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## Associative effects between two fibre sources on ileal and overall digestibilities of amino acids, energy and cell-wall components in growing pigs

BY J. P. LAPLACE AND BEATRICE DARCY-VRILLON

*Laboratoire de Physiologie de la Nutrition, INRA-Centre de Recherches de Jouy, F-78350 Jouy-en-Josas, France*

AND J. M. PÉREZ AND Y. HENRY

*Station de Recherches Porcines, INRA-St Gilles, F-35590 L'Hermitage, France*

AND SYLVIE GIGER AND D. SAUVANT

*Station de Nutrition et Alimentation, INA-Paris-Grignon, 16 rue Claude Bernard, F-75231 Paris Cedex 05, France*

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1. The associative effects of two fibre sources on ileal and overall digestibility of amino acids, energy and cell-wall components were studied by comparing wheat bran and soya-bean hulls in semi-purified diets given to growing pigs.

2. Castrated male pigs were prepared with ileo-rectal anastomosis to measure ileal digestibility, and overall digestibility was measured in pigs without anastomosis.

3. The three diets contained 190 g total fibre/kg dry matter (DM), derived from each fibre source or from a mixture of both fibre sources, so that each source provided half the amount of total fibre, and 170 g crude protein (nitrogen  $\times$  6.25)/kg DM by additions of casein.

4. The effects of fibre sources on the ileal digestibility of amino acids were additive for most amino acids; the only significant interactions were found for threonine, methionine and aspartic acid. In contrast with ileal digestibility, systematic negative interactions between fibre sources on overall digestibility of amino acids were noted.

5. There was no interaction between fibre sources in their effects on the digestibility of energy or of cell-wall components, irrespective of the site of digestion. The digestibility values were higher with soya-bean hulls than with wheat bran, especially at the faecal level.

6. It is concluded that ileal digestibility of amino acids provides a better estimate of amino acid availability, as ileal measurements allow a better discrimination between diets than faecal measurements when distinct fibre sources are used alone or in combination at the same total fibre content. In contrast, for energy, the measurement of digestibility at both faecal and ileal levels permits the estimation of the partition of available nutrients between the small and large intestines.

Dietary fibre in the diets of pigs has been given special attention during recent years (Low, 1985), in particular with regard to its effects on several physiological variables such as gastric emptying (Rainbird, 1986; Rainbird & Low, 1986*a, b*), food passage (Sambrook, 1979; Fioramonti & Bueno, 1980; Kuan *et al.* 1983; Bardon & Fioramonti, 1983), fermentation in the large intestine (Argenzio & Southworth, 1975; Imoto & Namioka, 1978; Ehle *et al.* 1982; Varel *et al.* 1984; Varel, 1987) and cholesterol and bile acid metabolism (Collings *et al.* 1979). In addition, information has accumulated on the digestion of fibre constituents in the different segments of the gastrointestinal tract (for example, Kass *et al.* 1980*a, b*).

The overall effect of dietary fibre on protein digestion and amino acid availability has been studied (Murray *et al.* 1977; Dierick *et al.* 1983; Drochner, 1984). However, there has been little documentation of this effect in relation to the type of fibre used and its chemical composition (Stanogias & Pearce, 1985*a-c*; Graham *et al.* 1986), and the possible

interactive effects of different fibre sources on amino acid availability have not been evaluated.

Therefore, the present work was designed to study the ileal and overall digestibilities of amino acids and fibrous materials in growing pigs fed on diets containing well-defined fibre sources (wheat bran, soya-bean hulls), either alone or in combination.

## MATERIALS AND METHODS

### *Diets*

Three semi-purified diets based on casein and maize starch were prepared to contain similar amounts of total fibre, originating from either wheat bran (diet B) or soya-bean hulls (diet H), or from a mixture of both fibre sources (diet BH) so that each source provided half the amount of total fibre. Total fibre, which amounted to about 190 g/kg dry matter (DM), was expressed as the sum of analysed cellulose + hemicellulose + uronic acids + acid detergent lignin, according to the methods described on p. 78. In addition to total fibre, the diets contained equivalent amounts of crude protein (nitrogen  $\times 6.25$ ; 170 g/kg DM), total fat and gross energy. The chemical composition of the dietary fibre sources is given in Table 1. The composition of the experimental diets and their chemical characteristics are shown in Table 2. The contents of amino acids in the three diets are given in Table 3.

Two separate trials were conducted with Large White castrated male growing pigs for overall and ileal digestibility measurements. Both overall and ileal digestibilities are expressed in the conventional way, i.e. (intake - excretion)/intake. Excretion was determined on the basis of total collection of either faeces or ileal digesta.

### *Ileal digestibility*

The mean initial live weight of the five pigs was 55.4 (SE 3.9) kg at surgery. Under halothane anaesthesia (Fluothan<sup>®</sup>, Coopers Vétérinaire S.A., Meaux, France) a mid-line incision from the navel to the pubis gave access to the distal digestive tract, to form an end-to-end ileo-rectal anastomosis after total bypass of the entire caeco-colic area as a Thiry loop. The ileum was cut 50 mm above the ileo-caeco-colic junction and the distal section was closed by a double invagination. The rectum was cut at the level of the pubic bone. A plastic cannula (PVC) was inserted into the proximal section of the caeco-colic area, and fixed by a purse-string suture. The cannula was then exteriorized through a small incision in the right flank of the animal to allow the extracorporeal emptying of residual digesta and gases from the caecum and colon. These segments were thus completely disconnected from the functional gastrointestinal tract. The proximal section of the ileum and the distal section of the rectum were anastomosed end-to-end by individual stitches. However, in most cases, the ileal part had to be cut obliquely in order to compensate for the difference in ileal and rectal diameters. Antibiotics were applied to the peritoneal cavity and two additional intramuscular injections were performed at 48-h intervals.

This surgical preparation allowed collection of ileal digesta directly excreted via the anus. Under the same general anaesthesia, the pigs' tails were amputated to avoid loss of digesta due to spreading or projection by tail movement. The pigs were adapted to individual housing in metabolism cages. In addition to the usual complete surgical and post-surgical care, specific additional attention included: (1) continuous supplementation of diets by sodium chloride (5 g/kg diet), sodium bicarbonate (5 g/kg diet) and a vitamin mix (1 g/kg diet) (previous experiments confirmed the adequacy of such a supplementation to compensate for the loss of colic function (Laplace *et al.* 1985 *b*)); (2) external application of a cream used for human babies (Mitosyl<sup>®</sup>) to avoid skin irritation due to the increased

Table 1. Chemical composition of dietary fibre sources (g/kg dry matter (DM))

	Dietary fibre source	
	Wheat bran	Soya-bean hulls
DM (g/kg diet)	871	881
Organic matter	929	952
Crude protein (nitrogen $\times$ 6.25)	150	120
Diethyl ether extract	39	20
Gross energy (MJ/kg DM)	19.46	18.33
Crude fibre	123	385
Neutral-detergent fibre	454	641
Acid-detergent fibre	141	450
Acid-detergent lignin	35	20
Hemicellulose*	257	210
Cellulose*	101	306
Uronic acids†	—	61
Total fibre‡	393	597
Individual monosaccharides in hemicellulose* and cellulose* fractions		
Rhamnose	1	5
Arabinose	100	50
Xylose	142	77
Mannose	4	53
Galactose	10	25
Glucose	101	306

\* Determined by gas-liquid chromatography according to Sawardeker *et al.* (1965).

† Determined according to El Rayah & Labavitch (1977).

‡ Cellulose + hemicellulose + uronic acids + acid-detergent lignin.

emission of digesta which were more liquid and acid than normal faeces. The colic cannula was kept clean several times daily until all residual colic contents were emptied. During the 8 d post-operative period, the pigs were fed on a standard diet containing the following amounts of ingredients (g/kg diet): barley 600, maize 150, soya-bean meal 150, lucerne (*Medicago sativa*) meal 65, minerals and vitamins 35. The crude protein and crude fibre contents (g/kg DM) were 159 and 67 respectively. The experimental period lasted 3 weeks, each pig being given the three diets consecutively in the following order: B, H, BH, with 4 d of adaptation and 3 d of collection for each diet. During the three experimental weeks, the pigs received 2 kg fresh food daily as two equal meals at 09.00 and 16.00 hours. The feed was mixed with water (1 part meal + 2 parts water). Water was given *ad lib*.

The excreted digesta were collected twice daily. After homogenization of the digesta collected over 24 h, a sample (10% of the total wet weight) was taken for determination of DM, nitrogen and gross energy. In addition, a representative sample of the 72 h collection was taken for each pig on each diet to determine amino acid and fibre contents.

#### Overall digestibility

Overall digestibility measurements were performed with eighteen pigs which were allocated to the three treatments (six pigs/diet) according to a completely randomized block design, taking age and live weight into account. After a 14 d adaptation period in digestibility crates, a total faecal collection was made during a 10 d period after the pigs reached a mean initial live weight of 47.9 (SE 3.0) kg. The level of feeding was adjusted to the initial live weight and was maintained constant during the entire collection period. The average daily

Table 2. *Ingredients (g/kg diet) in the experimental diets and their chemical composition (g/kg dry matter (DM))*

	Diet		
	B	BH	H
<b>Ingredients</b>			
Wheat bran	484.5	242.2	—
Soya-bean hulls	—	166.0	332.0
Casein (hydrochloride)	103.0	119.7	136.3
Maize starch	357.5	412.1	465.7
Maize oil	20.0	25.0	31.0
Mineral and vitamin mixture*	35.0	35.0	35.0
<b>Chemical composition</b>			
DM	884	883	888
Organic matter	934	945	956
Crude fibre	58	98	127
Neutral-detergent fibre	228	229	217
Acid-detergent fibre	71	112	148
Acid-detergent lignin	21	15	8
Total fibre†	190	194	198
Crude protein (nitrogen × 6.25)	171	169	171
Diethyl ether extract	44	43	41
Gross energy (MJ/kg DM)	19.04	19.20	19.12

\* Supplying the following quantities (/kg diet): dicalcium phosphate 10 g, calcium carbonate 4.5 g, sodium chloride 3.0 g, potassium chloride 4.0 g, magnesium carbonate 2.0 g, ferrous sulphate heptahydrate 50 mg, manganese sulphate pentahydrate 16 mg, copper sulphate pentahydrate 4 mg, zinc sulphate heptahydrate 44 mg, potassium iodide 45 µg, sodium selenite 60 µg, retinol 1.5 mg, cholecalciferol 25 µg,  $\alpha$ -tocopherol 28 mg, menadione 4.4 mg, thiamin hydrochloride 3.0 mg, riboflavin 6.0 mg, nicotinic acid 36 mg, calcium pantothenate 30 mg, pyridoxine hydrochloride 3.0 mg, pteroylmonoglutamic acid 2.0 mg, *meso*-inositol 200 mg, *p*-aminobenzoic acid 20 mg, biotin 3.0 µg, cyanocobalamin 30 µg, choline chloride 1.5 g.

† Calculated from the chemical analyses of dietary fibre sources: cellulose + hemicellulose + uronic acids + acid-detergent lignin.

feed supply was 2 kg (1.76 kg DM). The diets were offered twice daily in wet form (2 parts water + 1 part meal) as in the previous trial. Faeces were collected twice daily and stored at +1°.

#### *Chemical analyses*

The determinations of DM, ash and N (Kjeldahl) in feeds were made according to conventional methods. Gross energy content was measured with an adiabatic bomb calorimeter. Fat content was analysed with a Soxhlet-type semi-automatic apparatus (Soxtec). Amino acid analyses were made by ion-exchange chromatography as outlined by Pion & Fauconneau (1966). The contents of cell-wall components were determined with the following methods: crude fibre (EEC official method), neutral-detergent fibre (NDF), acid-detergent fibre (ADF) and acid-detergent lignin (ADL), according to the procedure described by Van Soest (1963), Van Soest & Wine (1967), and adapted by Giger *et al.* (1987*b*). Individual cell-wall monosaccharides liberated by acid-hydrolysis (Saeman *et al.* 1954) were analysed by gas-liquid chromatography of their alditol acetates (Sawardeker *et al.* 1965). Cell-wall uronide content was determined in fibrous sources according to the method of El Rayah & Labavitch (1977).

#### *Statistical analysis*

The methodology used to test digestive interactions has been fully described elsewhere by Giger *et al.* (1987*a*). Briefly, the method involved two steps. First, the 'diet' effect was

Table 3. Amino acid composition of experimental diets\* (g/kg dry matter)

	Dietary fibre source		
	Wheat bran	Wheat bran and soya-bean hulls†	Soya-bean hulls
Sum of seventeen amino acids	175.0	177.0	185.0
Essential amino acids			
Lysine	10.8	11.6	13.0
Methionine	3.3	3.8	3.8
Cystine	2.4	1.9	1.6
Threonine	7.5	7.2	7.9
Isoleucine	8.0	8.1	8.8
Leucine	14.3	14.4	15.3
Histidine	5.4	5.2	5.4
Valine	11.2	11.0	11.4
Phenylalanine	8.3	8.2	8.9
Tyrosine	8.3	8.8	9.9
Arginine	9.7	8.3	7.7
Non-essential amino acids			
Glutamic acid + glutamine	35.7	33.9	34.3
Aspartic acid + asparagine	12.2	12.1	11.6
Serine	8.9	9.2	10.0
Proline	16.5	15.5	17.1
Glycine	6.1	5.9	6.2
Alanine	6.7	6.1	6.1

\* For details, see Tables 1 and 2.

† Each fibre source provided half the amount of total fibre.

tested by an analysis of variance also taking into account any 'animal' effect. For repeated measurements, it was also possible to test the 'animal' × 'diet' interaction. If there was no 'diet' effect, there was no digestive interaction. If a significant 'diet' effect was found, it was necessary to use a regression method as a second step, as described by Giger & Sauvant (1983). The residual sums of squares obtained from the variance analysis and from the regression were compared by using the Fisher test (Snedecor & Cochran, 1967). By demonstrating a non-linear regression, this procedure allowed the existence of any digestive interaction to be shown.

## RESULTS

### *Protein and amino acid digestibility*

There were no significant differences between animals in the ileal digestibility measurements for amino acids and total N. As appears from the results given in Table 4, the digestibility of total N and of the sum of amino acids was not significantly affected ( $P > 0.05$ ) by diet composition. Diet effect, depending on the origin of cell-wall components, was significant for ten of the seventeen amino acids. For only three of them was there a significant interaction between fibre sources, which was positive for methionine and aspartic acid and negative for threonine. In addition to these three amino acids, digestibility values for cystine, histidine tyrosine, serine, glycine and aspartic acid were higher with diet B (wheat bran) than with diet H (soya-bean hulls). Conversely, amino acid digestibility was lower with diet B for leucine, phenylalanine, methionine and threonine.

The results of overall digestibility are reported in Table 5. There was only a small difference between diets B and H. The statistical importance of the overall diet effect is essentially explained by the association between the two dietary fibre sources (wheat bran

Table 4. *Ileal apparent digestibility of crude protein (nitrogen  $\times$  6.25) and amino acids of experimental diets†*  
(Mean values and standard deviations)

	Dietary fibre source						Statistical significance of	
	Wheat bran		Wheat bran and soya-bean hulls‡		Soya-bean hulls		Treatment effect	Non-linear effect
	Mean	SD	Mean	SD	Mean	SD		
Crude protein	0.808	0.011	0.807	0.008	0.800	0.003	NS	
Sum of seventeen amino acids	0.850	0.009	0.850	0.008	0.843	0.006	NS	
Essential amino acids								
Lysine	0.866	0.011	0.858	0.006	0.856	0.004	NS	
Methionine	0.887	0.004	0.914	0.004	0.906	0.009	***	**
Cystine	0.682	0.017	0.627	0.019	0.536	0.019	***	NS
Threonine	0.800	0.016	0.798	0.014	0.827	0.004	**	*
Isoleucine	0.844	0.008	0.844	0.010	0.832	0.014	NS	
Leucine	0.863	0.007	0.874	0.006	0.875	0.007	*	NS
Histidine	0.871	0.006	0.860	0.009	0.861	0.005	*	NS
Valine	0.839	0.007	0.841	0.009	0.836	0.009	NS	
Phenylalanine	0.864	0.005	0.872	0.008	0.881	0.008	**	NS
Tyrosine	0.885	0.007	0.875	0.004	0.866	0.010	**	NS
Arginine	0.856	0.013	0.850	0.009	0.850	0.008	NS	
Non-essential amino acids								
Glutamic acid + glutamine	0.888	0.016	0.889	0.009	0.887	0.008	NS	
Aspartic acid + asparagine	0.817	0.009	0.812	0.008	0.779	0.007	***	**
Serine	0.831	0.013	0.800	0.017	0.761	0.022	**	NS
Proline	0.903	0.013	0.896	0.011	0.890	0.006	NS	
Glycine	0.670	0.026	0.587	0.017	0.542	0.011	***	NS
Alanine	0.742	0.011	0.741	0.012	0.739	0.007	NS	

NS, not significant.

\*  $P < 0.05$ , \*\*  $P < 0.01$ , \*\*\*  $P < 0.005$ .

† For details, see Tables 1–3.

‡ Each fibre source provided half the amount of total fibre.

and soya-bean hulls) which resulted in a systematic negative interaction ( $-3$  points on average) for all the amino acids, especially in the case of threonine ( $-5$  points). The same phenomenon was observed for crude protein and for the sum of the seventeen amino acids.

#### *Digestibility of energy and cell-wall components*

As shown in Table 6, for all variables there was no animal effect on ileal digestibility and no block effect on overall digestibility. In addition, no interaction was observed. The diet effect was significant, with uniformly higher digestibility values in diet H than in diet B. This difference between the two diets was particularly important at the faecal level.

The calculation of the digestibility of the dietary non-nitrogenous cellular fraction indicates a slightly but significantly lower ( $P < 0.01$ ) digestibility at the terminal ileum with soya-bean hulls (diet H), while the opposite effect, moderate but significant ( $P < 0.05$ ), was observed for overall digestibility. On the whole, this corresponded to a significantly higher ileal digestibility of organic matter and energy in diet H containing soya-bean hulls.

Table 5. Overall apparent digestibility of crude protein (nitrogen  $\times$  6.25) and amino acids of experimental diets†  
(Mean values and standard deviations)

	Dietary fibre source						Statistical significance of	
	Wheat bran		Wheat bran and soya-bean hulls‡		Soya-bean hulls		Treatment effect	Non-linear effect
	Mean	SD	Mean	SD	Mean	SD		
Crude protein	0.872	0.015	0.835	0.019	0.864	0.006	**	**
Sum of seventeen amino acids	0.895	0.015	0.867	0.014	0.890	0.006	**	**
Essential amino acids								
Lysine	0.884	0.017	0.858	0.022	0.887	0.011	*	**
Methionine	0.867	0.019	0.843	0.015	0.883	0.009	***	***
Cystine	0.806	0.023	0.743	0.024	0.746	0.009	***	***
Threonine	0.855	0.022	0.804	0.020	0.858	0.005	***	***
Isoleucine	0.874	0.023	0.845	0.018	0.877	0.009	*	***
Leucine	0.894	0.013	0.870	0.013	0.902	0.006	***	***
Histidine	0.921	0.010	0.896	0.016	0.907	0.007	**	**
Valine	0.885	0.017	0.860	0.018	0.888	0.009	*	***
Phenylalanine	0.889	0.015	0.853	0.014	0.886	0.006	***	***
Tyrosine	0.905	0.013	0.877	0.012	0.898	0.008	***	***
Arginine	0.897	0.015	0.862	0.015	0.887	0.010	***	***
Non-essential amino acids								
Glutamic acid + glutamine	0.941	0.009	0.918	0.010	0.932	0.004	***	***
Aspartic acid + asparagine	0.840	0.023	0.795	0.022	0.821	0.011	***	***
Serine	0.898	0.014	0.864	0.012	0.882	0.004	***	***
Proline	0.952	0.006	0.938	0.006	0.946	0.003	***	***
Glycine	0.806	0.024	0.693	0.025	0.699	0.011	***	***
Alanine	0.805	0.017	0.740	0.030	0.778	0.010	***	***

NS, not significant.

\*  $P < 0.05$ , \*\*  $P < 0.01$ , \*\*\*  $P < 0.005$ .

† For details, see Tables 1–3.

‡ Each fibre source provided half the amount of total fibre.

#### Ileal v. overall digestibility

Except for one amino acid (phenylalanine) (see Table 7) the digestibility coefficients of all dietary components (energy and N as well) were significantly affected by the site of digesta collection (ileum v. faeces). For all these components, there was a significant interaction between treatment and collection site, except for three amino acids (lysine, histidine, proline).

#### DISCUSSION

With regard to the methodological approach, two points need to be considered: the choice of ileo-rectal anastomosis and the design of experimental diets.

The ileo-rectal anastomosis has been used in the past few years concurrently with various techniques based on ileal fistulation: i.e. 'T'-shaped, re-entrant or ileo-colic-post-valvular fistulation (Laplace *et al.* 1985*b*). This procedure became widely used relatively quickly, since the surgical preparation of an ileo-rectal anastomosis and the collection of digesta are very easy compared with fistulation techniques. In addition, the ileo-rectal anastomosis is well tolerated for long-term periods. Despite these advantages, the possible consequences



Table 6. *Ileal and overall digestibilities of non-nitrogenous components of experimental diets†*

(Mean values and standard deviations)

	Dietary fibre source						Statistical significance of	
	Wheat bran		Wheat bran and soya-bean hulls‡		Soya-bean hulls		Treatment effect	Non-linear effect
	Mean	SD	Mean	SD	Mean	SD		
<i>Ileal digestibility (five pigs)</i>								
Dry matter	0.647	0.022	0.658	0.014	0.681	0.027	**	NS
Organic matter (OM)	0.678	0.012	0.681	0.008	0.703	0.007	**	NS
Gross energy	0.675	0.021	0.691	0.014	0.713	0.025	**	NS
Crude fibre	-0.014	0.027	0.050	0.031	0.044	0.022	**	NS
Neutral-detergent fibre (NDF)	0.060	0.033	0.093	0.023	0.152	0.025	**	NS
Acid-detergent fibre	0.051	0.033	0.033	0.030	0.039	0.023	NS	
OM-Sum AA-NDF§	0.887	0.007	0.875	0.007	0.871	0.006	**	NS
<i>Overall digestibility (six pigs)</i>								
Dry matter	0.778	0.007	0.821	0.023	0.877	0.013	***	NS
Organic matter (OM)	0.800	0.007	0.837	0.021	0.888	0.013	**	NS
Gross energy	0.785	0.008	0.825	0.022	0.878	0.013	**	NS
Crude fibre	0.235	0.026	0.543	0.134	0.688	0.074	***	NS
Neutral-detergent fibre (NDF)	0.393	0.013	0.564	0.084	0.731	0.048	***	NS
Acid-detergent fibre	0.213	0.027	0.518	0.139	0.690	0.068	***	NS
OM-Sum AA-NDF§	0.940	0.006	0.942	0.003	0.948	0.004	*	NS

NS, not significant; AA, amino acids.

\*  $P < 0.05$ , \*\*  $P < 0.01$ , \*\*\*  $P < 0.005$ .

† For details, see Tables 1-3.

‡ Each fibre source provided half the amount of total fibre.

§ Non-nitrogenous cellular content; Sum AA = sum of seventeen amino acids.

of the ileo-rectal anastomosis have not been fully evaluated, as pointed out by Laplace *et al.* (1985*b*): hydro-electrolytic disturbances and vitamin deficiencies due to the bypass of the colon, reduced energy supplies due to the absence of colic fermentation, possible adaptations or changes in digestive and absorptive functions in long-term experiments, as well as possible hepatic dysfunctions or metabolic disorders which are well known to occur after small intestine resection or bypass (Laplace, 1975). In the present experiments, the problems mentioned previously have been treated in the following manner: the animals were given supplementary electrolytes and vitamins; the measured weight gain appeared within normal limits despite the absence of colic fermentation; the entire experiment was completed within 4 weeks following the operation, thus avoiding any long-term digestive or metabolic changes. Consequently, the period of adaptation to experimental diets was reduced to 4 d, which may seem brief. However, it should be underlined that bypassing the colon takes out the part of the intestinal tract which needs the longest time to clear residues of previous diets. In pigs after ileo-rectal anastomosis, 4 d are sufficient for renewing the contents of the stomach and small intestine (Laplace, 1981). Finally, the ileo-rectal anastomosis provides a particular advantage in the case of high-fibre diets, by avoiding possible blockage of cannulas and consequent erroneous estimates of food passage and digestibility.

The diets were designed to contain the same amounts of total fibre, crude protein and gross energy. Since the two fibre sources differed in their crude protein contents, this resulted in some difference in the level of casein incorporation, so that the proportion of

Table 7. Effects of the site of digesta collection and of its interaction with the treatment (diets) on the digestibility of dietary components

	Statistical significance of effects			Pooled SE
	Site of digesta collection	Treatment	Interaction site × treatment	
Crude protein (nitrogen × 6.25)	***	**	**	0.012
Sum of seventeen amino acids	***	*	*	0.010
Essential amino acids				
Lysine	***	*	NS	0.014
Methionine	***	**	***	0.012
Cystine	***	***	***	0.020
Threonine	***	***	**	0.015
Isoleucine	***	NS	*	0.015
Leucine	***	***	***	0.010
Histidine	***	***	NS	0.010
Valine	***	NS	*	0.013
Phenylalanine	NS	***	***	0.010
Tyrosine	***	***	**	0.010
Arginine	***	***	*	0.012
Non-essential amino acids				
Glutamic acid + glutamine	***	NS	*	0.010
Aspartic acid + asparagine	*	***	***	0.015
Serine	***	***	***	0.014
Proline	***	*	NS	0.008
Glycine	***	***	*	0.020
Alanine	***	**	**	0.020
Non-nitrogenous components				
Dry matter	***	**	**	0.013
Organic matter (OM)	***	**	**	0.014
Gross energy	***	***	**	0.037
Crude fibre	***	**	**	0.069
Neutral-detergent fibre (NDF)	**	**	**	0.046
Acid-detergent fibre	***	***	***	0.070
OM-Sum AA-NDF†	***	NS	**	0.006

NS, not significant.

\*  $P < 0.05$ , \*\*  $P < 0.01$ , \*\*\*  $P < 0.005$ .

† Non-nitrogenous cellular content; Sum AA = sum of seventeen amino acids.

dietary protein from casein in total protein was 0.57, 0.68 and 0.75 in diets B, BH and H respectively. In diet H the lower digestibility of protein in soya-bean hulls compared with wheat bran (Institut National de la Recherche Agronomique, 1984) was compensated by a higher dietary content of highly-digestible protein in the form of casein. Therefore, there was little difference in protein and amino acid digestibility between diets B and H at ileal and faecal levels. Nevertheless, since the experiment was focused on the interactions between fibre sources, it was necessary to formulate the diets on the basis of the same overall fibre content. Notwithstanding, this shows the difficulty of interpreting the effects of cell-wall polysaccharides on amino acid digestibility with fibrous feeds containing different proportions of protein. On the other hand, the equilibration of dietary energy contents resulted in a slight increase in the levels of maize starch and oil with the increasing level of soya-bean hulls. This provides a partial explanation of the higher ileal digestibility of energy in diet H.

From previous results using ileal-fistulated pigs, there is clear evidence that amino acid ileal digestibility allows a better discrimination between diets than overall digestibility, as reported by Low (1982), Tanksley & Knabe (1984) and Laplace *et al.* (1985*b*). This observation is confirmed in the present work which was based on ileo-rectal anastomosis with pigs given diets containing two distinct fibre sources alone or in combination at the same total dietary fibre content. In addition, it was clearly demonstrated in the present study that the association of different fibre sources resulted in a highly significant negative interaction with the overall digestibility of amino acids, while additive effects were observed in most cases at the end of the small intestine. The notable depression of overall amino acid digestibility in pigs fed on diet BH could have resulted from two possible effects: (1) a stimulation of bacterial growth in the large intestine, responsible for an increased microbial protein synthesis; (2) an increased flow of endogenous N originating from digestive secretions and cell-wall desquamation.

The fibre fraction originating from soya-bean hulls could be responsible for the first effect. Indeed, the difference between overall and ileal digestibility coefficients for diet H shows that a larger amount of NDF from soya-bean hulls than from wheat bran is fermented in the large intestine. Compared with wheat bran, soya-bean hulls contain a fairly high amount of pectins which have been shown to increase microbial protein synthesis in the large intestine (Rotenberg *et al.* 1982). However, if this was the only mechanism for a reduced N apparent digestibility of diet BH, this would lead to an even larger effect, i.e. a lower digestibility with diet H, and this was not observed.

The second possible effect also needs careful examination. Both quantity (Ecknauer *et al.* 1981) and quality (Jacobs, 1983) of fibre are known to affect the various components of cell-wall proliferation in the small intestine as well as in the colon (Jacobs & Schneeman, 1981). In addition, fibre effects on digestive secretions have been seen, mainly in the case of pancreatic secretion (Schneeman *et al.* 1982; Langlois *et al.* 1987). However, if there was an increased flow of endogenous N into the small intestine, either from the pancreatic flow due to wheat bran (36% increase of daily protein output according to Langlois *et al.* 1987) or from the mucosa of the small intestine, this did not result in a lower ileal N apparent digestibility (diet B *v.* diet H or BH). Therefore, if an increased flow of endogenous N was involved in the depression of overall digestibility of amino acids, it could only take place in the colon. Nevertheless, if such a bran effect was responsible for a reduced N apparent digestibility of diet BH, this would lead to an even larger effect, i.e. a lower digestibility, with diet B, and this was not observed. Thus neither of the two factors considered above seems to be sufficient in itself to explain the negative interaction of fibre-mixing on overall digestibility.

The proportions in which additional faecal amino acids were excreted with diet BH (compared with the average excretion with diets B and H) were close to those of bacteria isolated from faeces of pigs fed on a standard diet (Laplace *et al.* 1985*a*) as shown by  $\chi^2$  distance ( $\chi^2 = 59$ ), calculated according to Guilloteau *et al.* (1983). In contrast, they were different ( $\chi^2 = 383$ ) from that of total faeces of germ-free pigs, taken as an endogenous reference (Laplace *et al.* 1985*a*). Accordingly, the negative interaction in pigs fed on diet BH would result from a stimulated bacterial growth, under several complex influences. Among these, one could consider an increased retention of wheat-bran residues due to some decrease in the rate of passage in the presence of soya-bean hulls (Stanogias & Pearce, 1985*a*).

In contrast to protein and amino acids, there was no interaction between fibre sources on the digestibility of the other dietary constituents (energy, fibrous components) irrespective of the site of digestibility measurement. Furthermore, as expected from the literature (Keys & De Barthe, 1974), the magnitude of the difference between diets was larger in the case of overall digestibility. This suggests that overall digestibility provides a

better discrimination between diets for energy utilization than does ileal digestibility, due to the positive contribution of the hind-gut to fibre digestion.

The comparison of the digestibility coefficients of fibrous components (crude fibre, NDF, ADF) in the dietary treatments shows a higher disappearance of fibre in soya-bean hulls than in wheat bran, especially for the ADF (or lignocellulose) fraction. This is in agreement with previous results obtained in this laboratory (J. M. Pérez, unpublished results) or in the literature (Stanogias & Pearce, 1985*a-c*). It is noteworthy that the digestibility of the NDF fraction in pigs fed on diet H (soya-bean hulls) reached a relatively high value (0.15) at the end of the small intestine, and this is in agreement with recent findings (Lin *et al.* 1987).

From the results of the present study, as well as from the literature, there is no doubt that amino acid digestibility should preferably be measured at the distal end of the small intestine, while energy utilization may be measured at both faecal and ileal levels, in order to assess the partition of digestion between the small and large intestines. But positive or negative interactions between dietary components are likely to occur, depending on the type of fibre sources, their proportions and their associations in the diet. Therefore, the reported effects only refer to the specified fibrous materials used in the experiment. The general occurrence of interactions between the site of digestibility measurement and the dietary treatments, except for lysine, histidine and proline, is evidenced by the fact that the difference in digestibilities between the two sites varies considerably with the type of diet, although in a non-linear fashion. Consequently, the extrapolation from one to the other mode of digestibility measurement may be misleading.

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