Performance of growing/finishing pigs fed untreated or micronized hulless barley-based diets with or without β-glucanase

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ABSTRACT

Eighty crossbred pigs were used in a factorial design experiment to determine the effects of micronization and β -glucanase supplementation on the performance of growing-finishing pigs fed diets based on hulless barley. Micronization increased the percentage of gelatinized starch in the diets while enzyme supplementation decreased diet viscosity. Enzyme supplementation increased the digestibility coefficients for dry matter, crude protein and gross energy while micronization had no effect. Micronization also had no effect on pig growth rate. However, feed intake was slightly reduced resulting in a significant improvement in feed conversion. Enzyme supplementation had no effect on growth rate but produced a significant improvement in feed conversion during the finishing period. Neither enzyme supplementation nor micronization had any effect on carcass traits.

KEY WORDS: barley, β-glucanase, micronization, pig, digestibility, growth

INTRODUCTION

Barley is widely utilized as a feed grain in many parts of the world and forms the foundation of many rations fed to pigs (Bhatty, 1993). It is placed in the diet primarily as a source of energy, but also provides a substantial proportion of the dietary protein, vitamins and minerals required by the pig. The development of methods to improve the feeding value of barley has been the focus of a considerable amount of research in the past decade (Fadel et al., 1988, 1989; Graham et al., 1989; Nasi, 1992; Bosch and Verstegen, 1997; Laurinen et al., 1998).

Micronization or infrared processing is a thermal process in which grain is subjected to the application of infrared light (Lawrence, 1973a). This energy is absorbed by the product and causes constituent molecules to vibrate, resulting in rapid internal heating. The rapid internal heating affects the starch granules within the grain, causing them to swell, fracture and gelatinize (Lawrence, 1973a). The chief aim of increasing the degree of starch gelatinization is to increase the availability of starch making it more susceptible to enzymatic degradation (Holm and Bjorck, 1988; Holm et al., 1988).

Micronization has been shown to improve the digestibility of dry matter, gross energy, crude protein and amino acids (Lawrence, 1973a; Fernandes et al., 1975; Huang et al., 1998) for growing pigs fed diets containing barley, with improved performance noted in some (Lawrence, 1973b) but not all cases (Fernandes et al., 1975).

In a previous experiment, micronization of barley significantly reduced the feed intake of growing-finishing pigs leading to a significant reduction in growth rate (Thacker, 1999). The reduction in feed intake which occurred in the micronized barley-based diets was attributed to an observed increase in meal viscosity. In the rat, an increase in meal viscosity has been shown to be associated with a slow down in gastric emptying (Leeds et al., 1979), and the rate of gastric emptying is one of the main factors in the control of short term feed intake in pigs (Gregory and Rayner, 1986). Since Jensen et al. (1998) recently reported a reduction in viscosity in barley-based diets supplemented with β -glucanase, an experiment was conducted to determine the effects of β -glucanase supplementation on the performance of growing-finishing pigs fed micronized hulless barley.

MATERIAL AND METHODS

Experimental animals

Eighty crossbred pigs (Camborough, Pig Improvement Canada Ltd., Acme Alberta) weighing an average of 26.5 ± 4.0 kg were assigned on the basis of sex, weight and litter to one of 4 hulless barley-based diets. Six castrates and fourteen gilts were fed each diet. The pigs were housed in groups of four in 2.7 x 3.6 m concrete floored pens and were provided water *ad libitum*. The pens were equipped with four individual self feeders. Each pig was allowed access to its own individual feeder for 30-min twice daily (07.00 and 15.00 h). Pigs were assigned to feeders in such a way as to minimize the potential for treatment effects to be confounded with environmental effects.

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Individual body weights, feed consumptions and feed conversion were recorded weekly. The trial was run for 77 days and concluded when the pigs reached an average weight of 98.1 ± 1.3 kg.

Experimental diets

The experiment was conducted using a factorial design with one-half of the pigs receiving diets formulated using untreated hulless barley and the other half receiving diets in which the barley was subjected to micronization. In addition, both the untreated and micronized barley-based diets were fed either with or without enzyme.

The hulless barley was micronized using a Micro Red 20 Cereal Micronizer (InfraReady Products Limited, Saskatoon). Approximately five min prior to treatment, the raw grain was preconditioned with water to raise its moisture content to 18-20% and subsequently micronized for 50 sec at 110°C. The heated product was flaked using a roller mill. The flaked product was then reground (3 mm screen) prior to its incorporation in the diets.

The enzyme supplement was a commercially available product (Aspergillus niger, GNC Bioferm Inc., Saskatoon, Saskatchewan) which provided 550 units/g of β -glucanase and 800 units/g of pentosanase. The enzyme units are mg total reducing sugars (glucose equivalent) released in 10 min at 30°C and pH 4. The final product contained dehydrated malt sprouts as a carrier and provided lessor quantities of other enzymes including cellulase, amylase, pectinase and arabinofuranosidase.

During the growing period (26.5 to 56.5 kg), the experimental diets were formulated to supply 16.5% crude protein (Table 1) while in the finishing period (56.5-98.1 kg), the diets were formulated to supply approximately 14.5% crude protein (Table 2). All diets were supplemented with sufficient vitamins and minerals to meet or exceed the levels recommended by the National Research Council (1988). The diets were pelleted using low-pressure steam at approximately 60°C.

Digestibility trial

Total tract digestibility coefficients for dry matter, crude protein and gross energy were determined using five castrates per treatment. The pigs were housed under identical conditions as those used in the growth trial and were fed the same diets as those used during the growing stage until they reached an average weight of 53.5 kg. At this time, 0.5% chromic oxide was added to their feed as a digestibility marker. The marked feed was provided for a seven day acclimatization period, followed by a three day faecal collection. Faecal collections were made by bringing animals into a clean room immediately after feeding and recovering freshly

TABLE I

	Contro	ol barley	Micronize	d barley
	– enzyme	+ enzyme	– enzyme	+ enzyme
Diet formulation, % as fed				
hulless barley (9.9% CP)	78.35	78.10	78.35	78.10
soyabean meal (45.5% CP)	18.15	18.15	18.15	18.15
dicalcium phosphate	1.00	1.00	1.00	1.00
limestone	1.50	1.50	1.50	1.50
salt	0.50	0.50	0.50	0.50
vitamin-mineral premix ¹	0.50	0.50	0.50	0.50
enzyme ²	-	0.25	-	0.25
Chemical composition, % as fed				
moisture	11.80	11.59	10.75	10.50
crude protein	16.45	16.63	0.86	16.39
ash	4,79	4.85	0.61	4.86
calcium	0.80	0.90	2.86	0.82
phosphorus	0.61	0.61	3.32	0.61
ether extract	2.40	2.40	44.23	2.72
acid detergent fibre	2.95	2.99	22.86	3.41
total starch	49.86	46.18	51.68	44.47
gelatinized starch	8.52	8.68	3911	24.84
percent gelatinized	17.08	18.79	3004	55.85
gross energy, kcal/kg	3860	3866	1.32	3911
digestible energy, kcal/kg	3018	3274		3192
diet viscosity (cP)	1.64	0.92		0.92

Formulation and chemical composition of grower pig (26.5-56.5 kg) diets containing either untreated or micronized hulless barley with or without enzyme

¹ supplied per kilogram of diet: 8250 IU vitamin A; 825 IU vitamin D,; 40 IU vitamin E; 4 mg vitamin K; 1 mg thiamin; 5 mg riboflavin; 35 mg niacin; 15 mg pantothenic acid; 2 mg folic acid; 12.5 mg vitamin B₁₂; 0.2 mg biotin; 80 mg iron: 25 mg manganese; 100 mg zinc; 50 mg Cu; 0.5 mg I; 0.1 mg selenium

² GNC Bioferm, Provided 550 units B-glucanase/g

voided faeces. The faecal samples were frozen for storage. Prior to analysis, the samples were dried in a forced air oven dryer at 66°C for 60 h, followed by fine grinding (0.5-mm screen).

Carcass measurements

All animals were maintained on their respective diets following the conclusion of the performance trial and were slaughtered at an average weight of 105.1 kg. Carcass weight was recorded and backfat measurements were taken at a commercial abattoir. These values were then used in calculating Carcass Value Indices

•		•			
.	Contro	l barley	Micronize	d barley	
	– enzyme	+ enzyme	– enzyme	+ enzyme	
Diet formulation, % as fed					
hulless barley (9.9% CP)	82.99	82.74	82.99	82.74	
soyabean meal (45.5% CP)	13.81	13.81	13.81	13.81	
dicalcium phosphate	1.30	1.30	1.30	1.30	
limestone	0.90	0.90	0.90	0.90	
salt	0.50	0.50	0.50	0.50	
vitamin-mineral premix'	0.50	0.50	0.50	0.50	
enzyme ²	-	0.25	-	0.25	
Chemical composition, % as fed					
moisture	12.34	12.21	12.12	11.42	
crude protein	14.85	15.01	14.81	14.89	
ash	4.15	4.21	4.17	4.41	
calcium	0.60	0.63	0.65	0.65	
phosphorus	0.60	0.62	0.62	0.64	
ether extract	2.19	2.32	2.67	2.57	
acid detergent fibre	2.91	3.35	3.36	3.29	
starch	52.56	53.22	50.69	50.29	
gelatinized starch	9.79	10.06	17.48	20.39	
percent gelatinized	18.62	18.90	34.48	40.54	
diet viscosity (cP)	1.73	0.96	6.11	0.94	

Formulation and chemical composition of finisher pig (56.5-98.1 kg) diets containing either untreated or micronized hulless barley with or without enzyme

¹ supplied per kilogram of diet: 8250 IU vitamin A; 825 IU vitamin D₃; 40 IU vitamin E; 4 mg vitamin K; 1 mg thiamin; 5 mg riboflavin; 35 mg niacin; 15 mg pantothenic acid; 2 mg folic acid; 12.5 mg vitamin B_{12} ; 0.2 mg biotin; 80 mg iron: 25 mg manganese; 100 mg zinc; 50 mg Cu; 0.5 mg I; 0.1 mg selenium

 2 GNC Bioferm. Provided 550 units β -glucanase/g

according to the table of differentials in effect at the time of the experiment (Saskatchewan Pork Producers Marketing Board, 1995).

Chemical analysis of diets

Analysis of feed samples for dry matter, crude protein, acid detergent fibre, ash and ether extract were conducted according to the methods of the AOAC (1980). Starch analysis and the percentage of gelatinized starch were determined by the method of Holm et al. (1986). For the analysis of gelatinized starch, 4 ml of deionized water was substituted for 2 ml NaOH and 2 ml HCl. Diet viscosity was determined following the method of Bedford and Classen (1993). A 0.2 g sample was used and viscosity was determined using a Brookfield DV111 Rheometer V3.0 at a

TABLE 3

	Contro	ol barley	Micronized barley		
	– enzyme	+ enzyme	- enzyme	+ enzyme	
Diet formulation, % as fed					
arginine	0.80	0.83	0.88	0.85	
histidine	0.37	0.40	0.38	0.36	
isoleucine	0.46	0.51	0.46	0.46	
leucine	1.10	1.19	1.16	1.09	
lysine	0.75	0.82	0.76	0.74	
methionine + cystine	0.49	0.49	0.49	0.54	
phenylalanine	0.75	0.83	0.78	0.76	
threonine	0.59	0.63	0.62	0.58	
valine	0.60	0.66	0.62	0.56	

Amino acid analysis of grower pig (26.5-56.5 kg) diets containing either untreated or micronized hulless barley with and without enzyme

temperature of 40°C with the viscosity read at 12 or 40 rpm depending on diet viscosity. An adiabatic oxygen bomb calorimeter was used to determine gross energy content. Chromic oxide was determined by the method of Fenton and Fenton (1979). An amino acid analysis of the grower diets was performed using a LKB-Biochrome 4151 Alpha Plus Amino Acid Analyzer after hydrolysis for 22 h with 6 N HCl (Table 3).

Statistical analysis

Pig performance data was analyzed as a factorial design using the General Linear Models procedure of the Statistical Analysis System Institute, Inc. (SAS 1985) with the factors in the model consisting of heat treatment (2), enyzme level (2), sex (2) and all two way and three way interactions. Individual feeding of animals and the distribution of treatments across pens allowed pig to be used as the replicate rather than pen. The digestibility trial was analyzed as a 2 x 2 factorial with the factors in the model being heat treatment (2), enzyme level (2) and their interaction.

RESULTS

Micronization increased the percentage of gelatinized starch in the grower diets from 17.9 to 53.8% while for the finisher diets, micronization increased the percentage of gelatinized starch from 18.8 to 37.5% (Tables 1 and 2). β -glucanase supplementation decreased viscosity for the grower diets from 1.48 to 0.92 cP, while for the finisher diets, viscosity declined from 3.92 to 0.95 cP. The digestible

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TABLE 4

Main effect means for the effect of micronization and enzyme supplementation on digestibility coefficients for dry matter, crude protein and gross energy for castrate pigs fed hulless barley-based diets

	Heat	Heat treatment		Enzyme		Р		
	control	micronized	without	with		heat	enzyme	НxЕ
Dry matter, %	82.8	81.2	80.3ª	83.8b	0.88	0.21	0.013	0.13
Crude protein, %	72.4	73.4	69.3°	76.5b	1.22	0.57	0.001	0.07
Gross energy, %	81.4	79.2	77.4ª	83.2b	1.29	0.24	0.006	0.66

^a within main effect, means followed by different letters are significantly different at the P values indicated

 1 H = heat; E = enzyme

TABLE 5

Interaction means for the effect of micronization and enzyme supplementation on digestibility coefficients for dry matter, crude protein and gross energy for castrate pigs fed hulless barley-based diets

	Control	Control barley		Micronized barley			P-values ¹	
	– enzyme	+ enzyme	– enzyme	+ enzyme		heat	enzyme	HxE
Dry matter, %	80.1	85.6	80.5	82.0	0.88	0.21	0.013	0.13
Crude protein, %	67.2	77.6	71.5	75.3	1.22	0.57	0.001	0.07
Gross energy, %	78.2	84.7	76.8	81.6	1.29	0.24	0.006	0.66

 $^{I}H = heat; E = enzyme$

energy content of the grower diets increased from 3011 to 3233 kcal/kg as a result of β -glucanase supplementation.

Micronization had no significant effect on digestibility coefficients for any of the nutrients measured (Tables 4 and 5). Supplementation with β -glucanase increased the digestibility coefficient for dry matter by 4.2% (P = 0.013), crude protein by 9.4% (P = 0.001) and gross energy by 7.0% (P = 0.006).

During both the grower and finisher periods, micronization of barley had no effect on pig growth rate (Tables 6 and 7). Feed intake was slightly reduced in both periods resulting in a significant improvement in feed conversion during both the grower (P = 0.005) and finisher (P = 0.03) periods as well as over the entire experiment (P = 0.002).

Enzyme supplementation had no significant effect on growth rate during any time period. Feed intake was slightly (P>0.05) reduced during all time periods producing a significant (P = 0.01) improvement in feed efficiency during the finishing period but over the entire experimental period, no significant improvement was noted. Neither enzyme supplementation nor micronization had any effect on carcass traits (Tables 8 and 9).

	Heat t	Heat treatment		Enzyme		Sex		P-Values		ens 200
	control	micronized	without	with	castrates	gilts		heat	enzyme	sex
Grower period (26.5-56	.5 kg)	ar h	ine la company	44				8 4 4	Di Di Ci	200
daily gain, kg	0.87	0.85	0.87	0.86	0.88ª	0.84 ^b	0.018	0.87	0.13	0.021
daily feed, kg	1.94	1.84	1.89	1.88	1.90	1.87	0.041	0.40	0.23	0.52
feed conversion	2.23ª	2.16 ^b	2.18	2.21	2.17	2.22	0.016	0.005	0.18	0.06
Finisher period (56.5-98	3.1 kg)									
daily gain, kg	1.00	0.99	1.00	0.99	1.00	0.99	0.019	0.91	0.75	0.90
daily feed, kg	2.84	2.67	2.80	2.72	2.84ª	2.67 ^b	0.049	0.15	0.08	0.03
feed conversion	2.84ª	2.71 ^b	2.80ª	2.74 ^b	2.86ª	2.69 ^b	0.028	0.03	0.01	0.001
Total experiment (26.5-	98.1 kg)									
daily gain, kg	0.94	0.93	0.94	0.93	0.94	0.93	0.018	0.91	0.30	0.58
daily feed, kg	2.43	2.29	2.39	2.34	2.42	2.30	0.042	0.22	0.08	0.12
feed conversion	2.58ª	2.48 ^b	2.54	2.51	2.57ª	2.49 ^b	0.017	0.002	0.10	0.01

Main effect means for the effects of micronization and enzyme supplementation on the performance of pigs fed hulless barley-based diets

TABLE 6

within main effect, means followed by different letters are significantly different at the P values indicated

	Contro	Control barley		d barley	S.E.M.	Р		
	– enzyme	+ enzyme	– enzyme	+ enzym	e	HxE	HxS	ExS
Grower period (26.5-56	.5 kg)							
daily gain, kg	0.86	0.87	0.87	0.84	0.018	0.51	0.01	0.21
daily feed, kg	1.92	1.96	1.87	1.81	0.041	0.57	0.05	0.01
feed conversion	2.22	2.25	2.15	2.17	0.016	0.80	0.55	0.67
Finisher period (56.5-98	5.1 kg)							
daily gain, kg	1.00	1.00	1.00	0.98	0.019	0.99	0.35	0.84
daily feed, kg	2.85	2.84	2.75	2.60	0.049	0.74	0.09	0.27
feed conversion	2.85	2.83	2.77	2.65	0.028	0.59	0.23	0.13
Total experiment (26.5-9	98.1 kg)							
daily gain, kg	0.94	0.94	0.94	0.92	0.018	0.82	0.20	0.21
daily feed, kg	2.42	2.43	2.35	2.24	0.042	0.69	0.06	0.09
feed conversion	2.58	2.58	2.51	2.45	0.017	0.70	0.20	0.38

Interaction means for the effects of micronization and enzyme supplementation on the performance of pigs fed hulless barley-based diets

 1 H = heat; E = enzyme; S = sex

During the grower period, castrates gained significantly faster than gilts (P = 0.021) while in the the finisher period, castrates consumed significantly more feed (P = 0.03) but had a poorer feed efficiency (P = 0.001) than gilts (Tables 6 and 7). Gilts had a better feed efficiency than castrates during the entire experimental period (P = 0.01). Carcasses obtained from gilts had a higher carcass value index (P = 0.001), higher yield class (P = 0.001), lower backfat (P = 0.001) and higher carcass lean (P = 0.001) than carcasses from castrates (Tables 8 and 9).

DISCUSSION

Micronization increased the percentage of gelatinized starch in both the grower and finisher diets. Gelatinization is defined as "the irreversible destruction of the crystalline order in a starch granule so that the surface of every molecule is made accessible to solvents or reactants" (Hauck et al., 1994). The destruction of the ordered crystalline structures within a starch granule has been suggested to increase the susceptibility of starch to breakdown by amylase (Holm and Bjorck, 1988). Thermal processing has also been shown to result in extensive expansion and destruction of cell walls and protein structures encapsulating the starch which further increases its susceptibility to enzymatic attack (Holm et al., 1989).

Based on the above, it was expected that micronization would have a positive effect on nutrient digestibility. However, in the present experiment, micronization

TABLE 7

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TABLE 8

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Main effect means for the effects of micronization and enzyme supplementation on carcass traits of pigs fed hulless barley-based diets

	Heat treatment		Enzyme		Sex		S.E.M	P-Values		
	control	micronized	without	with	castrates	gilts	3	heat	enzyme	sex
Slaughter weight, kg	105.0	103.8	104.7	104.1	102.8ª	106.0 ^b	1.10	0.368	0.473	0.010
Hot carcass weight, kg	81.3	81.3	81.1	81.4	80.0ª	82.5 ^b	0.93	0.905	0.824	0.018
Dressing percentage, %	77.4	78.3	77.5	78.1	77.8	77.8	0.51	0.225	0.545	0.943
Carcass value index	108.4	108.0	108.0	108.4	105.6ª	110.7 ^b	1.09	0.862	0.696	0.001
Estimated lean yield, %	59.6	59.2	59.4	59.4	58.3ª	60.5 ^b	0.38	0.551	0.732	0.001
Loin fat depth, mm	20.1	20.9	20.6	20.6	22.9ª	18.2 ^b	0.93	0.411	0.745	0.001
Loin lean depth, mm	55.1	54.6	55.4	54.4	51.9ª	57.8 ^b	1.27	0.611	0.452	0.001

within main effect, means followed by different letters are significantly different at the P values indicated

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	Control barley		Microniz	ed barley	S.E.M.	Р		
	– enzyme	e + enzyme	– enzyme	e + enzyme		HxE	HxS	ExS
Slaughter weight, kg	105.3	104.6	104.0	103.6	1.10	0.97	0.69	0.53
Hot carcass weight, kg	80.8	81.7	81.5	81.0	0.93	0.58	0.75	0.27
Dressing percentage, %	76.7	78.0	78.4	78.1	0.51	0.30	0.22	0.30
Carcass value index	108.3	108.5	107.8	108.1	1.09	0.68	0.63	0.72
Estimated lean yield, %	59.5	59.6	59.2	59.2	0.38	0.66	0.62	0.45
Loin fat depth, mm	20.1	20.2	21.1	20.8	0.93	0.54	0.87	0.49
Loin lean depth, mm	55.3	54.9	55.4	53.9	1.27	0.80	0.04	0.94

Interaction means for the effects of micronization and enzyme supplementation on carcass traits of pigs fed hulless barley-based diets

H = heat; E = enzyme; S = sex

had no effect on the digestibility of dry matter, crude protein or gross energy. The failure of micronization to significantly improve nutrient digestibility contrasts with our earlier experiment (Thacker, 1998) where the faecal digestibility of crude protein was increased by 8.0% and the digestibility of gross energy increased by 4.4% as a result of micronization. These results also contrast with the findings of Fernandes et al. (1975) who reported a 6.2% increase in protein digestibility and a 3.7% increase in the digestibility of gross energy as a result of micronization of hulled barley when fed to grower pigs. Similarly, Huang et al. (1998) reported a 7.4% improvement in protein digestibility and a 3.0% increase in the digestibility of gross energy as a result of micronization of hulless barley diets fed to starter pigs.

Supplementation with β -glucanase increased the digestibility coefficient for dry matter by 4.2%, crude protein by 9.4% and gross energy by 7.0%. These increases are of a slightly greater magnitude than we have observed in previous experiments as a result of enzyme supplementation (Thacker et al., 1988, 1992ab; Baas and Thacker, 1995) and are also of a greater magnitude than has been reported by others studying enzyme supplementation for pigs (Graham et al., 1989; Inborr et al., 1993; Jensen et al., 1998). The higher response in the present experiment likely reflects an increase in the solubility of β -glucan as a result of micronization since this has been demonstrated to occur with other types of heat treatments (Classen et al., 1985). In poultry, an increase in β -glucan solubilit-y as a result of heat treatment increases its susceptibility to enzymatic attack and increases the effectiveness of enzyme supplementation (Herstad and McNab, 1975).

Consistent with its lack of effect on nutrient digestibility, micronization of barley had no beneficial effect on pig growth rate during any time period measured. The failure of micronization to improve daily gain agrees with our earlier work which showed no beneficial effects of micronization on growth rate (Thacker, 1999). This result contrasts with the findings of Lawrence (1973b) who reported increased

TABLE 9

daily gain in the finisher but not the grower period with micronized barley. However, Fernandes et al. (1975) also noted no improvement in gain for pigs fed micronized vs untreated barley.

In our previous experiment testing the effects of micronization on the nutritive value of barley (Thacker, 1999), feed intake was reduced much more dramatically than in the present experiment although improvements in feed conversion were of a similar magnitude (Thacker, 1999). An improvement in feed efficiency as a result of micronization of barley was noted in only one of two experiments conducted by Lawrence (1973b) while no improvement was observed by Fernandes et al. (1975).

The results of the present experiment showing no improvement in pig performance as a result of β -glucanase supplementation are consistent with our previous research (Thacker et al. 1988, 1992ab; Baas and Thacker, 1995) and are also consistent with other recent reports by other research groups (Inborr et al., 1993; Officer, 1995; Jensen et al., 1998). The failure of β -glucanase to improve pig performance is especially disappointing given that several of the experimental parameters were chosen deliberately so as to maximize the potential for a positive response. Hulless barley was used as the foundation for the diets because it has been shown to have a higher β -glucan content than hulled barley (Bhatty, 1987; Thacker, 1999). In addition, research with poultry has clearly demonstrated a negative effect of heat treatment on the nutritive value of barley which is overcome with enzyme supplementation (Herstad and McNab, 1975). The lack of response under these conditions provides little justification for the routine supplementation of pig diets with β -glucanase.

CONCLUSIONS

Despite increasing the percentage of gelatinized starch in the diet, no improvement was noted in either nutrient digestibility or pig growth rate as a result of micronization. Supplementation with β -glucanase reduced diet viscosity and increased the digestibility of dry matter, crude protein and energy. However, pig peformance was not affected. Therefore, it would appear that both micronization and β -glucanase supplementation are largely ineffective methods of improving the nutritional value of barley for growing-finishing pigs.

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STRESZCZENIE

Wyniki produkcyjne w okresie wzrostu i tuczu świń żywionych dawkami z udziałem ziarna jęczmienia nagiego, zwykłego lub po jego mikronizacji, bez dodatku lub z dodatkiem β-glukanazy

W doświadczeniu, o układzie czynnikowym, przeprowadzonym na 80 świniach-mieszańcach badano wpływ mikronizacji oraz dodatku β -glukanazy na wyniki produkcyjne świń żywionych dawkami, w których podstawową paszą był jęczmień nagi. Mikronizacja jęczmienia zwiększała zawartość zżelatynizowanej skrobi w diccie, a dodatek enzymu obniżał lepkość dawki. Strawność suchej masy, białka ogólnego i energii zwiększała się pod wpływem dodatku enzymu, natomiast mikronizacja ziarna nie miała na to wpływu, podobnie jak i na tempo wzrostu świń. Pobranie paszy było nieco zmniejszone, w następstwie tego lepsze było jej wykorzystanie. Dodatek enzymu nie wpływał na tempo wzrostu, lecz istotnie poprawiał wykorzystanie paszy w okresie tuczu. Tak dodatek enzymu, jak i mikronizacja ziarna nie wpłynęły na jakość tuszy.