportionately (Mosenthin et al., 2000; Stein et al., 2007). The estimated standardized digestibility of P (97.4%) generated by simple linear regression technique corrected the values for endogenous basal flux of P, which remain unchanged, independent of the P inclusion level (Yang et al., 2007).

The standardized digestibility value of P in the SDPP found in our experiment was 5 percentage points higher than that cited by Rostagno et al. (2011), but close to the value of

Figure 2. Estimated values of endogenous phosphorus losses mg/kg dry matter intake (DMI) from phosphorus free diet or the regression method with six observations per treatment.

Figure 3. Effect of ingested phosphorus mg/kg dry matter intake (DMI) in standardized and apparent digestibility with spray-dried porcine plasma with six observations per treatment.
98% of standardized digestibility presented in the NRC (2012). This result demonstrates that the P in SDPP has high digestibility that is associated with low content, meaning that it can be recommended as a substitute for GEL in P-free diet.

The apparent digestibility values of P obtained at different levels of inclusion of SDPP were corrected according to the equation suggested by Mosenthin et al. (2000): STTD of P in SDPP(%) = ATTD(%) + [(EBLP mg/kg DMI×P content in the diet mg/kg DM)×100] by the basal diet loss obtained by the method of P-free diet, with GEL as a protein source; here, the values of 96.9%, 98.7%, and 95.9% (data are not shown) were found for the standard P protein source; here, the values of 96.9%, 98.7%, and 95.9% (data are not shown) were found for the standard P protein source.

Stein et al. (2007) recommend that the basal amino acid flux be measured routinely in studies aiming to evaluate ileal digestibility. The suggestion comes from the observation that even with standardized experimental conditions and analytical procedures, discrepancies are detected. This same recommendation can be extended to studies whose purpose is to determine the EBLP.

CONCLUSION

SDPP can replace GEL as a protein source in P-free diets designed to determine endogenous losses of P. The EBLP showed an inevitable minimal loss of P for pigs of 141.3 mg/kg DM intake. The STTD of P in the SDPP was 97.2%, as estimated by P-free diet and 97.4% using SDPP as a protein source.

CONFLICT OF INTEREST

We certify that there is no conflict of interest with any financial organization regarding the material discussed in the manuscript.

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