

# Effect of fat tail docking on meat quality of Awassi sheep in comparison with Lacaune sheep

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## 1 ABSTRACT

This experiment was conducted to study the effect of fat tail docking on Awassi lamb sheep's meat in comparison with thin-tail Lacaune sheep's meat (imported breed in Lebanon). Twenty seven ram lambs were divided into three groups, intact fat-tail Awassi (IA), docked fat-tail Awassi (DA), and imported thin-tail Lacaune (IL). Docking was performed at one week of age, by applying tight rubber rings on the tail. Animals having the same body weight were slaughtered at one year of age. Three samples of muscles were excised from each carcass; *Biceps Femoris (Bf)*, *Longissimus Dorsii (Ld)* and *Gluteus Medius (Gm)*. Samples were wrapped in an oxygen permeable film and stored at 4°C and -30°C for the assessment of pH, and meat quality traits. The ultimate pH was significantly higher ( $P<0.01$ ) in Lacaune breed as compared to both docked and intact Awassi breed. DA animals had significantly higher ( $P<0.01$ ) fat content in *Ld* muscle than IA and IL lambs. Both Awassi groups presented higher  $L^*$  values than Lacaune animals ( $P<0.05$ ). In addition for Awassi breed,  $L^*$  values of *Bf* and *Gm* muscles were significantly greater ( $P<0.05$ ) than that of *Ld* muscle.  $b^*$  values were significantly greater ( $P<0.05$ ) in *Gm* of all animals under study. Lacaune group had lower drip loss values than Awassi groups ( $P<0.05$  in *Bf* and *Gm* muscles and  $P<0.01$  in *Ld* muscle). However for Awassi breed, the drip loss in *Ld* muscle of DA animals was significantly lower ( $P<0.01$ ) than IA ones. Furthermore, thawing and cooking losses were neither affected by breed nor by docking. Regarding the PND values, the *Ld* muscle of DA group presented higher values ( $P<0.05$ ) than *Bf* and *Gm* muscles of both intact groups. In addition, cooked meat PND values were significantly higher ( $P<0.05$ ) in *Ld* muscle of DA when compared to intact groups. Due to the scarcity of information in this area of research, further investigations are needed.

## 2 INTRODUCTION

Awassi is the most widespread sheep breed of non-European origin (Galal *et al.*, 2008). They are large framed, fat tailed sheep mostly grown and consumed in the Middle East countries (Abdelqader *et al.*, 2017). The fat tail plays an important role in adaptation of sheep raised under the harsh feeding conditions of arid and semi-arid regions, where the availability of

food-stuff is seasonal (Vatankah and Talebi, 2008). Awassi sheep are known to deposit 20 % of their carcass weight as fat in the tail which has traditionally been used as source of cooking fat (Yousefi *et al.*, 2012). Awassi sheep have desirable carcass traits and meat quality (Holloway *et al.*, 1994), especially because the presence of the fat tail which allows for leaner

carcasses and makes it easier to trim any undesirable fat. The fat stored in the tail may contribute to the lower level of fat found elsewhere in various cuts compared to more traditional sheep breeds (Safdarian *et al.*, 2008). Fat tailed lambs store fat in their tail and they tend to have lower amount of carcass and muscular fat (Webb and O'Neill, 2008). Awassi lamb's meat had lower dry matter and crude fat percentages than other genotypes (Abdullah *et al.*, 2011). Despite their high demand in the Mediterranean and Arabian Gulf region yet they produce a very small proportion of lamb and mutton. The self-sufficiency of highly preferred locally produced lambs in the area of fat-tailed sheep ranges from 10 to 12 % (Nik-Khah, 1984). This deficit obliges the market to import thin-tail lamb from Europe and Australia. Moreover, the market price of fat-

tailed sheep meat is two to three times higher than that of thin-tailed sheep (FAO, 2015). Many researchers *e.g.* Abi Saab *et al.* (2010) have studied the effect of fat tail on the reproductive characteristics of Awassi breed. The objectives of docking fat tail in lambs are to improve reproductive skills, feed efficiency, live body weight gain and to facilitate natural mating with non-fat-tailed breeds (Farah *et al.*, 2019). To the best of our knowledge, experimental data on the effect of fat tail docking on meat quality in ovine and especially in Awassi sheep are scarce. Furthermore, fat tail docking is thought to be desirable for increasing the organoleptic properties of lamb's meat (Moharrery, 2007). Therefore, the present experiment was carried out on Awassi and Lacaune sheep to study the effect of fat tail docking on meat quality and compare it with thin-tail sheep.

### 3 MATERIALS AND METHODS

**3.1 Animals:** A total of 27 ram lambs were raised at the Center of Research and Agriculture Formation (CRAF) in Ghazir, till one year of age (2018-2019) under conventional breeding conditions and nutrition. Males were divided into 3 equal groups; 2 groups of fat tail Awassi sheep (intact "IA" and docked "DA") and a group of imported thin-tail Lacaune sheep (IL). Docking was performed at one week of age, by applying tight

rubber ring on the tail at the distal end of the caudal fold (between the first and the second vertebrae down the tail). Animals were withdrawn for 24 hours, transported for 1.5 h and allowed to rest for 1 h before being slaughtered according to standard commercial procedure (captive bolt stunning, no electrical stimulation). Table 1 summarizes some characteristics of the animals within groups at slaughter.

**Table 1:** Summary of animals<sup>1</sup> at slaughter.

Breeds	Number of lambs	Fasting (h)	Transportation (h)	Live body weight (Kg)	Carcass weight (kg)
<b>Awassi</b>					
IA	9	24	1.5	51.54± 3.74	22.76± 2.17
DA	9			52.32± 4.01	23.0± 2.54
<b>Lacaune</b>					
IL	9	24	1.5	53.71± 3.99	23.15± 2.33

<sup>1</sup> IA, intact Awassi lambs; DA, docked Awassi lambs; IL, intact Lacaune lambs.

**3.2 Muscle sampling:** After 30 minutes of dressing, the right femoral muscle (*Biceps femoris* "Bf"), the loin muscle (*Longissimus dorsi* "Ld") and the sirloin muscle (*Gluteus medius* "Gm") were excised from the carcass of each animal. Two similar size cuts of Ld and Gm muscles were packed and stored at 4°C and -30°C, respectively. Whereas, 2 slices of approximately 300 g each with similar size and shape were cut from the right femoral

muscle; one slice was stored at 4°C, while the other one was vacuum-packed, frozen and conserved at -30°C to determine the meat quality after freezing/defrosting.

**3.3 Ultimate pH measurements:** At 2 different *post mortem* times (24 and 48 hours) and after cooking, the ultimate pH of each muscle was determined as described by Jeacocke (1977). Approximately, 2 g of muscle were homogenized in

18 ml of 5 mM iodoacetate buffer. The pH of the homogenate was measured using a pH meter (WTW 720 pH meter) equipped with a combined electrode (Sen Tix 21) and an average of 2 replications by sample was calculated.

### 3.4 Meat quality

**3.4.1 Fat:** Petroleum ether extraction method was performed to determine the meat fat content as described by the Association for Official and Analytical Chemists (AOAC, 1995). Approximately, 5 grams of muscle were grinded until obtaining a homogeneous mass, and put in a filter paper then in a drying oven (SELECTA 295057; from 0 to 250°C) at 105°C for 60 minutes. Dried samples were then cooled for 15 minutes in a desiccator (KARTTEL) and weighed. Later on, filter papers were transferred into a Soxhlet equipment and placed in the extractor flask containing the petroleum ether solvent. After heating and cooling, the solvent trickles dissolve the fats and within 10 to 12 hours the fat extraction finishes. Total fat were expressed in gram/ per 100 g of fresh tissue.

**3.4.2 Colour:** Meat color was determined using a chromameter (ADCI-60-C; Beijing, China). The instrument was set to measure using the CIE system values of luminance ( $L^*$ ), redness ( $a^*$ ), and yellowness ( $b^*$ ) using illuminate D and 65° standard observer (CIE, 1986). All measurements (3 replicates on each sample) were carried out at 2 different *post mortem* times (24 and 48 hours) and after cooking on the surface of each sample, in an area free of obvious color defects (over scalding, blood spots, and haemorrhages).

**3.4.3 Water holding capacity (WHC):** Drip, thawing and cooking losses were determined as described by Honikel (1998). Cuts of *Bf*, *Ld* and *Gm* were weighed at the time of collection (approximately 30 minutes *post mortem*) placed in a polystyrene tray, wrapped in an oxygen permeable film and kept at 4°C. Meat cuts were reweighed at 48 hours *post mortem* and the drip loss was expressed

as percentage of initial weight. After 12 h thawing at 4°C, cuts from *Bf*, *Ld* and *Gm* muscles were taken from bags, dried with filter paper, and reweighed before cooking. Thawing loss was expressed as a percentage of the frozen weight. Cooking loss was determined immediately after thawing; cuts (from *Ld* and *Gm* muscles) were vacuum-packed in polyethylene bags and cooked in a water bath at 80°C for 15 minutes corresponding to an internal temperature of 70°C. Samples were cooled for 45 minutes under running tap water at room temperature. After that, they were taken from the bags, dried with filter paper and weighed. Cooking loss was expressed as the percentage loss relative to the weight immediately before cooking.

**3.4.4 Texture:** Texture was measured using a penetrometer (interface RS232C) with a needle of 2.5g based on a weight of 97.5 g, thus attaining a total weight of 100 g. The measurements were performed at different *post mortem* times (3, 12, 24, 48, 72, 144 hours and after cooking). The penetration was carried out on meat slices (3cm x 2cm x 1cm) prepared such that the longest dimension was parallel to the fiber axis. Slices were placed on a horizontal support and the force of the needle was applied perpendicularly to the muscle fibers for 5 seconds (Becila, 2002). The penetrometer needle depth (PND; in mm) was recorded and an average of 3 replications by sample was calculated.

**3.5 Statistical analysis:** ANOVA was carried out using the GLM procedure of SPSS 16.0. The model included the effects of breed, docking and muscle type. When significant effects were recorded, mean values were compared using Duncan's multiple range tests. All results were reported as mean  $\pm$  standard deviation ( $\mu \pm Sd$ ). In graphs, vertical bars indicate standard deviation (Sd).

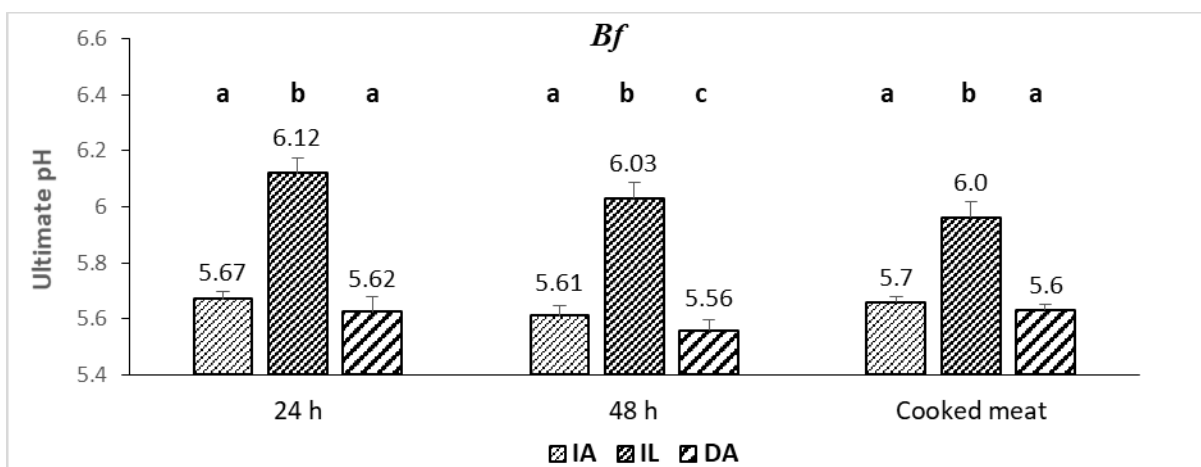
## 4 RESULTS AND DISCUSSION

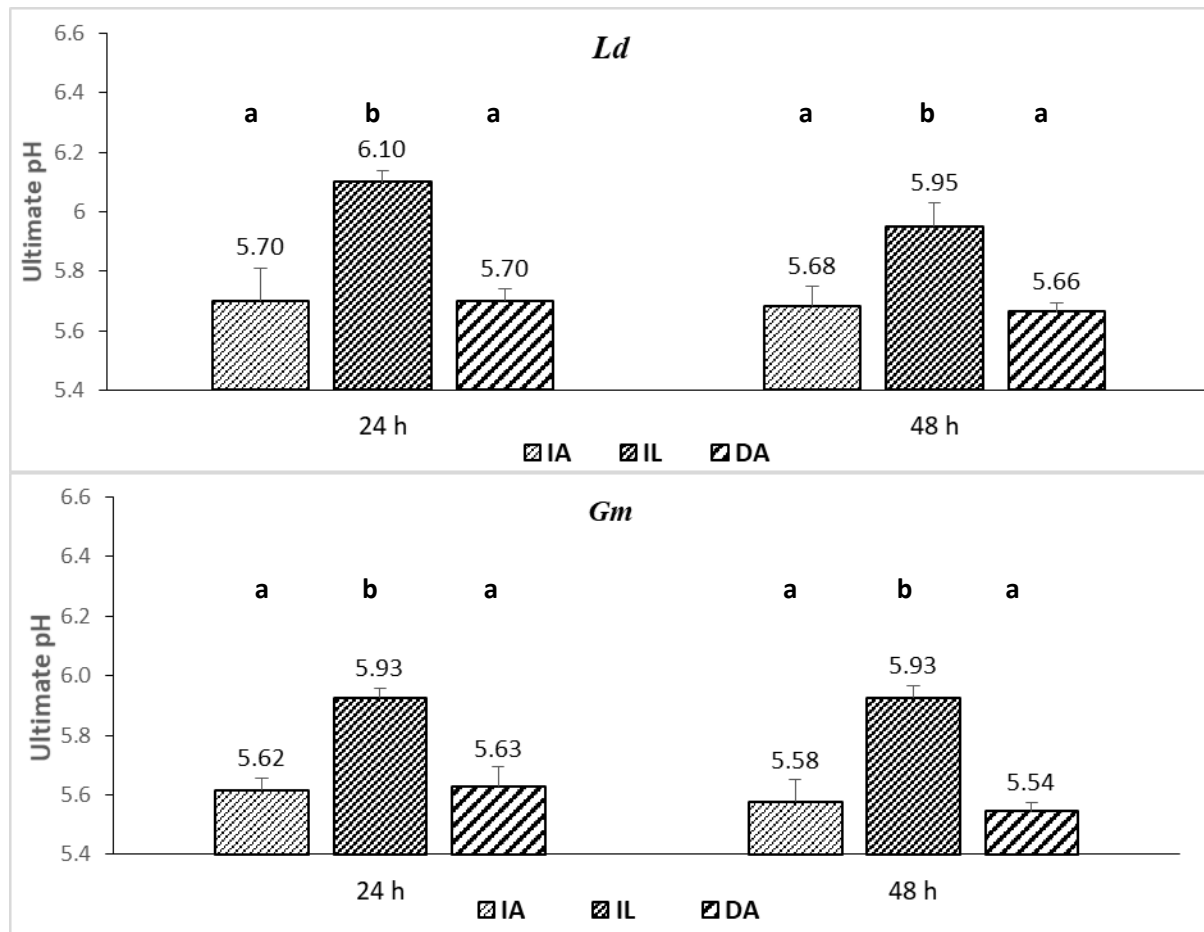
**4.1 Muscle ultimate pH:** As shown in figure 1, the ultimate pH of all muscles in IL animals, was significantly higher than in IA and DA sheep ( $P < 0.01$ ). In fact, at 48 h *post mortem* the *Bf* muscle of DA was significantly the lowest (DA:  $5.56 \pm 0.04$  vs. IA:  $5.61 \pm 0.04$  and IL:  $6.03 \pm 0.06$ ,  $P < 0.01$ ), in addition and after cooking, both Awassi groups (Intact and docked) had lower *Bf* muscle's pH then Lacaune breed ( $5.96 \pm 0.05$  vs.  $5.66 \pm 0.11$  and  $5.63 \pm$

$0.06$ ;  $P < 0.01$ ). Moreover, at 48 h *post mortem* *Ld* muscle's pH of docked lambs was significantly lower than Lacaune, and slightly lower than IA ( $5.66 \pm 0.03$  vs.  $5.95 \pm 0.08$  and  $5.68 \pm 0.07$ ;  $P = 0.07$ ). Finally at 48h *post mortem* for the *Gm* muscle's pH, the values of IL breed were significantly higher than Awassi both docked and intact groups ( $5.93 \pm 0.04$  vs.  $5.58 \pm 0.07$  and  $5.54 \pm 0.03$ ;  $P < 0.05$ ). In conclusion, for the present study the mean ultimate

pH in all muscles of Lacaune Breed was above 5.8 and values ranged between 5.60 and 6.79, whereas, Awassi groups (IA and DA) presented lower ultimate pH values varying between 5.51 and 5.77. Results of the present research suggested that ultimate pH of meat in Awassi sheep is not affected by fat tail docking. Those results are in agreement with Wang *et al.* (2018) who stated that tail docking does not affect the ultimate pH (measured at 24 h *post mortem*) of *Ld* and *Gm* muscles in sheep. However Awassi lambs (IA or DA) have a significant lower ultimate pH (at 24 and 48 h *post mortem*) in *Bf*, *Ld* and *Gm* muscles than Lacaune breed, thus affected by breed. These variations in the ultimate pH of different breeds could be explained by the fact that sheep of various genetic types had different *peri mortem* glycogen content. Indeed, as muscle is converted into meat, the stored glycogen is anaerobically degraded to lactic acid and the extent of *post mortem* pH decline is determined by the accumulation of lactate and consequently the content of muscle glycogen at slaughter. In agreement with this assertion, the high mean ultimate pH value detected in the present study in Lacaune breed meat could be attributed to the low glycogen muscle concentration in IL animals as compared to Awassi breed. Those results agreed with Hoffman *et al.* (2003) and Jandasek *et al.* (2014) who confirmed the effect of breed on the ultimate pH of meat; they reclaim that some sheep breeds are less resistant to pre-slaughter stresses than others leading to different rates and extents of *post mortem* pH fall. Similarly, while studying meat quality in different lamb breeds, Martinez- Cerezo *et al.* (2005) found pH differences. In contrast those results don't agree with, Hernandez- Cruz *et al.*

(2009) who reported that the ultimate pH measured at 24 h *post mortem* is not affected by ovine breeds. Furthermore Hopkins *et al.* (2011) assumed that values of red meat ultimate pH greater than 5.8 affect negatively the bacteriological stability, the shelf life, the flavor and the aroma of meat. Since the ultimate pH of Awassi and Lacaune sheep meat were, respectively, lower and higher than 5.8 (ranging between 5.51 and 5.77 for Awassi breed and 5.60 and 6.79 for Lacaune sheep), it could be suggested that Awassi sheep meat has a longer shelf life and a better flavor and aroma than Lacaune sheep. Finally this experiment result's agree with Esenbuga *et al.* (2009) and Abdullah *et al.* (2011), who reported that after the cessation of blood circulation and oxygen supply, lactic acid accumulates in *Ld* muscle of fat tailed sheep resulting in a pH drop from 6.33 to 5.64 at 24 hours *post mortem*. Concerning the effect of muscle type on ultimate pH in lamb's meat (Tab.2.), the values differed significantly among the 3 muscle types ( $P < 0.05$ ). In Awassi lambs (IA and DA), the *Ld* muscle showed the highest ultimate pH while, in Lacaune animals, the ultimate pH was the lowest in the *Gm* muscle. These findings are consistent with the work of Ablikim *et al.* (2016) who noticed significant differences in the ultimate pH of *Bf*, *Ld* and *Gm* muscles in different sheep breeds. These variations in muscle pH may result from differences in the proportion of red (STO: Slow Twitch Oxidative) and white (FTOG: Fast Twitch Oxidative Glycolytic; FTG: Fast Twitch Glycolytic) muscle fibers which had different contractile and metabolic properties and different glycogen content and ATP stores.





**Figure. 1:** Effect of breed and fat tail docking on ultimate pH (measured at 24 and 48 hours *post mortem* and after cooking) in *Bf*, *Ld*, and *Gm* muscles of sheep lambs. <sup>a, b, c</sup> different letters indicate significant difference at  $P < 0.01$ .

**Table. 2:** Effect of muscle type<sup>1</sup> on ultimate pH (measured at 24 and 48 h *post mortem*) in Awassi (IA), tail docked Awassi (DA) and Lacaune (IL) sheep.

	IA				DA				IL			
	<i>Bf</i>	<i>Ld</i>	<i>Gm</i>	<i>S</i>	<i>Bf</i>	<i>Ld</i>	<i>Gm</i>	<i>S</i>	<i>Bf</i>	<i>Ld</i>	<i>Gm</i>	<i>S</i>
24h	5.67± 0.03 a	5.74± 0.11 b	5.62± 0.04 a	**	5.62± 0.05 a	5.70± 0.04 b	5.63± 0.07 a	**	6.12± 0.06 a	6.08± 0.04 a	5.79± 0.03 b	**
48h	5.61± 0.04 a	5.68± 0.07 b	5.58± 0.07 a	**	5.56± 0.04 a	5.66± 0.03 b	5.54± 0.03 a	***	6.03± 0.06 a	5.95± 0.08 ab	5.93± 0.04 b	*

<sup>1</sup>. *Bf*, *Ld*, *Gm*, *Biceps femoris*, *Longissimus dorsii*, *Gluteus medius*;

<sup>a, b</sup>, within the same row, indicate significant differences at  $P < 0.05^*$ ,  $P < 0.01^{**}$ , and  $P < 0.001^{***}$ ; NS,  $P > 0.05$ .

## 4.2 Meat quality attributes

**4.2.1 Fat:** The present work (Tab. 3.) showed that the breed did not affect the meat fat concentration in sheep. Whereas, tail docking influenced the deposition of fat in Awassi sheep muscles. Moreover, in *Ld* muscle, DA animals had a significant higher fat content ( $P < 0.01$ ) than IA and IL groups ( $5.18 \pm 0.15$  Vs.  $3.76 \pm 0.16$  and  $4.08 \pm 0.5$  g/ 100 g, respectively;  $P < 0.01$ ). However, no significant effect of docking on muscle fat

deposition was detected in *Bf* and *Gm* muscle. That is to say, docking affected significantly fat content in the *Ld* muscles mainly. Findings of the current work coincided with the observations of Kyanzad (2001) who published that the loss of fat tail in sheep is compensated by an increase in sub-cutaneous muscle and internal fat percentages. Moreover, Safdarian *et al.* (2008) reported that fat-tailed sheep generally had lower proportions of carcass and muscle fat depots than docked lambs. Indeed Webb

and O'Neill (2008), confirmed that tail docking increases the carcass fat accumulation including skeletal muscle. Likewise, Wang *et al.* (2018) observed that rump and epiploic fat weights were increased in docked lambs as were *Ld* and *Gm* muscles fat comparing to undocked sheep. Besides, when evaluating the effect of muscle type on fat content, this trial detected that, in IA and IL groups, *Gm* muscle showed the highest fat concentration, whereas *Ld* the lowest and *Bf* muscle an

intermediate value, ( $P < 0.001$ ). Whereas, in DA lambs, the *Gm* muscle exhibited the highest fat content ( $8.06 \pm 0.73$  g/ 100 g) followed by *Bf* muscle ( $5.82 \pm 0.08$ ) and after the *Ld* ( $5.18 \pm 0.15$ ) with no significant difference between *Bf* and *Ld* muscles. These results were consistent with the findings of Abdullah *et al.* (2011) who noticed that meat fat content was affected by muscle types and that Awassi sheep were considered to have the lowest *Ld* muscle fat % in comparison with other breeds.

**Table. 3:** Fat content in *Bf*, *Ld* and *Gm* muscles<sup>1</sup> of intact and tail docked Awassi (IA and DA, respectively) and Lacaune (IL) lambs sheep.

	Fat (g/ 100 g)			S
	<i>Bf</i>	<i>Ld</i>	<i>Gm</i>	
<b>Awassi</b>				
IA	5.96 ± 0.84 <sup>x</sup>	3.76 ± 0.16 <sup>a<sup>y</sup></sup>	8.19 ± 0.29 <sup>z</sup>	***
DA	5.82 ± 0.08 <sup>x</sup>	5.18 ± 0.15 <sup>b<sup>x</sup></sup>	8.06 ± 0.73 <sup>y</sup>	*
<b>Lacaune</b>				
IL	6.13 ± 0.05 <sup>x</sup>	4.08 ± 0.50 <sup>a<sup>y</sup></sup>	8.60 ± 1.28 <sup>z</sup>	***
<b>S</b>	NS	**	NS	

<sup>1</sup> *Bf*, *Ld*, *Gm*, *Biceps femoris*, *Longissimus dorsi*, *Gluteus medius*; <sup>a</sup>, <sup>b</sup>, and <sup>x</sup>, <sup>y</sup>, <sup>z</sup>, within the same column and row, respectively, indicate significant differences at  $P < 0.05^*$ ,  $P < 0.01^{**}$ , and  $P < 0.001^{***}$ .

**4.2.2 Colour:** The present experiment showed that the breed affected significantly the color measurements (CIE: L\*, a\*, b\*) of raw meat in sheep ( $P < 0.05$ ). However, no significant influence of docking was detected on L\*, a\* and b\* values (Tab. 4). For a clearer presentation, the trichromatic coordinates (CIE values) measured on cooked meat are not shown in the table, since these parameters were not affected by treatments under study. Concerning L\* values at all *post mortem* time, only breed (not docking) affected luminosity. Indeed Awassi lambs (docked and intact) presented higher values in all studied muscles than IL animals ( $P < 0.05$ ). Muscle effect on luminosity showed that in Awassi sheep (IA and DA), luminosities of *Bf* and *Gm* muscles measured at all *post mortem* time were significantly greater ( $P < 0.05$ ) than that of *Ld* muscle, with no significant difference recorded in Lacaune breed. Concerning the redness of meat, neither breed nor docking or type of muscle influenced a\* values. Only a trend toward muscle effect was recorded in Lacaune breed, where *Ld* muscle tended to have higher a\* values than *Bf* and *Gm* muscles ( $P = 0.08$ ). Regarding yellowness, b\* values were significantly greater ( $P < 0.05$ ) in *Gm* of all animals under study, as a result of muscle effect. In conclusion, this study confirmed that L\* is affected by both breed and muscle type, but not by

docking; whereas, a\* is neither affected by breed nor by docking, only muscle type had an effect on redness of Lacaune's meat; finally, yellowness is only affected by muscle type. Those findings don't agree with Esenbuga *et al.* (2009) who reported that lamb sheep genotypes did not affect the L\* and a\* values of meat. In contrast, our results agreed with Abdullah *et al.* (2011) who noted a significant difference among lamb breeds in lightness and redness of meat (L\* and a\* values). Some authors admitted that the luminance of meat (L\*) indicates the superficial light scattering and depends not only on *post mortem* pH fall but on redness and yellowness of meat as well, which are mainly considered as indicators of myoglobin content and oxidation (Santé, 1993). In agreement with this statement, the present experiment showed that sheep breeds affected significantly the luminance of meat ( $P < 0.05$ ) and that higher L\* values were associated with lower ultimate pH (see Tab. 4. and Fig. 1). Although the current study showed that lightness was numerically higher in *Ld* muscle of DA lambs than that of IA animals (Tab. 4.) which agree with Wang *et al.* (2018) who declared that docked lambs presented higher L\* values in *Ld* and *Gm* muscles than intact sheep with an insignificant effect of docking on redness (a\*) and yellowness (b\*) values. However, the relationship between luminosity and

fat content in *Ld* muscle in docked animals could be due to the fact that, in this work, the high *Ld* ultimate pH of docked lambs compensated the positive effect of high fat content on meat lightness. Finally, it is noteworthy that  $b^*$  values were significantly higher ( $P < 0.05$ ) in *Gm* muscle than *Bf* and *Ld* muscles of all lambs under study (see Tab. 4.). This result was consistent with the finding that *Gm* muscle had greater fat content than *Bf* and *Ld* muscles (see Tab. 3). We suggest then i) that not only the lightness of meat is affected by the intramuscular fat but also the yellowness and ii) that intramuscular fat is not the only reason for differences found in the lightness of sheep meat; several factors seem to play an important role e.g. ultimate pH. Due to the scarcity of information in this area of research, more investigations are needed to clarify the effect of fat tail docking sheep on muscular fat accumulation and consequently lightness of meat by determining the intensity of relationship between fat concentration, ultimate pH and CIE color readings ( $L^*$ ,  $a^*$ ,  $b^*$ ).

**4.2.3 Water holding capacity:** Experimental results obtained on water holding capacity of meat within breed, fat tail docking animals and muscle type are presented in table 5. Current results showed that breed affected significantly the drip loss and animals of IL group had lower values than Awassi sheep (IA and DA groups), ( $P < 0.05$  in *Bf* and *Gm* muscles and  $P < 0.01$  in *Ld* muscle). These results were consistent with the fact that high ultimate pH induced low drip losses and consequently high water holding capacity (WHC). Moreover, Lacaune breed (IL) presented the highest ultimate pH in the three studied muscles. Indeed, as pH decreases and approaching the iso- electrical point (pH= 5.5) of proteins, the repulsion forces between myofibrils decrease and thick and thin

filaments move closer together pushing the cell water out. Thus, fluid is lost from muscle fibers leading to a reduction in meat WHC. The current study showed that docking influenced the WHC of Awassi sheep meat. The drip loss in *Ld* muscle of DA animals was significantly lower than that of IA group ( $1.42 \pm 0.03$  vs.  $1.49 \pm 0.03$  %;  $P < 0.01$ ). It is very likely that the positive effect of docking on the WHC was related to the high fat content and exacerbated by the high ultimate pH measured in the *Ld* muscle of DA lambs. Thawing and cooking losses in *Bf*, *Ld* and *Gm* muscles of lambs were neither affected by breed nor by tail docking. Nevertheless, a trend toward a significant effect of docking on thawing loss was recorded in *Ld* muscle; thawing loss tend to be lower in *Ld* muscle of DA animals than that of IA and IL groups ( $P = 0.06$ ). Those findings agree with El Rammouz *et al.* (2004) who recorded a significant correlation between drip loss and ultimate pH ( $r = -0.56$ ;  $P < 0.01$ ) in turkey toms. Moreover this experiment result's agreed with Wang *et al.* (2018) who reported that drip loss of lamb meat is closely related to muscle fat concentration and confirmed that drip losses were decreased in *Ld* and *Gm* muscles by 2.3 and 9.9 %, respectively, in docked sheep, finally he confirmed that neither the thawing loss nor the cooking loss were affected by docking. Some authors stated that fat concentration is indeed a potential contributor to the WHC of raw meat by surrounding the muscle fibers and the water, the deterioration of fat by the presser and the heat when thawing and cooking (Lawrie and Leward, 2006). Regarding cooking loss, Hoffman *et al.* (2003) reported similar results and detected insignificant difference between sheep breeds. In contrast, Abdullah *et al.* (2011) published a significant effect of breed on cooking loss in ovine meat.



**Table. 4:** Meat color ( $L^*$ ,  $a^*$ ,  $b^*$ ; measured at 24 and 48 h *post mortem*) in *Bf*, *Ld* and *Gm* muscles<sup>1</sup> of intact and tail docked Awassi (IA and DA, respectively) and Lacaune (IL) lambs sheep.

	$L^*$				$a^*$				$b^*$			
	Awassi		Lacaune		Awassi		Lacaune		Awassi		Lacaune	
	IA	DA	IL	S	IA	DA	IL	S	IA	DA	IL	S
<b>24 h p.m.</b>												
<i>Bf</i>	42.22± 3.01a*	43.04± 3.88a*	38.21± 3.22 <sup>y</sup>	*	19.51± 3.35	19.45± 4.95	18.90± 4.11	NS	10.36± 2.80a	9.68± 3.01a	10.55± 3.18a	NS
<i>Ld</i>	38.96± 2.41b*	39.52± 1.44b*	36.31± 2.59 <sup>y</sup>	*	20.75± 3.91	18.92± 4.16	20.79± 2.70	NS	12.01± 1.45a	11.95± 2.81a	11.84± 2.81a	NS
<i>Gm</i>	42.24± 3.11a*	43.25± 3.36a*	36.34± 2.26 <sup>y</sup>	*	18.87± 3.47	17.80± 1.71	18.19± 3.71	NS	14.71± 2.09b	14.56± 1.79b	14.37± 1.50b	NS
<b>S</b>	*	*	NS		NS	NS	$P=0.08$		*	*	*	
<b>48 h p.m.</b>												
<i>Bf</i>	44.31± 4.00a*	44.30± 3.45a*	39.65± 2.00 <sup>y</sup>	*	17.19± 2.38	18.20± 3.38	18.30± 2.57	NS	10.38± 2.81a	11.10± 2.66a	11.65± 2.62a	NS
<i>Ld</i>	39.96± 1.41b*	40.21± 2.10b*	37.48± 2.41 <sup>y</sup>	*	18.89± 1.49	18.85± 3.46	18.98± 3.11	NS	12.49± 2.40ab	12.02± 1.74a	12.00± 1.76a	NS
<i>Gm</i>	42.12± 0.87a*	43.61± 2.79a*	39.15± 3.39 <sup>y</sup>	*	18.43± 1.79	18.63± 1.96	18.60± 1.78	NS	14.11± 1.64b	14.98± 2.40b	14.45± 2.36b	NS
<b>S</b>	*	*	NS		NS	NS	NS		*	*	*	

<sup>1</sup> *Bf*, *Ld*, *Gm*, *Biceps femoris*, *Longissimus dorsi*, *Gluteus medius*; a, b, and <sup>x, y</sup> within the same column and row, respectively, indicate significant differences at  $P < 0.05$ ; NS, not significant.

**Table. 5:** Effect of breed and fat tail docking on water holding capacity (WHC; drip, thawing and cooking losses) in *Bf*, *Ld* and *Gm* muscles<sup>1</sup> of sheep lambs.

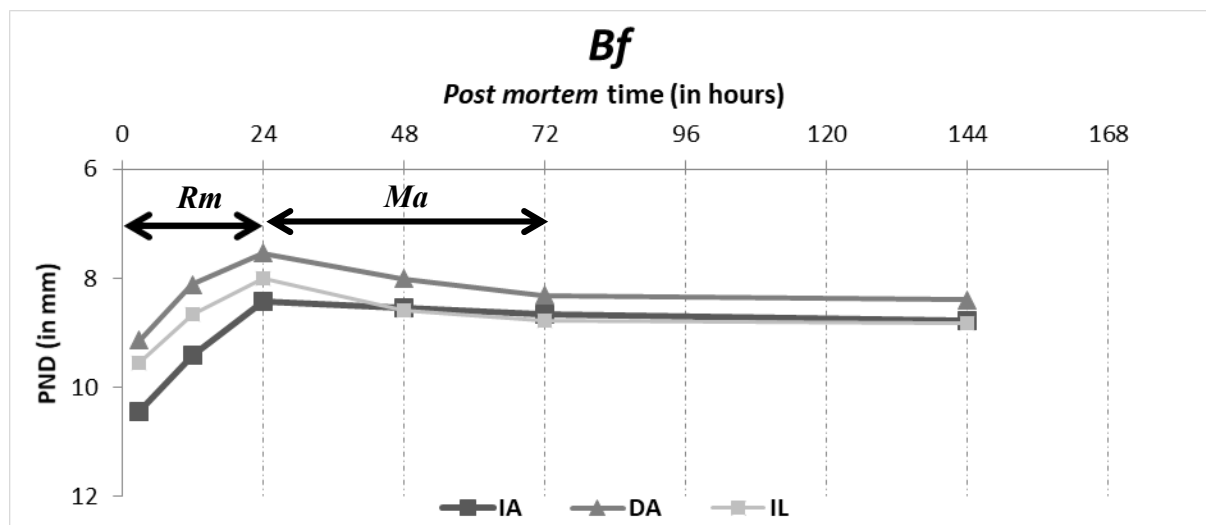
	Drip loss (%; 48 h p.m.)				Thawing loss (%)				Cooking loss (%)			
	Awassi		Lacaune		Awassi		Lacaune		Awassi		Lacaune	
	IA	DA	IL	S	IA	DA	IL	S	IA	DA	IL	S
<i>Bf</i>	1.55± 0.04a*	1.53± 0.03a*	1.43± 0.02a <sup>y</sup>	*	4.81± 0.13	4.74± 0.18	4.73± 0.12	NS	27.21± 0.63	27.44± 0.41	27.00± 0.74	NS
<i>Ld</i>	1.49± 0.03b*	1.42± 0.03b <sup>y</sup>	1.42± 0.03a <sup>z</sup>	**	4.82± 0.12	4.72± 0.11	4.81± 0.14	$P=0.06$	27.48± 0.60	27.23± 0.66	27.31± 0.79	NS
<i>Gm</i>	1.53± 0.03a*	1.52± 0.03a*	1.46± 0.02b <sup>y</sup>	*	4.78± 0.15	4.77± 0.12	4.77± 0.15	NS	27.33± 0.71	27.12± 0.61	26.88± 0.53	NS
<b>S</b>	*	*	*		NS	NS	NS		NS	NS	NS	

<sup>1</sup> *Bf*, *Ld*, *Gm*, *Biceps femoris*, *Longissimus dorsi*, *Gluteus medius*; a, b, and <sup>x, y, z</sup> within the same column and row, respectively, indicate significant differences at  $P < 0.05$ ; NS, not significant.



**4.2.4 Texture:** The *post mortem* evolution of meat tenderness in *Bf*, *Ld* and *Gm* muscles of intact, docked Awassi and Lacaune sheep (Fig.2.) showed that the time of *Rigor mortis* establishment in Awassi and Lacaune sheep is 24 hours and that meat ageing begins at 24 h and achieves at about 72 h *post mortem*. In addition, the optimum sheep meat tenderness (Awassi and Lacaune sheep) occurred at 72 h *post mortem* when stored at 4°C. However, the tenderness of raw meat in *Bf*, *Ld* and *Gm* muscles measured at different *post mortem* times was not affected by breed. Moreover, docking affected meat tenderness of Awassi sheep. Indeed, at all *post mortem* times, the *Ld* muscle of DA group presented higher PND values than IA and IL lambs ( $P < 0.05$ ). Concerning the cooked meat, the current study recorded a positive effect of docking on *Ld* muscle's tenderness (DA:  $5.33 \pm 0.58$  v.s. IA:  $4.48 \pm 0.66$  and IL:  $4.39 \pm 0.53$  mm;  $P < 0.05$ ), (Fig. 3). Findings of this experiment agrees with different author's results. When anaerobic glycolysis ceased, the pH fall stops as well as the producing of adenosine triphosphate (ATP); actin and myosin form rigid chain inducing an inextensible state of muscle and the establishment of *Rigor mortis* (Bate-Smith and

Bendall, 1947). After that, meat ageing occurs (Abdullah *et al.*, 2011) which is associated with *post mortem* proteolysis degradation by endogenous enzymes (Prado and De Felicio, 2010). Moreover, Cetin *et al.* (2012) reported that the muscle pH fall stopped and the *Rigor* achieved at 24 h *post mortem* followed by the tenderization of muscle (meat ageing) which lied between 24 and 72 h *post mortem* in lambs. However results of this study were slightly different than those reported by Burke and Apple (2007) who noted a significant difference between sheep breeds in terms of instrumental meat tenderness. However, concerning docking effect, this experiment agreed with Wang *et al.* (2018) who reported that the shear force of *Ld* muscle, in relation with fat level, was significantly reduced in docked lambs compared to intact animals (reduction of 12.15 %;  $P < 0.01$ ). In accordance with our results and those of Wang *et al.* (2018), tenderness and drip loss are significantly correlated with the meat fat content. Moreover, Hopkins *et al.* (2006) mentioned that muscle fat concentration is an important indicator in meat quality related to smoothness and has a great influence on freshness and tenderness.



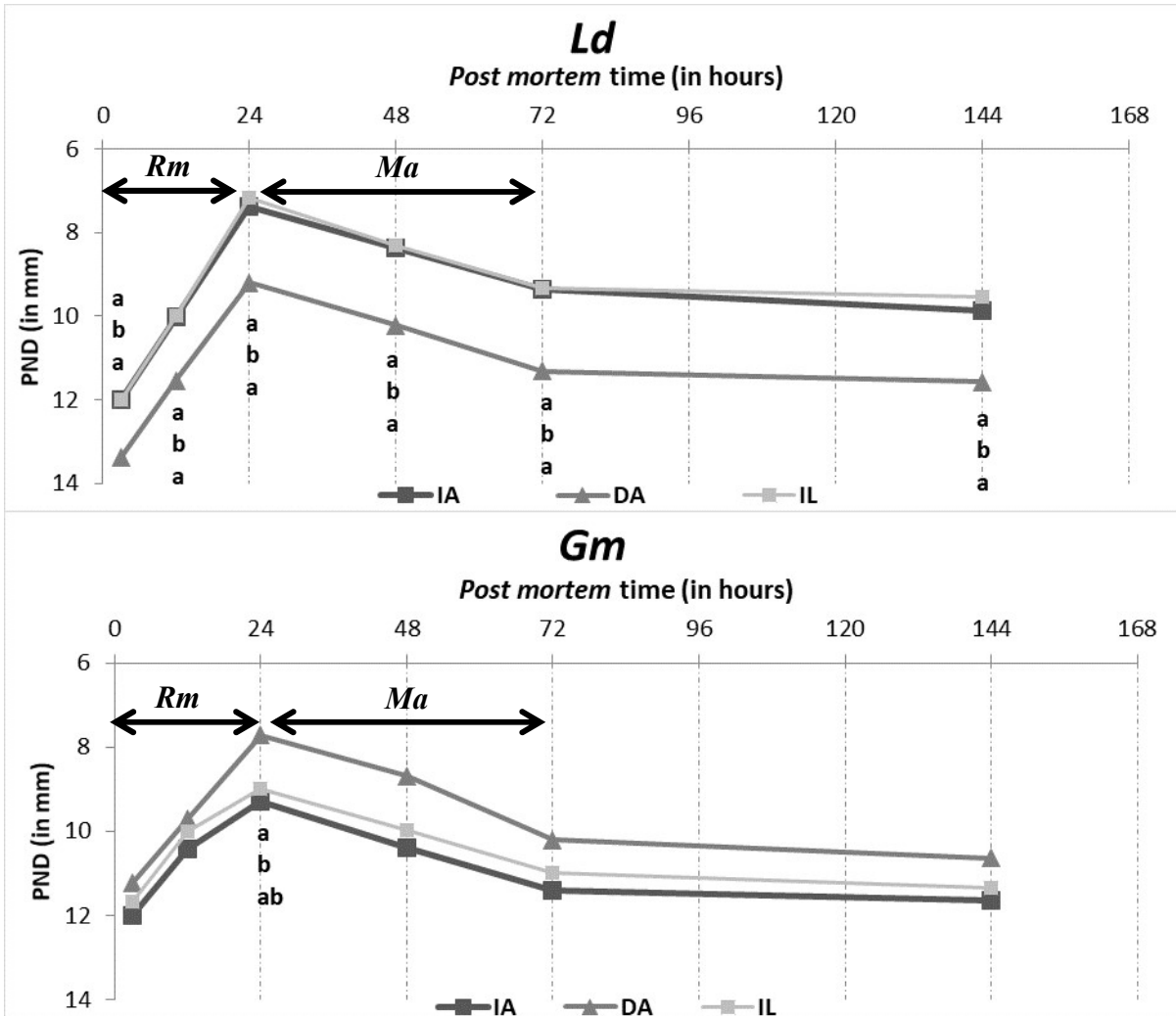


Figure 2: Post mortem evolution of meat tenderness in *Bf*, *Ld* and *Gm* muscles of intact (IA), fat tail docking Awassi (DA) and Lacaune (IL) sheep. <sup>a, b</sup> different letters indicate significant difference at  $P < 0.05$ . PND, penetrometer needle depth (mm); *Rm*, Rigor mortis; *Ma*, Meat ageing.

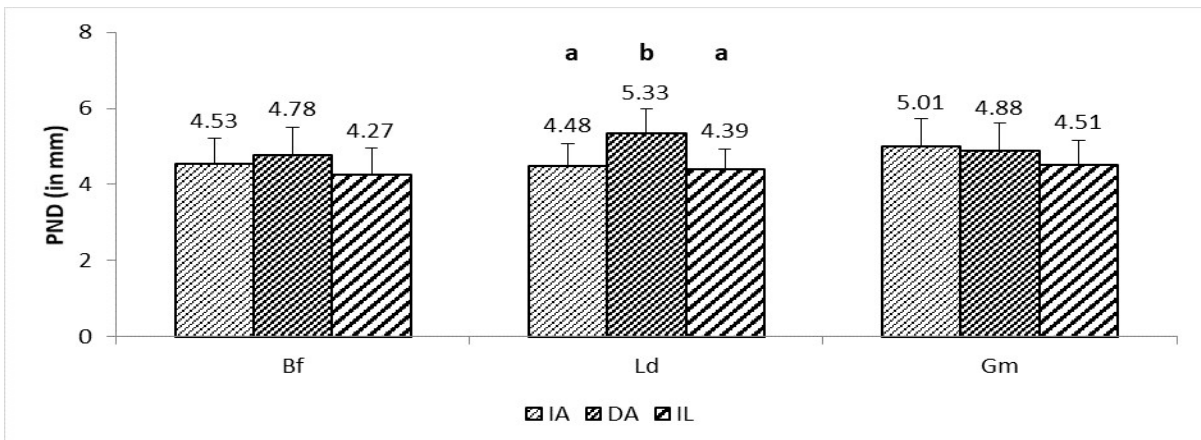


Figure 3: Effect of breed and fat tailed docked on PND (Penetrometer needle depth; mm) of cooked meat in *Bf*, *Ld* and *Gm* muscles of intact (IA), fat tail docking Awassi (DA) and Lacaune (IL) sheep. <sup>a, b</sup> different letters indicate significant difference at  $P < 0.05$ .

## 5 CONCLUSION

Results of the current experiment showed that lamb's breed affected meat quality. It seems that Awassi sheep meat has a longer shelf life and a better flavor and aroma than Lacaune lambs. Awassi tail docking enhances the accumulation of fat in *Ld* muscle and induces better WHC and tenderness.

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