

Effect of roasted cereals and soybean in feed mixtures on fattening and slaughter traits as well as blood composition in fattening lambs

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Abstract

The study examined the effect of raw and roasted cereals (corn, oat and barley) and soybean in diet on growth performance and blood parameters of lambs fattening. Totally 60 lambs after ab lactation at average of 50 days, were tested and equally divided into three groups: an experimental I (E I), experimental II (E II) and a control (C) group. The fattening period lasted 56 days. Lambs of the C group were fed on mixture with crude forages, those from the E I on mixture with roasted cereals and lambs of the E II on mixture with both roasted cereals and soybean. The lambs from E I and E II groups grew faster (by 13.62 and 19.40%), gained higher final body weight (by 6.72 and 8.08%) and spent less food per kg of daily gain weights (by 14.03 and 13.24%) although feed consumption in E I was lower (by 2.33%) but in E II higher (3.49%) in comparison with the C group. The blood serum concentrations of cholesterol-total and cholesterol-HDL were lower for lambs E I groups compared with lambs C and E II groups. ALP activity was higher for lambs E I and E II groups compared with C group. Urea concentrations were significantly ($P < 0.05$) lower, and that of albumin higher in E II group compared to C group. There were no differences among groups for the serum minerals content, other biochemical indicators and enzymes. Obtained differences in slaughter properties of lambs (carcass weight and dressing) indicate their better productive effects. Concerning the gained production results may be concluded that the use of roasted cereals and soybean in feed mixture in lambs fattening is justified.

Keywords: lambs, roasted cereals, soybean, fattening traits, slaughtering traits, blood parameters

Zusammenfassung

Einfluss von gerösteten Getreiden und Sojabohnen in Futtermischungen auf Mast- und Schlachteigenschaften von Lämmern

Verglichen wurde der Einfluss von unbehandelten und gerösteten Getreiden sowie Sojabohnen in Futtermischungen auf Wachstums-, Schlachteigenschaften und Blutparameter von Lämmern. Der 60 Lämmer umfassende Versuch mit drei jeweils identischen Gruppen dauerte vom Absetzen der Lämmer im Alter von 50 Tagen

insgesamt 56 Tage. Die Kontrollgruppe C erhielt Futtermischungen aus unbehandeltem Getreide, die Versuchsgruppe E I geröstetes Getreide und die Gruppe E II geröstetes Getreide sowie geröstete Sojabohnen. Die Lämmer der Gruppen E I und E II erreichten signifikant höhere Tageszunahmen (13,62 bzw. 19,40%), Endgewichte (6,72 bzw. 8,08%) und eine bessere Futtereffizienz von 14,03 bzw. 13,24% bei tendenziell geringfügig geringerem Futterverbrauch (2,33 bzw. 3,49%) als Tiere der C Gruppe. Tiere der Gruppe E I wiesen gegenüber E II und C geringere Blutwerte für Gesamtcholesterin und HDL Cholesterin auf. Im Serum der Gruppen E I und E II wurde gegenüber C eine größere Alkaline Phosphatase Enzymaktivität festgestellt. Signifikant niedrigere Harnstoffkonzentration und höhere Albuminkonzentration ergab sich bei E II gegenüber C Lämmern. Keine Unterschiede zwischen den Gruppen fanden sich beim Mineralstoffgehalt, biochemischen Indikatoren und Enzymen. Außer den Schlachtgewichten wurden keine Unterschiede in den Schlachtmerkmalen beobachtet. Es wird geschlussfolgert, dass die Verwendung von geröstetem Getreide und gerösteten Sojabohnen sich vorteilhaft auf die Produktionsmerkmale von Lämmern auswirkt.

Schlüsselwörter: Lamm, Getreide geröstet, Sojabohnen geröstet, Mastmerkmale, Schlachteigenschaften, Blutparameter

Introduction

Young lambs have high energy and protein requirements and consume limited amounts of dry matter. Main energy source in fattening lambs rations are cereals, primarily corn and oat whereas soybean is the proteins source. Due to some harmful ingredients (anti-nutritive factors) which reduce the soybean feeding value and cause undesired health consequences its ration portion is limited. In lambs, the rumen does not function fully until about 7 weeks of age, thus, the very young lambs are functionally non-ruminants (SCHWULST 1996, ANTUNOVIĆ *et al.* 2000) and are more sensitive to various stress conditions (ablactation, feed changes etc.). They may also cause diverse rumen microflora disorders leading to development of some harmful microorganisms and lambs growth decrease. Forage heat treatment is known to reduce harmful bacteria (DOMAĆINOVIĆ *et al.* 1996) thus contributing to easier transition from milk feeding to dry food. Crude soybean anti-nutritive factors (especially tripsine and kimotripsine) can significantly reduce nitrogen utilization and have a depressive effect on body weight (LOESCHE *et al.* 1992). Heat treatments of legume seeds can inactivate anti-nutritional factors (VAN DER POEL *et al.* 1990), reduce activity of urease, being a thermo-sensitive enzyme, and improve feeding value (MONARI 1990). Heat treatments of cereals improve utilization of starch affecting positively feed efficiency (GRUBIĆ 1987, PATTANAIK *et al.* 2000). In cereal grains, the effect of heat treatments is usually to increase rumen degradability of starch (THEURER *et al.* 1999). Blood characteristics settlement is a worthwhile indicator of animals' state according to nutrition, digestion and resorption of nutrients. There are no many scientific papers about application of heat-treated forage in lamb rations. Therefore, the aim of our investigation was to establish application of heat-treated forage in lamb rations and their effect on fattening and slaughter traits as well as blood composition.

Material and methods

Animals and diets

Biological investigations were carried out with 60 lambs of Tsigai breed after ablation at average age of 50 days. The experimental lambs were classified into three groups with 20 lambs in group of which control (C) and two experimental ones (EI and EII). Lambs were of the equal age, body weight, sex and origin. They were kept in the same conditions during the experiment. The fattening period lasted 56 days. Lambs of the control group were fed on mixture with grounded forages, those from the experimental group 1 on mixture with roasted cereals (corn, oat and barley) and lambs of the experimental group 2 on mixture with both roasted cereals and soybean (Table 1). Meadow hay and water were consumed *ad libitum*. Cereals and soybean roasted using a Jet-Pro Roaster (Jet-Pro Co. Atchison, KS) to an exit temperature of either 135°C during 6 min. Considering the aim of the investigation, forages used in feed mixtures, their raw material and chemical composition (analysed according to AOAC 1995) can be seen in Table 1. Metabolic energy was determined according to DLG (1993).

Table 1
Raw material and chemical composition of feed mixtures
Bestandteile und Inhaltstoffe der Futtermischungen

Composition, %	Feed mixtures		
	Control	Experiment I	Experiment II
Raw corn	25.00	–	–
Roasted corn	–	25.00	25.00
Oat	16.00	–	–
Roasted oat	–	16.00	16.00
Barley	20.00	–	–
Roasted barley	–	20.00	20.00
Soybean	31.00	31.00	–
Roasted soybean	–	–	31.00
Wheat forage meal	5.00	5.00	5.00
Phosphonal	1.00	1.00	1.00
Limestone	1.00	1.00	1.00
Salt	0.50	0.50	0.50
VAM (vitamin and mineral mixture)*	0.50	0.50	0.50
Total	100.00	100.00	100.00
Dry matter	85.96	85.94	85.97
Crude fat	3.80	3.72	3.78
Crude ash	5.31	5.38	5.34
Crude fibres	8.13	8.17	8.15
Crude proteins	15.50	15.52	15.47
Crude proteins/dry matter	18.03	18.06	17.99
Metabolic energy, MJ ME/kg	12.33	12.33	12.33
g crude proteins/1 MJME	12.6	12.6	12.5
Calcium	0.92	0.91	0.93
Phosphorus	0.44	0.46	0.45

*contents in 1 kg = vitamin A: 1 000 000 IU/g, vitamin D3: 150 000 IU/g, vitamin E: 1 500 mg; vitamin K3: 50 mg, vitamin B1: 100 mg, vitamin B2: 200 mg, nicotin acid: 1 000 mg, pantothenic acid: 500 mg, vitamin B6: 200 mg, vitamin B12: 1.2 mg; cholin chloride: 20 000 mg, Fe: 4 000 mg, Cu: 800 mg, Mn 3 500 mg, Zn 5 000 mg, I: 80 mg, Co: 20 mg, Se: 15 mg, Mg: 10 000 mg, S: 10 000 mg, antioxidant: 10 000 mg.

Sampling and analyses

Body weight, average daily gain, daily feed consumption and feed efficiency (expressed as kg of body weight gain per kg feed intake) during postweaning period were determined for lambs on 28 and 56 days of fattening. The lambs were weighed at the beginning of the experiment, on 28th day and at the end of the experiment (56th day).

The blood was collected from the jugular vein (10ml) into the sterile vacuum tubes Venoject (Sterile Terumo Europe, Leuven, Belgium) from all lambs to hours after feeding. After that, the serum was separated by centrifugation (20 min) at 3 500 revolutions/min. Within the blood serum there have been found the concentrations of the mineral indicators (Ca – Calcium, P – inorganic Phosphorus, K – Potassium, Na – Sodium, Cl – Chloride and Fe – Iron, concentrations of the biochemical indicators (urea, glucose, total proteins, albumin, cholesterol, high-density lipoproteins or HDL-cholesterol, triacylglycerols and bilirubin total) and enzyme activity (ALT – alanine aminotransferase, AST – aspartate aminotransferase, alpha-amylase, ALP – alkaline phosphatase, CK – creatine kinase, GGT – γ -glutamyl transferase, cholinesterase and LDH – lactate dehydrogenase). All biochemical values were determined with Olympus System Reagents (OSR), manufactured and distributed by Olympus Diagnostic GmbH (Irish Branch), Lismeehan, Ireland, manufactured for Olympus Diagnostic GmbH, Hamburg, using OLYMPUS AU 600 apparatus.

Animals used in this study were maintained in facilities approved by the Croatian Association for Accreditation of Laboratory Animal Care, and in accordance with current regulations and standards issued by the Croatian Ministry of Agriculture.

Slaughter and meat analysis

Ten lambs from each group were randomly chosen by the method of random sample and slaughtered at the end of fattening. Warm carcass weight with viscera, edible viscera weight (heart, lungs, liver and spleen) were determined after performed slaughtering processing of carcasses (within 60 min). Weight of cold carcass with viscera was determined after 24 h of cooling at a temperature of 4°C in the ham depth. Dressing percentage 1 and 2 were calculated on the basis of body weight before slaughtering and weight of warm and cold carcasses with viscera. Cooling loss was calculated on the basis of warm carcass weight with viscera and weight of cold carcass with viscera. Meat quality was determined on the *musculus longissimus dorsi* (MLD) samples taken between 12th and 13th rib. Meat values of pH₀, pH₁ and pH₂ were determined by the contact pH meter Mettler Toledo immediately after carcass dressing, 45 min later and 24 h again *post mortem*. Water holding capacity was measured based on method proposed by SIERRA (1973). In duplicate, muscle samples weighing about 300 mg were placed between two pieces of filter paper previously weighed (*P1*), and pressed for 5 min, using a weight of 3.4 kg. The muscle samples were then removed, and the paper was re-weighed (*P2*). Water holding capacity was determined as:

$$WHC(\%) = \frac{P2 - P1}{S \cdot 100} \quad (1)$$

where *S* represents the sample weight.

Slaughtering quality indicators of lamb carcasses: carcass length 1 (*os pubis - atlas*) and 2 (*os pubis - first rib*); carcass circumference 1 (by shoulder) and 2 (by ham); ham length (*tuber calcanei - tuberculum ossis ischii*); ham circumference (the widest part) and ham index was calculated on the basis of ham circumference and its length) were measured after lamb carcass processing on slaughtering line according to Figure 1.

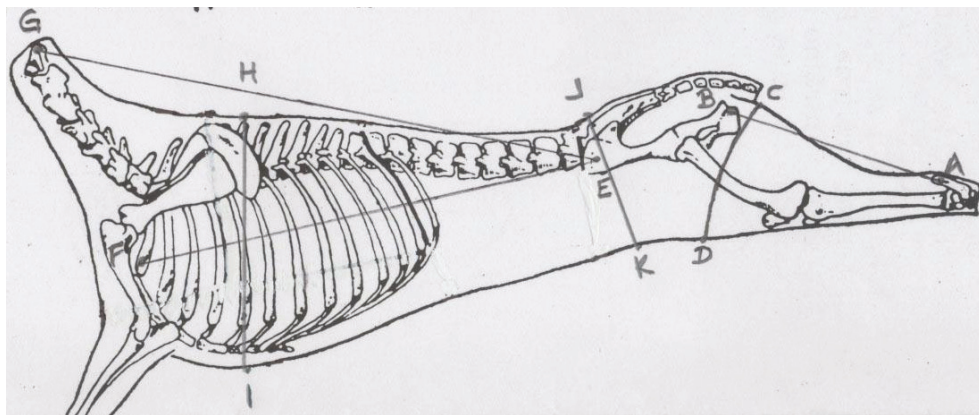


Figure 1

Measuring scheme of indicators of lamb carcass slaughtering quality

A-B = ham length (*tuber calcanei - tuberculum ossis ischii*), C-D = ham circumference (the widest part), E-G = carcass length 1 (*os pubis - atlas*), E-F = carcass length 2 (*os pubis - first rib*), H-I = carcass circumference 1 (by shoulder), J-K = carcass circumference 2 (by ham)

Maße am Schlachtkörper

Of the indicators of lamb carcass slaughtering quality which were estimated by subjective method, conformation and fatness including kidney knob and channel fat were analysed. They were evaluated according to ŽIVKOVIĆ *et al.* (1981) using points from 1 to 5 (mark 1 = extremely deficient, mark 2 = deficient, mark 3 = moderate, mark 4 = good and mark 5 = excellent). Chemical analyses of meat (MLD) after freeze-drying determined by AOAC (1995).

Statistical analysis

Statistical analysis of data was performed by computer program Statistica 7.1 (StatSoft Inc 2005). Levels of significance of differences between groups of lambs were determined by a one-way ANOVA. Data were statistically evaluated using Duncan's post hoc multiple range tests. A value of $P < 0.05$ was considered significant.

Results

Lambs fed on mixture with roasted cereals and soybean were characterized by significantly ($P < 0.05$) higher body weight on 28th day of fattening (by 6.85%) compared to the control groups (Table 2).

Lambs of the experimental group II had very significantly ($P < 0.01$) higher, those of the experimental group I statistically significant ($P < 0.05$) higher body weight at the end of the experiment by 6.72% and 8.08% compared to the control group lambs. Statistically

very significant ($P < 0.01$) higher daily gains obtained on the 28th (by 23.51 and 26.12%), from 28th to 56th day (by 9.44 and 12.66%) as well as on the average from 1st to 56th day (by 13.62 and 19.40%) were accomplished by the lambs from the experimental groups. At the same time, lambs of the experimental group II had significantly higher ($P < 0.05$) average daily gains from 1st to 56th fattening day compared to the lambs from the experimental group I.

Daily consumed feed and its efficiency can be seen in Table 3.

Table 2
Average body weights and daily gains
Körpergewichte und tägliche Zunahmen

Indicators	Statistical parameters	Groups		
		Control (n=20)	Experiment I (n=20)	Experiment II (n=20)
Body weight, kg				
1 st day	Mean	16.82	16.84	16.77
	SD	2.30	2.26	2.25
	%	100.00	100.12	99.71
28 th day	Mean	23.08 ^a	24.46 ^{ab}	24.66 ^b
	SD	2.63	2.77	2.99
	%	100.00	105.98	106.85
56th day	Mean	29.32 ^{Aa}	31.29 ^b	31.69 ^B
	SD	3.01	2.88	2.93
	%	100.00	106.72	108.08
Daily gain, g				
1st to 28th day	Mean	223.57 ^A	276.13 ^B	281.96 ^B
	SD	39.26	44.75	47.29
	%	100.00	123.51	126.12
29th to 56th day	Mean	222.86 ^{Aa}	243.89 ^b	251.07 ^B
	SD	40.66	39.61	29.77
	%	100.00	109.44	112.66
1st to 56th day	Mean	223.21 ^A	253.60 ^{Bb}	266.52 ^{Cb}
	SD	28.89	23.57	27.81
	%	100.00	113.62	119.40

Means with different superscript letters differ significantly: ^{a,b} $P < 0.05$, ^{A,B,C} $P < 0.01$. SD standard deviation

Table 3
Average daily feed consumption and conversion
Futteraufnahme und -verwertung

Indicators	Statistical parameters	Groups		
		Control (n=20)	Experiment I (n=20)	Experiment II (n=20)
Daily food consumption, kg/day				
Feed mixture	Mean	0.86	0.84	0.89
	%	100.00	97.67	103.49
Feed efficiency, kg/kg				
Feed mixture	Mean	3.85	3.31	3.34
	%	100.00	85.97	86.76

Lambs of the E I group received less feed by 3.33% whereas those from the E II group consumed more feed by 3.49% compared to the C group. Feed efficiency was better by 14.13 and 13.24% in lambs from the E I and E II groups compared to the C group.

Table 4
Mineral indicators in lambs blood serum
Mineralstoffgehalt im Blutserum

Indicators	Statistical parameters	Groups		
		Control (n=20)	Experiment I (n=20)	Experiment II (n=20)
Ca, mmol l ⁻¹	Mean	2.28	2.32	2.28
	SD	0.22	0.21	0.19
P-inorganic, mmol l ⁻¹	Mean	2.97	3.24	3.21
	SD	0.46	0.40	0.36
Na, mmol l ⁻¹	Mean	132.90	134.40	129.90
	SD	9.07	8.06	8.56
K, mmol l ⁻¹	Mean	4.42	4.49	4.32
	SD	0.28	0.52	0.48
Cl, mmol l ⁻¹	Mean	93.60	93.20	90.70
	SD	6.04	5.87	6.06
Fe, µmol l ⁻¹	Mean	27.99	28.73	30.65
	SD	5.27	6.33	4.91

SD standard deviation

Table 5
Biochemical indicators in lambs blood serum
Biochemische Indikatoren im Blutserum

Indicators	Statistical parameters	Groups		
		Control (n=20)	Experiment I (n=20)	Experiment II (n=20)
Urea, mmol l ⁻¹	Mean	11.23 ^a	10.23 ^{ab}	9.45 ^b
	SD	2.45	1.46	1.61
Glucose, mmol l ⁻¹	Mean	1.93	1.96	2.09
	SD	0.44	0.39	0.34
Bilirubin, µmol l ⁻¹	Mean	4.90	4.30	3.80
	SD	2.02	1.06	1.03
Albumins, g l ⁻¹	Mean	28.49 ^a	29.63 ^{ab}	30.61 ^b
	SD	2.21	2.03	2.12
Total proteins, g l ⁻¹	Mean	55.42	55.50	59.17
	SD	5.19	5.37	6.63
Cholesterol- total, mmol l ⁻¹	Mean	0.99 ^A	0.72 ^{Ba}	0.96 ^b
	SD	0.18	0.18	0.26
Cholesterol- HDL, mmol l ⁻¹	Mean	0.62 ^A	0.46 ^B	0.59 ^{AC}
	SD	0.10	0.12	0.14
Triacylglycerols, mmol l ⁻¹	Mean	0.39	0.30	0.36
	SD	0.25	0.31	0.18

Means with different superscript letters differ significantly: ^{a,b}P<0.05, ^{A,B,C}P<0.01. SD standard deviation

Analysis of blood serum biochemical indicators (Table 5) in lambs fed on roasted cereals determined very significantly ($P<0.01$) lower concentration of total and HDL cholesterol compared to the lambs from other treatments. Similar trend was also determined for triglyceride concentration, and differences between treatments were not significant. Urea concentrations were statistically significant ($P<0.05$) lower, and that of albumin higher in lambs fed on rations with roasted cereals and soybean compared to the control group. Slightly lower concentration glucose obtained in blood serum of lambs feeding with raw cereals and soybean from lambs feeding with roasted cereals and soybean. Very significantly ($P<0.01$) higher activity of ALP and CK enzymes appeared to be in lambs fed on roasted cereals and soybean compared to the control lambs group (Table 6).

Table 6
Enzyme activities in lambs blood serum
Enzymaktivitäten

Enzymes, U l ⁻¹	Statistical parameters	Groups		
		Control (n=20)	Experiment I (n=20)	Experiment II (n=20)
Alpha amylase	Mean	12.50	13.70	14.60
	SD	3.21	3.30	4.65
Alanine aminotransferase	Mean	36.10	38.50	30.10
	SD	10.32	18.26	18.90
Aspartate aminotransferase	Mean	160.50	150.00	174.50
	SD	31.98	56.55	60.62
Alkaline phosphatase	Mean	196.40 ^{Aa}	249.00 ^b	266.50 ^B
	SD	40.57	35.77	66.50
Creatine kinase	Mean	344.80 ^a	462.00 ^{ab}	526.20 ^b
	SD	138.50	161.11	185.36
γ-Glutamyl transferase	Mean	67.20	71.70	68.10
	SD	14.59	13.48	13.17
Cholinesterase	Mean	118.50	113.90	124.50
	SD	14.00	19.14	12.19
Lactate dehydrogenase	Mean	598.90	685.70	683.30
	SD	116.18	72.73	149.22

Means with different superscript letters differ significantly: ^{a,b} $P<0.05$, ^{A,B} $P<0.01$. SD standard deviation

Analysis of the slaughter indicators showed very significantly ($P<0.01$) higher body weight of the lambs prior slaughter, weights of both warm and cold trunks with innards in lambs from the E I and E II groups compared to the lambs from the C group (Table 7).

Very significantly ($P<0.01$) higher dressing of the warm trunk with innards was characterized by the lambs of the experimental groups as well as higher digestive system weight in lambs of the experimental group II compared to the control group. As for the linear measures and lamb trunks quality, no significant differences ($P>0.05$) were found between feeding treatments although lambs fed on roasted cereals and soybean had somewhat more qualitative trunks (Table 8 and 9).

Table 7
Slaughtering characteristics of lambs' carcasses
Schlachtmerkmale

Indicators	Statistical parameters	Groups		
		Control (n=10)	Experiment I (n=10)	Experiment II (n=10)
Slaughter weight, kg	Mean	29.20 ^A	31.06 ^B	31.49 ^B
	SD	1.51	1.44	1.34
Weight of warm carcass, kg	Mean	15.10 ^A	16.50 ^B	17.04 ^B
	SD	0.74	0.85	0.34
Dressing percentage 1, %	Mean	51.76 ^{Aa}	53.12 ^b	54.17 ^B
	SD	2.17	1.22	1.54
Weight of warm innards, kg	Mean	1.14	1.17	1.25
	SD	0.16	0.24	0.23
Weight of digestive tract, kg	Mean	5.80 ^a	6.12	6.47 ^b
	SD	0.37	0.58	0.49
Weight of cold carcass, kg	Mean	14.58 ^A	15.86 ^B	16.32 ^B
	SD	1.27	0.83	0.27
Dressing percentage 2, %	Mean	50.00	51.06	51.89
	SD	4.43	0.99	1.63
Cooling loss, g/kg	Mean	34.91	38.82	42.19
	SD	7.53	6.32	3.86
Weight of cold innards, kg	Mean	1.15	1.08	1.19
	SD	0.17	0.24	0.23

Means with different superscript letters differ significantly: ^{a,b} $P < 0.05$, ^{A,B} $P < 0.01$. SD standard deviation

Table 8
Linear indicators of lambs' carcasses
Maße an der Schlachthälfte

Indicators	Statistical parameters	Groups		
		Control (n=10)	Experiment I (n=10)	Experiment II (n=10)
Carcass length 1, cm	Mean	76.00	77.20	79.10
	SD	3.08	2.04	1.40
Carcass length 2, cm	Mean	50.00	51.20	52.00
	SD	1.49	2.04	2.67
Carcass circumference 1, cm	Mean	60.20	61.40	62.60
	SD	1.69	3.10	0.52
Carcass circumference 2, cm	Mean	47.20	49.10	49.80
	SD	4.59	2.53	0.79
Ham length, cm	Mean	32.80	33.60	34.20
	SD	0.79	0.52	1.81
Ham circumference, cm (the widest part)	Mean	33.20	34.00	34.80
	SD	3.12	2.40	2.25
Ham index, %	Mean	99.64	99.24	98.55
	SD	10.17	6.84	7.07

Carcass length 1: *os pubis* - *atlas*, carcass length 2: *os pubis* - first rib, carcass circumference 1: by shoulder, carcass circumference 2: by ham, ham length: *tuber calcanei* - *tuberculum ossis ischii*

Table 9
Quality indicators of lambs' carcasses
Qualitätsmerkmale des Schlachtkörpers

Indicators	Statistical parameters	Groups		
		Control (n=10)	Experiment I (n=10)	Experiment II (n=10)
pH ₀	Mean	6.30	6.41	6.33
	SD	0.12	0.16	0.31
pH ₁	Mean	5.99	6.02	6.04
	SD	0.16	0.12	0.09
pH ₂	Mean	5.73	5.78	5.79
	SD	0.25	0.08	0.18
pH change 0 to 45 min	Mean	0.26	0.40	0.31
	SD	0.07	0.12	0.08
pH change 0 to 24 h	Mean	0.57	0.63	0.51
	SD	0.24	0.18	0.30
Water holding capacity, %	Mean	33.87	35.72	36.73
	SD	5.30	8.22	7.85
Conformation, 1-5	Mean	3.60	3.80	3.95
	SD	0.79	0.52	0.42
Fatness including kidney knob and channel fat, 1-5	Mean	3.20	3.55	3.60
	SD	0.42	0.52	0.67

pH₀ pH immediately after carcass dressing, pH₁ pH at 45 min *post mortem*, pH₂ pH at 24 h *post mortem*

Lamb meat quality indices (pH₀, pH₁, pH₂, water holding capacity and other) did not differ significantly depending on a treatment (Table 9). Chemical composition of the lamb meat did not show significant differences in terms of the feeding treatments (Table 10).

Table 10
Chemical content of lambs' meat
Chemische Zusammensetzung des Lämmerfleisches

Indicators	Statistical parameters	Groups		
		Control (n=10)	Experiment I (n=10)	Experiment II (n=10)
Dry matter	Mean	27.57	26.70	26.58
	SD	4.99	1.81	3.09
Crude proteins	Mean	18.29	19.09	18.55
	SD	1.70	1.63	1.76
Crude fat	Mean	3.11	3.48	3.65
	SD	0.38	1.56	1.21
Crude ash	Mean	1.00	0.99	1.01
	SD	0.08	0.04	0.05

Discussion

Chemical analysis of feed mixtures showed that forage heat treatment did not affect changes of their composition (Table 1). The results showed that heat treatments did not modify chemical composition, but significantly reduced the content of trypsin inhibition and urease activity, as well as decreased protein degradability (EWEEDAH *et al.* 1997).

RAMANZIN *et al.* (1991) obtained that roasted reduced the trypsin inhibitor content and urease activity of soybean, without modifying its chemical composition. Improvement of both daily gains and feed conversion indicates approval of heat treated forages application.

Lambs feeding with roasted cereals-raw soybean (E I) and roasted cereals-roasted soybean (E II) respectively affected fattening results (final body weight, daily gain), some slaughtering values, some biochemical values (urea, albumins, total cholesterol and HDL cholesterol) as well as enzyme activities (ALP, CK) which in the experimental groups (E I or E II) differed significantly or tended to compared to the control group.

The primary objective of most cereal grain-processing methods is to improve starch availability, which results in enhanced digestion and feed efficiency (GAEBE *et al.* 1998). Roasted corn resulted in greater body weight gains and better feed efficiency for finishing beef cattle than did raw corn (THOMAS *et al.* 1975). Lambs and bull-calves fed on heat-treated cereals brought about statistically significant higher daily gains, better feed conversion and reduced feed consumption compared to those fed on untreated ones (GRUBIĆ 1987). Occurrence of decreased consumption of the rations contained heat-treated cereals is probably the result of increased grain bulkiness as well as ration being composed in such a way. It is so because gelatinous starch is characterized by increased volume compared to the crude one (GRUBIĆ 1987). Heat treatment can successfully inactivate anti nutritional factors and also provides a means of protecting dietary proteins from rumen degradation. This result is more protein of dietary origin reaching the small intestine which may improve animal growth (BEEVER and THOMSON 1977, STERN *et al.* 1985). The amount and quality of crude protein reaching the small intestine is largely dependent on the roasting time and temperature used (YU *et al.* 2001a). In investigation ABDELGADIR *et al.* (1996) roasted corn enhanced calves performance when used with roasted soybean on 138°C. Results of this experiment demonstrated that interactions between carbohydrate and protein sources in calf starters are important and that synchrony of carbohydrate and protein ruminal availabilities may improve body weight gain and efficiency of utilization of calf starters. Namely, the harmful components inhibit digestive enzymes which caused disturbances of metabolism with animals fed full fat soybean, i.e. growth decrease, and even death of the animal (LIENER 1980). Apart from the above mentioned, crude soybean contains higher portion of highly unsaturated oils that can affect reduction of digestibility and energy absorption in ruminants (LOESCHE *et al.* 1992, SCOTT *et al.* 1991). Heat treatment has been used to decrease the rate of protein and (or) starch degradation in the rumen and increase the supply of dietary protein and (or) starch at the duodenum (TAMMINGA and JANSMAN 1993, GOELEMA 1999). Dry roasting of lupine and fabe beans tended to increase average daily gain and gain feed ration of lambs. Increased daily gains in lambs feeding with roasted lupin seeds and fabe beans were for 9-10 g as well as gain to feed ration for 8 g/kg DM (YU *et al.* 2001b). Daily body weight gains of lambs fed extruded lupine seeds were higher ($P < 0.05$) and better feed efficiency than those of lambs fed the untreated lupine-barley mixture (234 vs. 186 g) (PAJAK *et al.* 2001). Calves fed the diet containing soybeans roasted at 143°C consumed more feed, gained more body weight, and were more efficient in converting feed and energy than those fed soybeans roasted at 99 or 127°C (REDDY and MORRILL 1993).

However, YU *et al.* (2001a) not obtained significant differences in final live weight lambs, average daily gain and feed conversion efficiency between the raw and roasted whole narbon bean. Lambs feeding ratio with roasted whole narbon bean had higher daily gain for 9.0% and better feed conversion from lambs feeding with raw whole narbon bean. The inclusion of significant amounts of raw materials such maize, either as ensiled forage or as grain, in ruminant diets, however, has been shown to result in the flow of undegraded starch to the small intestine (SEAL and PARKER 2000). It can be assumed that, in this way, amount of the absorbed glucose in E I and E II groups in our trial was continuously slightly lower compared to the one from C group resulting in somewhat lower insulin level. The major control of insulin secretion is exerted by a feedback effect of plasma glucose, but orally administered glucose exerts a greater insulin-stimulating effect than intravenously administered glucose (GANONG 2001).

Conditions that increase insulin will increase cholesterol synthesis – conditions that decrease insulin or increase glucagon will decrease cholesterol synthesis. Transcriptional control of apolipoprotein A-I gene expression, the major apolipoprotein of HDL is dependent on insulin (MOORADIAN *et al.* 2004). Based on our results it can be concluded that total cholesterol drop (C:E I, $P<0.01$) was caused by the assumed lower insulin level in E I compared to C group. This occurred partially due to HDL cholesterol decrease (C:E I, $P<0.01$) composing greater part of the serum cholesterol in ruminants (DRACKLEY 2000) and because of decreasing other lipoprotein fractions. Values of tryacylglycerols are also in favour of the aforesaid.

Enzymatic starch decomposition in a small intestine is lower in ruminants due to insufficient pancreas amylase activity (WALKER and HARMON 1995) and approximately 45% of starch entering the intestine is not absorbed as glucose and in terms of overall ME yield, grain starch is best used when it is fermented in the rumen (HUNTINGTON 1997). Better provision with amino acids and energy in the group E I compared to group C significantly affected final body weight ($P<0.05$), daily gains ($P<0.05$) and proportions relative to slaughtering measures. All the above mentioned was obtained with lower daily food consumption, better conversion and significant ALP activity increase ($P<0.05$). The aforesaid results suggesting slight anabolic stimulation in E I compared to C are clearly visible in E II compared to C but not to E I. Changes suggesting intensive anabolic stimuli were caused by feeding lambs with roasted cereals and roasted soybean (E II). Started anabolic stimulation trend, due to better supply of amino acids and volatile fatty acids in E I, has been proceeded and more clearly pronounced in E II. This stimulus was caused by the soybean proteins heat treatment. Effect of heat treatment (toasting) on soybean protein degradability in the rumen was numerically increased duodenal flow of amino acid N (AAN) due to an increase in flow of undegraded feed AAN (LUND *et al.* 2004). There are a number of antinutritional factors present in soybean (protease inhibitors, phytohemagglutinin), among those factors, the major part are heat-labile (BAU *et al.* 2001).

The best fattening E II results could be attributed to feed effect on endocrine system and reaching maximum anabolic activity.

Conditions in which there are increased amounts of certain amino acids in the plasma increase growth hormone (GH) secretion (GANONG 2001). Circulating GH is a major

regulator of metabolism and growth during postnatal life and it regulates key metabolic pathways of intermediary metabolism (BREIER *et al.* 2000). GH separately increases the maximal capacity for protein accretion and the efficiency with which amino acids are used for protein accretion (BELL *et al.* 1998). This is mediated by insulin-like growth factor system (IGF-I ; IGF-II) (FLORINI *et al.* 1996). However, the above mentioned solid anabolic results can also be found in E I group. Differences detected between E I and E II could be attributed to diverse level of GH and IGH-1 being manifested as the trial results. IGF-1 level is considerably affected by GH which is completely dependent on nutritional factors. There are significant endocrine changes associated with malnutrition characterized by GH serum evaluation and IGF-1 decrease whereas obesity is associated with decreased GH and normal IGF-1 (STINE 1997). Our trial could be also characterized by IGF-1 level which was different between C, E I and E II affecting proportional significant increase of the values (feeding results, urea, albumins, ALP, CK, slaughtering results), i.e. ($P>0.05$) (glucose, total albumins) denoting anabolic processes. Significant level increase in total cholesterol, mainly due to increase of HDL cholesterol and HDL cholesterol between E I and E II could also be attributed to increase of GH i.e. IGF-1. Higher level of GH and IGF-1 increased glucose level stimulating insulin secretion (REIMERS 2003). Growth hormone replacement therapy decreased total cholesterol and LDL-cholesterol and increased significantly HDL-cholesterol ($P<0.05$) (TANRIVERDI *et al.* 2005). In this way insulin together with GH and IGF-1 significantly increased ($P<0.01$) HDL cholesterol level in E II compared to E I. Obtained differences in slaughter properties of lambs (trunks weight and dressing) indicate their better productive effects (Table 2 and 7). However, lambs meat quality (linear indices of lamb trunks and their quality indicators) did not significantly differ in terms of feeding treatment in spite of somewhat worse water fixing ability and higher meat water and fat percent in both experimental group lambs (Table 9 and 10). There were no significant differences in carcass measurements (hot carcass weight, cold carcass weight, dressing percentage and fat depth between 12th and 13th rib) between the raw and roasted whole narbon beans treatments YU *et al.* (2001a).

In conclusion, application of roasted cereals and soybean in feed mixtures led to better fattening traits and trunks quality in fattened lambs. Changes in lamb productive traits were followed by changes in some blood parameters composition (total-cholesterol, cholesterol HDL, urea, albumin and activity of ALP and CK). Concerning the gained production results may be concluded that the use of roasted cereals and soybean in feed mixture in lambs fattening is justified.

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