Carcass characteristics and meat quality attributes in lambs reared indoors, on cultivated pasture, or on semi-natural pasture

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This study evaluated the effects of different lamb production systems on live weight gain (LWG), carcass quality and meat quality. Four production systems for weaned intact male lambs were examined: indoor feeding with grass silage and concentrate (group 1), grazing on cultivated pasture with (group 2) or without (group 3) concentrate, and grazing on semi-natural pasture (group 4). Live weight, carcass weight, dressing percentage, carcass conformation, fatness and pH decline were recorded at slaughter, and M. longissimus thoracis et lumborum was analysed for colour, thawing and cooking loss, pH after 24 hours and 6 days, and Warner-Bratzler shear force. LWG was strongly affected by production system, being highest for group 1 and lowest for group 4 (p<0.001). Group 4 had the lowest conformation (p=0.002) and fat scores (p<0.001). Hence, production system affected age at slaughter, live weight gain, weight at slaughter, carcass conformation and fatness scores, but caused no differences in meat quality attributes in intact male lambs.

Key words: production system, intact male lamb, live weight gain, pH, colour, Warner-Bratzler shear force

Introduction

Domestic lamb and sheep meat production was in 2019 5.090 tonne, which corresponds to 30.7% of Sweden's total consumption in 2019 (Lannhard Öberg 2020). This suggests that there is potential for expansion of Swedish lamb meat production. Rising demand for high-quality Swedish lamb means that data are needed on the optimal way to rear lamb under Swedish conditions, so as to produce meat with high and consistent eating quality. It is well known that the eating quality and size of valuable cuts of Swedish lamb currently vary more than those of imported lamb and sheep (Carlsson and Arvidsson Segerkvist 2018). Several factors may explain this variation, including some related to primary production such as production system, choice of breed and/or cross, age at slaughter and carcass weight. The Swedish lamb meat production is characterized by few large and many small farms, with a large verity of breeds and production systems (Carlsson and Arvidsson Segerkvist 2018, Lannhard Öberg 2020). The average herd size was 33 ewes and/or rams in 2019 (Jordbruksverket 2019). It has also been shown that feedstuffs and feeding regime can also affect lamb meat quality (Watkins et al. 2013). In particular, pasture and concentrate may have different effects on meat flavour (Fisher et al. 2000, Arsenos et al. 2002, Priolo et al. 2002, Resconi et al. 2009). Feeding strategy can also influence glycogen storage in muscles, which in turn affects post-mortem muscle metabolism and thereby meat quality. Glycogen in muscles is converted into lactic acid under anaerobic conditions after slaughter, which reduces the pH of the muscle tissue. Since glycogen serves as ‘fuel’ in this process, it is essential to ensure that glycogen storage in the muscles prior to slaughter is sufficient to enable an adequate decline in pH (Bendall 1973). The official recommendation in Australia is for crossbred lambs to gain 100–150 g day⁻¹ in the last two weeks pre-slaughter, to ensure that the animals are in positive energy balance (growth phase) and have adequate glycogen depots in muscle (MSA 2015a). The effects of different feeding strategies on lamb meat quality can be evaluated by measuring the carcass pH, which is a useful indicator of various meat quality parameters. Specifically, the pH at 24 hours after slaughter (pH₂₄) is commonly used as an indicator of tenderness in meat (Geesink et al. 2000, Thompson et al. 2005, Toohey et al. 2006). Carcass pH (and thus meat quality) is sensitive to many factors pre-slaughter, during slaughter and post-slaughter, and can therefore vary between production systems (Sañudo et al. 1998). It is thus very important to understand how different production systems affect meat quality, in order to help producers deliver the consistent meat quality demanded by consumers.

The aim of this study was to determine how the four most commonly used lamb production systems in Sweden affect live weight gain (LWG), carcass quality and meat quality. The hypotheses tested were that: 1) higher feeding intensity improves growth rate and carcass and meat quality; and 2) concentrate allowance increases LWG, and thus carcass and meat quality.
Material and methods

Animals and experimental design

The experiment was performed between 29 June and 26 October 2016, at SLU Götala Beef and Lamb Research, Swedish University of Agricultural Sciences (SLU), Skara, Sweden (58°42′N, 13°21′E) and at a private farm outside Skara, Sweden (58°20′N, 13°26′E). In total, 80 crossbred weaned intact ram lambs (Dorset x Fine Wool) were included in the study, 36 of which were 50:50 crosses and 44 were 75:25 Dorset:Fine Wool crosses. The experiment was approved by the Ethics Committee on Animal Experiments, Gothenburg, Sweden (Registration No. 53-2016). Immediately prior to the study, the lambs were weighed and divided into four groups of 20 individuals each, equally balanced by breed crosses. All lambs were weaned just prior to the study. Average live weight (26.4±2.7, 26.8±2.8, 26.4±3.1 and 26.0±2.7 kg for group 1, 2, 3 and 4, respectively) and age (85.2±5.9, 84.4±5.3 and 84.4±6.0 days for group 1, 2, 3 and 4, respectively) were similar for the four groups at the start of the experiment. Each group was assigned a unique feeding treatment and followed that treatment throughout the experiment, with all groups starting treatment on 29 June. The treatments were: 1) indoor rearing; 2) cultivated pasture with a concentrate supplement daily; 3) only cultivated pasture; and 4) semi-natural pasture (Table 1). Group 4 was reared on the private farm, while the other three groups were reared at SLU Götala Beef and Lamb Research farm.

Table 1. Feeding strategies used for groups 1–4

<table>
<thead>
<tr>
<th>Group</th>
<th>Treatment</th>
</tr>
</thead>
<tbody>
<tr>
<td>Group 1, indoor</td>
<td>Silage ad libitum + 0.8 kg concentrate per lamb daily</td>
</tr>
</tbody>
</table>
| Group 2, pasture | Cultivated pasture + 0.3 kg concentrate per lamb daily  
* | Group 3, pasture | Cultivated pasture with no concentrate  |
| Group 4, pasture | Semi-natural pasture with no concentrate  |

*Group 2 received 0.4 kg of concentrate per lamb daily between 28 September and 4 October, due to poor pasture availability.

Experimental diets

Feed values and chemical composition of the experimental feeds are presented in Table 2. Group 1 was fed a diet consisting of silage *ad libitum* and 0.8 kg of a standard commercial concentrate (Fårfor Lamm 500, Lantmännen, Västerås, Sweden) per lamb per day to promote rapid growth. The seed mix for the silage consisted of 76% timothy (*Phleum pratense* L.), 18% red clover (*Trifolium pratense* L.) and 6% white clover (*T. repens* L.). The silage was harvested on 27–31 May, and was fertilised in late April with around 30 tonne ha\(^{-1}\) of cattle manure, providing 1.5 kg N tonne\(^{-1}\). A commercial additive (mixture of formic acid and propionic acid) was added to the herbage before ensiling. The animals in group 1 had free access to water, salt and minerals, and were housed indoors in an enclosed pen made from metal gates and with wheat straw bedding.

Groups 2 and 3 were kept on cultivated pasture in two different enclosures of 1.0 ha, both were divided into three grazing paddocks of 0.3 ha each. Both groups had access to one of the three paddocks at a time and they were moved once a week. Groups 2 and 3 differed in feeding intensity (Table 1), but both groups had daily access to a salt and mineral block and were given free access to water in a tub in each paddock. The seed mix for the cultivated pastures consisted of 50% timothy, 20% meadow fescue (*Festuca pratensis* Huds.), 15% perennial ryegrass (*Lolium perenne* L.), 10% red clover and 5% white clover. The cultivated pasture for groups 2 and 3 was fertilised on 3 April with 250 kg of Axan (Yara AB, Malmö, Sweden) (total nitrogen 27.0%, nitrate nitrogen 13.5%, magnesium 0.4%, sulphur 3.7% and calcium 6.0%) and then on 5 April with approximately 30 tonne ha\(^{-1}\) of cattle manure (1.5 kg N tonne\(^{-1}\)). The forage was harvested on 10 June. The regrowth was not fertilised after harvesting.

Group 4 was kept on unimproved semi-natural pasture (Table 2) with daily access to salt and minerals. The semi-natural pasture contained trees and shrubs and was on hilly land. This group was moved to new pastures on three occasions (31 August, 8 September, 11 October) due to poor quality and growth of the pasture late in the season. Lambs on semi-natural pasture had free access to water in a pond, stream or tub.

Due to illness, two lambs were removed from the experiment, one from group 1 and one from group 2. Data on these animals were excluded from further analyses, so the results are based on data for 78 lambs in total.
Special treatments during the experiment

All animals were dewormed (ivermectin, 0.8 mg ml\(^{-1}\)) at the start of the experiment. All animals other than those in group 1 were dewormed again after four weeks. Animals in group 4 were dewormed a third time, six weeks after the second treatment.

Feed sampling and analyses

Throughout the experiment, silage samples were collected daily and pasture samples were taken once a week and stored at \(-20^\circ C\) until analysis. Pasture samples were cut with a handheld machine, along a W-shaped route in each pasture according to Frame (1993). Before analysis, samples were pooled to obtain representative samples for the whole experimental period. The height of the cultivated pastures was measured each week in conjunction with weighing the lambs, immediately before the lambs were released into a new paddock. Sward height was measured according to Frame (1993), following the W-shaped route, with a rising plate meter (0.3 × 0.3 m, weight 430 g). The same sampling procedure was used for the semi-natural pasture, although the lambs in this group were not moved to a new pasture each week.

Crude protein was analysed according to Dumas (1831) and digestible protein levels were calculated using the digestibility coefficient of Spörndly (2003). Ash content was analysed by combustion at 525 °C. Dry matter (DM) measurement was performed by drying samples at 60 °C for 16 hours and then at 130 °C overnight. Neutral detergent fibre (NDF) was analysed as described by Chai and Udén (1998), using 100% neutral detergent fibre; AAT=amino acids absorbed in the duodenum; PBV=protein balance in the rumen; \(\text{ME}\)=metabolisable energy (MJ kg\(^{-1}\) DM) was determined by incubation in rumen fluid and buffer for 96 hours (Lindgren 1979) and then calculating the ME concentration based on in vitro disappearance of rumen organic matter, as described by Lindgren (1983).

Weighing and body condition scoring of the lambs

All lambs were weighed once a week on a portable scale (Iconix 21, Iconix New Zealand Ltd, New Zealand). Body condition scoring (BCS) was performed according to Swedish standards, using five condition classes ranging from 1 (very lean) to 5 (very fat) (Eggertsen 2007). The target for the lambs for slaughter was BCS 3 and live weight 47–50 kg. The lambs were divided into 10 slaughter groups, with 6–8 animals in each group.

Slaughter

For practical reasons, all animals were gathered and kept indoors on the farm on the night before slaughter. All had free access to water and silage until transport to a commercial abattoir, located about 10 minutes’ drive from...
All animals were transported in a horse trailer driven by staff from the university (SLU) to the abattoir at approximately 08:00 h. All procedures at the abattoir, such as lairage before slaughter, were varied as little as possible. The lambs were rendered unconscious by captive bolt stunning and then exsanguinated within 6 ± 2 seconds. Carcass weight (hot carcass weight × 0.98) and carcass grade were recorded. Conformation and carcass fatness were assessed manually by a certified classifier using the EUROP-scale, which has 15 classes ranging from 1 (poor conformation/very low fat) to 15 (very excellent conformation/very high fat). Dressing percentage was calculated as carcass weight/live weight × 100. Muscle pH was recorded, in the *M. longissimus* muscle, 24 hours after slaughter of each animal (Seven2Go pro, Metler Toledo, Schwerzenbach, Switzerland). Between test occasions, the pH probe was cleaned with pepsin solution to remove residual protein and with ethanol to remove residual fat, in accordance with the manufacturer’s instructions. The probe was then re-calibrated with pH 4.0 and 7.0 buffer solutions.

**Meat quality analyses**

All carcasses were hung by the Achilles tendon for six days at 4 °C and then the right *longissimus* muscle was collected from the first eight slaughtered animals in each group (in total 32 lambs). Immediately after sampling, these meat samples were vacuum-packed and kept frozen at –20 °C until analysis.

For each meat sample, the colour of the thawed meat, weight loss after thawing, weight loss after cooking and Warner-Bratzler shear force (WBSF) of the cooked meat were determined. The preparations required to measure all these physical variables meant that it was only possible to process four samples per day. Therefore, we chose to analyse one sample from each sample group on each day of analysis. An additional seven analysis sessions were performed for the remaining samples. All analyses were completed within two weeks. Before each analysis session, four meat samples (one each from groups 1–4) were removed from the freezer, unpacked, weighed (start weight), repacked in a new vacuum bag and thawed for 15 hours at 4 °C. The samples were then tempered in a 20 °C water bath for one hour, unpacked and reweighed to get the weight loss after thawing. After removal of the fat cover, colour measurements were performed on the surface of the longissimus muscle, at eight locations, using DigiEye (VeriVide, Enderby, UK) and mean of the eight measurements was calculated. For WBSF measurements, repacked samples were placed in a water bath pre-heated to 75 °C for one hour or until the core temperature reached 70 °C. The samples were then allowed to cool in ice water for one hour, after which the meat was unpacked, weighed to get cooking loss and then placed in bags to reach room temperature. Cylindrical samples with diameter 15 mm (7–10 replicates/meat sample) were punched out in the longitudinal direction of the fibres for measurement of WBSF using an Instron 5542 instrument (Instron Ltd., High Wycombe, UK). A total of 7–10 measurements were performed and the mean value was calculated. Each sample was placed in a wedge-shaped recess under the cutting blade. The blade was 1 mm thick and moved downwards through a rectangular hole at a speed of 50 mm min\(^{-1}\). The maximum force measured was used as a measure of the cutting resistance of the sample.

**Statistical analysis**

Statistical analyses were performed using the Mixed procedure in SAS (SAS 9.4, SAS Inst. Inc., Cary, NC, USA). Two statistical models of the following forms were created, with production system (with four sub-classes) included as a fixed effect.

Live weight gain, age at slaughter and carcass characteristics were analysed using the model:

\[ Y_{ij} = \mu + P_i + e_{ij} \]

Thawing and cooking loss, colour and WBSF were analysed using a model which included day of analysis as a random effect:

\[ Y_{ij} = \mu + P_i + d_{ij} + e_{ik} \]

where \( Y_{ij} \) is the dependent variable, \( \mu \) is the grand mean, \( P \) is the fixed effect of the production system, \( d \) is the random effect day of analysis, and \( e \) and \( e_{ik} \) are the residual error (\(-N(0, \sigma^2))\). A general Satterthwaite approximation for the denominator degrees of freedom was performed, using the SATTERTH option in SAS.

Differences were considered significant at \( p<0.05 \) and indicative of tendencies at 0.05≤\( p<0.10 \).
Results

Feeding intensity and live weight gain

The amount of pasture available to group 4 (5.6 cm on average over the experimental period) was less than that for the other pasture groups (9.5 cm for group 2, 9.2 cm for group 3). Production system had an effect \((p<0.001)\) on LWG, which followed the intensity of the feeding treatments, i.e. group 1 lambs had the highest LWG, followed by group 2, group 3 and group 4, respectively (Table 3). As expected, LWG affected the age at slaughter, which differed between all four groups \((p<0.001)\). Group 4 lambs had the highest age at slaughter. Live weight at slaughter also differed between the groups \((p<0.011)\). Group 1 had the highest live weight at slaughter, while group 2 had higher live weight at slaughter than group 3 (Table 3). There were between-group differences in LWG during the last 14 days pre-slaughter. In group 4, growth rate per day in the last 14 days prior to slaughter exceeded the overall growth rate during the experiment.

Carcass quality

Production system had a significant effect on carcass weight \((p<0.001)\), with group 4 having lower carcass weight than the other three groups. Group 4 had lower conformation \((p<0.001)\), fat score \((p=0.039)\) and dressing percentage \((p<0.001)\) than all other groups (Table 4).

### Table 3. Live weight (LW), age and live weight gain (LWG) of lambs reared using four different production systems

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Group 1</th>
<th>Group 2</th>
<th>Group 3</th>
<th>Group 4</th>
<th>SEM</th>
<th>Significance</th>
</tr>
</thead>
<tbody>
<tr>
<td>n</td>
<td>19</td>
<td>19</td>
<td>20</td>
<td>20</td>
<td></td>
<td></td>
</tr>
<tr>
<td>LW at start, kg</td>
<td>26.4</td>
<td>26.8</td>
<td>26.4</td>
<td>26.0</td>
<td>0.65</td>
<td>NS</td>
</tr>
<tr>
<td>LW at slaughter, kg</td>
<td>50.6(^a)</td>
<td>50.3(^ab)</td>
<td>48.3(^c)</td>
<td>48.9(^d)</td>
<td>0.54</td>
<td>0.011</td>
</tr>
<tr>
<td>Days in experiment</td>
<td>65(^a)</td>
<td>82(^a)</td>
<td>91(^b)</td>
<td>109(^d)</td>
<td>2.7</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Age at slaughter</td>
<td>149(^d)</td>
<td>167(^c)</td>
<td>177(^b)</td>
<td>194(^a)</td>
<td>2.9</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>LWG, g day(^{-1})</td>
<td>377(^a)</td>
<td>287(^b)</td>
<td>244(^c)</td>
<td>211(^d)</td>
<td>7.9</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>LWG 14d(^{-1}), g day(^{-1})</td>
<td>322(^a)</td>
<td>287(^b)</td>
<td>244(^c)</td>
<td>319(^d)</td>
<td>0.0</td>
<td>0.043</td>
</tr>
</tbody>
</table>

### Table 4. Carcass quality and meat quality of lambs reared using four different production systems

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Group 1</th>
<th>Group 2</th>
<th>Group 3</th>
<th>Group 4</th>
<th>SEM</th>
<th>Significance</th>
</tr>
</thead>
<tbody>
<tr>
<td>Carcass quality</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>n</td>
<td>19</td>
<td>19</td>
<td>20</td>
<td>20</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Carcass weight</td>
<td>21.6(^a)</td>
<td>21.3(^a)</td>
<td>20.9(^a)</td>
<td>18.8(^c)</td>
<td>0.47</td>
<td>0.003</td>
</tr>
<tr>
<td>Conformation(^1)</td>
<td>9.2(^a)</td>
<td>8.7(^a)</td>
<td>8.7(^a)</td>
<td>7.9(^a)</td>
<td>0.24</td>
<td>0.002</td>
</tr>
<tr>
<td>Fatness(^2)</td>
<td>7.4(^a)</td>
<td>7.7(^a)</td>
<td>7.4(^a)</td>
<td>6.5(^a)</td>
<td>0.17</td>
<td>&lt; 0.001</td>
</tr>
<tr>
<td>Dressing, %</td>
<td>42(^a)</td>
<td>42(^a)</td>
<td>41(^a)</td>
<td>37(^a)</td>
<td>0.4</td>
<td>&lt; 0.001</td>
</tr>
<tr>
<td>(pH_{54})</td>
<td>5.83</td>
<td>5.66</td>
<td>5.77</td>
<td>5.59</td>
<td>0.098</td>
<td>NS</td>
</tr>
<tr>
<td>Meat quality</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>n</td>
<td>8</td>
<td>8</td>
<td>8</td>
<td>8</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Age at slaughter, d</td>
<td>146(^a)</td>
<td>163(^a)</td>
<td>172(^a)</td>
<td>193(^a)</td>
<td>3.5</td>
<td>&lt; 0.001</td>
</tr>
<tr>
<td>(pH_{54})</td>
<td>5.45</td>
<td>5.40</td>
<td>5.45</td>
<td>5.41</td>
<td>0.032</td>
<td>NS</td>
</tr>
<tr>
<td>Thawing loss, %</td>
<td>4.3</td>
<td>4.8</td>
<td>4.1</td>
<td>5.0</td>
<td>0.4</td>
<td>NS</td>
</tr>
<tr>
<td>Cooking loss, %</td>
<td>24.3</td>
<td>24.1</td>
<td>22.8</td>
<td>24.2</td>
<td>0.87</td>
<td>NS</td>
</tr>
<tr>
<td>Colour L*</td>
<td>37.1</td>
<td>37.2</td>
<td>36.5</td>
<td>35.7</td>
<td>0.69</td>
<td>NS</td>
</tr>
<tr>
<td>Colour a*</td>
<td>16.9</td>
<td>16.5</td>
<td>16.6</td>
<td>16.4</td>
<td>0.28</td>
<td>NS</td>
</tr>
<tr>
<td>Colour b*</td>
<td>7.5</td>
<td>7.0</td>
<td>6.8</td>
<td>7.1</td>
<td>0.31</td>
<td>NS</td>
</tr>
<tr>
<td>WBSF, N(cm(^2))(^{-1})</td>
<td>33.8</td>
<td>45.9</td>
<td>31.9</td>
<td>34.9</td>
<td>4.88</td>
<td>NS</td>
</tr>
</tbody>
</table>

Group 1 is reared by indoor feeding with silage and concentrate; group 2 on cultivated pasture with 0.3 kg concentrate per lamb daily; group 3 on cultivated pasture and group 4 on semi-natural pasture. SEM = standard error of the mean; NS = non-significant \((p>0.05)\); \(^{a-b}\) = Mean values within rows with different superscripts differ significantly \((p<0.05)\); \(^{1}\) = According the EUROP system, where 1=P-, 2=P, 3=P+, 4=O-, 5=O, 6=O+; 7=R-, 8=R, 9=U-, 11=U, 12=U+, 13=E, 14=E and 15=E+; \(^{2}\) = According the EUROP system, where 1=1-, 2=1, 3=1+, 4=2-, 5=2, 6=2+, 7=3-, 8=3, 9=3+, 10=4-, 11=4, 12=4+, 13=5, 14=5 and 15=5+; NS = non-significant \((p>0.05)\); WBSF = Warner-Bratzler shear force.
Meat quality indicators and technological meat quality attributes

There were no differences between groups with respect to the technological meat quality attributes, i.e. pH after 24 hours, pH\textsubscript{6d} (six days after slaughter), thawing loss, cooking loss, colour (L*, a* and b*) and WBSF (Table 4).

Discussion

In this study, higher feeding intensity resulted in higher LWG, with group 1 lambs having the highest LWG, but there was no effect on meat quality attributes. Live weight gain and/or live weight at slaughter differed from those in comparable studies (e.g., Young et al. 1994, Pethick et al. 2005, Campbell et al. 2012).

The LWG values were mostly higher than in previous studies, where some of the treatments described even gave negative LWG (Young et al. 1994, Pethick et al. 2005, Campbell et al. 2012). However, the nutrient content of the feeds used in those studies is only briefly described, so it is difficult to compare the results with those in the present study. The energy content of the cultivated pasture for group 3 was lower than that for group 2, even though both groups grazed the same field (albeit different parts). This difference in energy content was most likely due to differences in the establishment and relative abundance of various plant species, i.e. the sward composition in the pasture, influencing the nutritional content in different areas of the pasture. Although the nutrient content in the semi-natural pasture was good (Table 2), it should be noted that the amount of pasture available for group 4, based on sward height, was less than that for the other pasture groups (groups 2 and 3). This observation is important for understanding the comparatively low growth rate of group 4, which was most likely adversely affected by pasture availability, as the nutritional content of the pasture was high and should have resulted in higher growth. However, the LWG in group 4 lambs (211 g day\textsuperscript{-1}) was similar to that in a study by Lind et al. (2009), who observed LWG of 230 g day\textsuperscript{-1} on semi-natural pastureland in northern Norway. Thus, it can be concluded that rearing lambs on semi-natural pasture in the temperate climate zone is feasible. The LWG\textsubscript{6d} for group 4 was higher than the average LWG for the whole rearing period. The reason for this remains unclear. To increase the weight at slaughter for animals reared on semi-natural pastures (group 4), the lambs could: i) be given supplementary feed or ii) lambing time could be brought forward in the spring, in order to enable earlier release to pasture and ensure that the lambs have time to grow and reach slaughter maturity before the nutritional quality and availability of semi-natural pastures decreases in the autumn.

As expected, the high-intensity feeding systems (groups 1 and 2) and the intermediate production system (group 3) all yielded carcasses with higher conformation and fatness scores than the extensive feeding system (group 4). However, although group 4 lambs had the lowest scores for both conformation and fatness, the carcasses would still qualify for the highest payment per kg carcass weight according to the current price list at two Swedish abattoirs (HKScan 2020, KLS 2020). The dressing percentage was lower for group 4 (37%) compared with the other groups (41–42%). This could be explained by the higher age at slaughter for group 4 lambs, which is associated with greater loss of weight in terms of head, bones and intestines (Muir et al. 2008). This loss of weight probably did not derive from greater rumen fill at slaughter, since the amount of pasture available to lambs reared under semi-natural conditions was limited. As expected, feeding intensity affected both growth per day and number of days to slaughter. Although group 3 lambs were reared at a lower intensity than those in groups 1 and 2, they were ready for slaughter at around the same time as group 1 and 2 lambs, despite being raised solely on cultivated pasture. Overall, there were no appreciable differences between groups 1, 2 and 3 with respect to carcass conformation and fatness scores.

In meat from groups 1 and 3, the pH\textsubscript{6d} value exceeded 5.7 which is considered as the upper limit for a good eating quality (MSA 2015b). This outcome was unexpected, since high growth rates are associated with high glycogen storage in muscles, which normally causes the expected post-slaughter pH decline. However, our results are consistent with those of Pethick et al. (2005), who found that meat from animals fed a high-energy diet had higher pH\textsubscript{6d} than meat from animals fed low-energy diets. That study also found that the high-energy group unexpectedly lost a greater proportion of glycogen between farm and slaughter than lambs raised on pasture. Pethick et al. (2005) suggested that these animals may have been more predisposed to lose glycogen in response to stress, a trait which could be of metabolic or behavioural origin. In the present study, even though groups 1 and 3 meat had high pH\textsubscript{6d} values, there was no expected effect of high pH on the tenderness of the meat (WBSF values). Other studies have found that lower pH\textsubscript{6d} values are associated with more tender meat, e.g. Devine et al. (1993) found that meat is most tender when its final pH is 5.5–5.7. Moreover, two of the pH\textsubscript{6d} values observed in the present study were above the upper limit of 5.7 recommended by Meat Standards Australia (MSA 2015b). Based on this standard,
there was a risk of meat from groups 1 and 3 having lower eating quality than that from the other groups. Commercial research in New Zealand has established that the desirable pH<sub>24</sub> range for lamb is 5.4–5.8, with values of 5.8–6.0 being associated with intermediate quality (Alliance Group Ltd 2010). Based on these criteria, groups 2, 3 and 4 had more numerically preferable pH<sub>24</sub> values than group 1.

In the literature, the pH at 24 hours post mortem is often defined as “final” pH (e.g. Koohmaraie et al. 1991, Koohmaraie et al. 1995, Watanabe et al. 1996, McGeehin et al. 2001, Díaz et al. 2002, Priolo et al. 2002, Sañudo et al. 2003, Velasco et al. 2004, Pethick et al. 2005, Teixeira et al. 2005, Majdoub-Mathlouthi et al. 2013, Majdoub-Mathlouthi et al. 2015). However, in the present study a continuous decline in pH was seen between pH<sub>24</sub> and pH<sub>72</sub> (Table 4). In contrast, Koohmaraie et al. (1995) found that the pH of lamb carcasses 24 hours post mortem (pH<sub>24</sub> = 5.6) was identical to that six days later, and that carcass pH values then rose slightly (to 5.7) between seven and 21 days post mortem. It is thus not clear whether pH generally drops significantly after 24 hours or not. However, the results presented both in this study and in Koohmaraie et al. (1995), suggest that describing the pH at 24 hours post mortem as “final” or “ultimate” may be inaccurate.

Technological meat quality parameters determined in this work can only be compared with those in previous work to a limited extent. Comparisons between studies are hampered by differences in e.g. feeding intensity, LWG, breed, sex, intact or castrated ram lambs, carcass weight and slaughter method, or the fact that some of these factors are barely described. These differences derive from the many different production systems used for lamb, which have different prerequisites such as climate and tradition. Thus, previous studies have found no differences in WBSF (Berge et al. 2003, Rodrígues et al. 2008, Karaca et al. 2016), significant differences in WBSF (Sañudo et al. 2003), no differences in meat colour (Díaz et al. 2002, Pethick et al. 2005), significant differences in meat colour (Priolo et al. 2002) and no differences in water loss (Díaz et al. 2002, Rodrígues et al. 2008, Karaca et al. 2016) when comparing different lamb types and production systems.

Berge et al. (2003) did not find any differences in WBSF (2.17–3.69 kg, equal to 21.3–36.2 N) between indoor-reared entire male lambs fed concentrate compared with entire male lambs reared on different types of pasture, with or without concentrate. The results in Berge et al. (2003) are not fully comparable with ours, due to differences in animal age at slaughter (3.5–7 months), carcass weight (10.4–19.7 kg) and feeding systems, but were similar to those in our study. Karaca et al. (2016) studied the effect of two feeding systems where lambs were fed a finishing diet of either alfalfa hay (1750 g day<sup>−1</sup>) or alfalfa hay (1250 g day<sup>−1</sup>) + 500 g of barley per lamb and day, with both diets balanced to give equal energy intake in the two groups. Live weight gain for both groups (90 and 32 g day<sup>−1</sup>) was low compared with that in the present study, but these values were significantly different. The WBSF values did not differ between groups in that study (45.7 and 43.9 N), as also found in the present study. However, carcass weights reported by Karaca et al. (2016) (17.5 and 15.7 kg) were lower than those in the present study. Significant differences in average daily gain between groups were observed in both that and the present study, indicating that differences in growth rate do not automatically result in differences in WBSF. Further, Karaca et al. (2016) found no differences in either meat colour (L<sup>*</sup>, a<sup>*</sup> or b<sup>*</sup>) or cooking loss, corresponding to the results of the present study. A study by Rodrigues et al. (2008) found differences in LWG when comparing different feeding intensities (straw + pelleted commercial concentrate compared with whole barley grain + protein supplement), with a LWG per day of 272 g for the straw group and 371 g for the barley group. On the other hand, Rodrigues et al. (2008) did not find any differences in meat colour, WBSF or water-holding capacity. However, carcass weight was lower (by 12 kg) than in the present study, hampering direct comparison of the results.

Sañudo et al. (2003) compared different production systems from six European countries, based on either grass or concentrate or a combination of both. The WBSF results in that study revealed significant differences between systems related to different feeding strategies, age at slaughter, carcass weight and breeds. However, the results are not unequivocal, indicating a combined effect of the factors mentioned above on WBSF. Nevertheless, the results indicate that WBSF values are affected by age at slaughter when comparing entire male lambs with similar carcass weights. The youngest animals had the lowest WBSF values in the study by Sañudo et al. (2003), but no such effect was seen in the present study. The lack of differences in technological meat quality attributes in the present study is a positive finding, since it indicates that all four production systems compared can be used in practice without altering the technological properties of the lamb meat. Pethick et al. (2005) found significant differences in LWG when comparing different feeding intensities (pasture, moderate-energy pellet, high-energy pellet, straw). They found no differences in L<sup>*</sup> and b<sup>*</sup>, as in the present study, but observed differences in a<sup>*</sup>, with meat from lambs in the pasture and moderate-energy treatment having a darker red colour than the high-energy pellet group, in contrast to the present study. Pethick et al. (2005) attributed the differences in a<sup>*</sup> to elevated ultimate pH levels in the high-energy pellet (pH 5.66) and straw (pH 5.67) feeding treatments compared with pasture.
(pH 5.57) and moderate-energy pellet (pH 5.59). This could also explain the lack of differences in colour in the present study, since pH did not differ significantly between groups and may therefore not have influenced any of the colours (L*, a* or b*). As in the present study, Diaz et al. (2002) did not observe any differences in colour (L*, a* or b*) or water-holding capacity when comparing concentrate and pasture for fattening lambs. However, Priolo et al. (2002) recorded differences in L*, with a grass-fed group producing darker meat than the stall-fed group, and saw a tendency for differences in b*, with the grass-fed group having a lower yellowness index than a stall-fed group. Priolo et al. (2002) attributed the difference in L* to numerically higher ultimate pH for the grass-fed group (pH 5.62) compared with the stall-fed group (pH 5.57). However, this difference was non-significant, and it is thus questionable whether it can explain the difference in L* between grass-fed and stall-fed lambs.

Conclusions

There were differences in LWG between the four production systems studied here, but the results indicate that intact ram lambs can be reared under intensive or extensive conditions without any differences in meat quality attributes. Parameters affected by the production system included age at slaughter, live weight gain, carcass weight, carcass conformation and fatness scores. Lambs reared on cultivated pasture had better carcass classification (conformation and fatness), as well as carcass weights than those reared on semi-natural pasture. Further studies are needed to evaluate the existing recommendations of pH to be 5.7 for meat to identify if this recommendation applies for animal material and production systems used in this study. Studies are also needed on whether pH at 24 hours post mortem should be described as “final” or “ultimate” pH, since further pH decline in muscle beyond 24 hours has been observed. If pH measurements are to be carried out later than 24 hours after slaughter, possible correlations between pH and meat quality attributes, such as tenderness, should be examined.

Acknowledgement

We thank Jonas Dahl, David Johansson, Karin Wallin and Frida Dahlström for valuable technical support, the staff at Skara lammslakteri for help during slaughter, and Mr Lennart Pettersson, farmer, for good cooperation. We are also grateful to the funding bodies Stiftelsen Svensk Fårforskning, Interreg ÖKS [grant no. 20200994], Västra Götalandsregionen [grant no. RUN-610-0789-13], Agroväst and the Swedish University of Agricultural Sciences for base support.

References


