Effects of using siris (Albizia lebbeck) foliage in the diet of fattening lambs on nutrient digestibility, blood and rumen parameters, growth performance, and meat quality characteristics

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Abstract
This study aimed to investigate the effect of substitution of siris foliage with alfalfa forage in the diet of fattening lambs on digestibility, fermentation, and growth performance of fattening lambs. In the present experiment, 27 8-month-old Arabi lambs (31.3 ± 6) with an initial weight of 28.8 ± 1.99 kg were used in a completely randomized design. The effect of experimental diets on dry matter intake was not significant; however, the diets had a significant effect on the intake of neutral detergent fiber (NDF), acid detergent fiber (ADF), and crude protein (P < 0.05). The effect of experimental diets on the apparent digestibility of dry matter, organic matter, NDF, ADF, and crude protein was not significant (P < 0.05). Ammonia nitrogen concentration, pH, and a total population of ruminal fluid protozoa and blood parameters were not affected by experimental diets. Parameters of fattening performance such as feed intake, live weight changes, feed conversion ratio, some carcass traits such as mean weight and size of carcass parts, and colorimetric indices of muscle tissue in the order of fattening lambs were not affected by experimental diets. The use of foliage of siris in the diet of fattening lambs as a substitute with part of alfalfa had no adverse effect on the characteristics studied in the present experiment. Therefore, siris be recommended as part of the diet of fattening lambs.

Keywords  Daily weight gain · Carcass yield · Colorimetry · Tissue · Muscle

Introduction
Due to the high price of feed resources, including dietary protein sources, always many efforts have been made to use cheap and available sources to replace with common feeds in ruminant diets.

The siris plant, which is scientifically known as Albizia lebbeck, belonged to the family of Leguminosae and the sub-family of Mimosoideae. The leaves are small, green, and without thorns and hairs, and have 6 to 18 leaflets. The pods are thin, pale yellow, 15 to 30 cm long, and 2.5 to 5 cm wide, with 6 to 12 seeds (fruit) in each pod (Mozafarian 2005). The siris are widely cultivated and naturalized in some tropical and subtropical provinces of Iran, such as Khuzestan (Ahvaz, Bavi, Dezful, Shushtar, Abadan, Behbahan, and Gotvand), Bushehr, Fars, and Hormozgan (Mozafarian 2005). The siris is native to Africa and tropical Asia, northern Australia, and the tropical Americas (Balgees et al. 2009; Mozafarian 2005). In many tropical and subtropical regions of the world, it is used for feeding and other purposes, and also can be used in diet to reduce the costs of livestock feed (Babadi et al. 2017, 2018; Hassan et al. 2007). However, the use of this feedstuff may be limited due to the presence of some anti-nutritional substances such as tannins (El-Hawary et al 2011). Secondary metabolites of siris include saponins, alkaloids, terpenes, and flavonoids (El-Hawary et al. 2011; Rashid et al. 2003). Annual shedding of flowers, leaves, and pods of the siris tree over 122 days period can be a source of nitrogen for livestock (Kennedy et al. 2002).

The siris is a plant source with proper nutritional value (Babadi et al. 2017, 2018; Hassan et al. 2007), which contains low fiber and saturated fats. The foliage of versatile trees such as siris, Leucaena leucocephala, Morus alba, and
Azadirachia indica can be used as a useful and inexpensive source of protein in ruminant diets (Patra et al. 2003). Concentrations of tannins in the leaves and seeds of siris were reported to be 4% and 5.3%, respectively (El-Hawary et al. 2011). The main feature of tannins is binding to nutrients, especially proteins, which has an enzymatic inhibitory effect (Samtiya et al. 2020). The rapidly degradable (α coefficient) and potential of degradability of protein in leaves and pods of siris were more than alfalfa forage (Yousefi et al. 2017).

Replacing 5, 50, 75, and 100% siris pods or leaves instead of alfalfa in sheep diets showed that levels above 50% siris pods in the diet reduced the in vitro digestibility of neutral detergent fiber (NDF) (Yousefi et al. 2014). However, diets containing siris leaves had a higher dry matter digestibility than the control diet (containing alfalfa, without replacement with alfalfa) (Yousefi et al. 2014). Hassan et al. (2007), in a study on the effect of using siris pods and seeds in the diet of farm animals, concluded that seeds could be used as a protein supplement and pods as an important source of trace elements in diets. Kennedy et al. (2002) showed that the use of 15% siris leaves in the sheep diet increases feed intake and dry matter digestibility by 52% and 67%, respectively. It was reported that in goats fed beagasse supplemented with siris leaves, the nutrient intake and digestion were increased (Balgees et al. 2009). This study showed that siris could be used as a source of nitrogen effectively and economically to improve the use of fibrous materials during the dry season (Balgees et al. 2009).

The results of Babadi et al. (2017) with Najdi goats showed that dry matter and protein intake, digestibility of dry matter and NDF, and rumination time (kg/DMI) were significantly higher in diets containing 50% and 75% siris instead of alfalfa; and a diet 75% siris has the highest record (Babadi et al. 2017). The feeding of siris to livestock had no significant effect on glucose and cholesterol (Babadi et al. 2017, 2018). The inclusion of siris pod in the diet reduces in vitro ammonia nitrogen concentration (Yousefi et al. 2014). Most research with siris has been done in vitro, although there have been a few in vivo experiments that have often only measured digestibility. Therefore, considering that siris is abundant in tropical regions of Iran and the world and there were no studies on the effects of its use in fattening livestock, in this study, the effect of its use in the diet on digestion, fermentation, growth performance, and meat color of fattening lambs was investigated.

Materials and methods

The present experiment was performed in the educational-research station of Agricultural Sciences and Natural Resources University of Khuzestan (ASNRUKH) according to the Care and Use of Agricultural Animals in Research and Teaching guidelines (FASS 2010), and the study was approved by the ASNRUKH Animal Care Committee.

Twenty-seven Arabi male lambs 6 ± 1.3 months old and a weight of 28.8 ± 1.99 kg were selected for the present experiment. The feeding and management conditions of the selected lambs were the same, and lambs were randomly divided into three groups with no significant difference in mean initial weight. The duration of the experimental period was 75 days, including 15 days for adaptation to the diet and environment and 60 days for the sampling and data recording and collecting. The experimental diets (Table 1) were formulated according to the standard requirements of small ruminants (NRC 2007).

The three experimental treatments were diets 50 and 75% replacement of siris foliage instead of alfalfa forage in the control diet (no siris), including (1) control diet (no siris), (2) diet 50% replacement (or ration containing 15% siris), and (3) diet 75% replacement (or ration containing 22.5% siris). During the experimental period, the lambs were kept in metabolic cages, fed ad libitum, and had free access to drinking water.

Lamb feed intake was measured daily by weighing the feed, and the remaining amount was measured after 24 h. In order to determine the digestibility of feed nutrients, in the last days of the experimental period (days 49–56), the method of collecting whole feces was used. The feces of lambs and the rest of their feed were collected, weighed, and recorded every 24 h. A constant percentage of them was then stored in the cold storage for 7 days. At the end of the seventh day, samples from each animal were mixed and used for subsequent measurements. Nutrient digestibility was calculated from differences in feed, residue, and feces.

For in vitro digestion and fermentation measurement, the samples were dried (48 h, 60 °C) with an oven (Fanazm Tajhiz Gostar Co., 24 I, Iran) and ground (2-mm mesh) (AOAC 2012). Chemical composition of experimental diets containing siris, including crude protein (CP, FOSS kjeltec 2300 analyzer, Sweden; method 990.03-AOAC 2012) (Kjeldahl method, Automatic Kjeltec V50, Tehran Laboratory Industries, Iran. method 990.03-AOAC 2012), ether extract (Soxhlet method, method 954.02-AOAC 2012), dry matter, ash (method 942.05-AOAC 2012), acid detergent insoluble fibers (ADFom, AOAC 2012 # 973.18), and total tannin (for siris), was measured by standard methods (AOAC 2012). Neutral detergent insoluble fibers (NDFom, corrected for ash, without alpha-amylase, and with sodium sulfate) were measured by the usual method (Van Soest et al. 1991).

In the last days of the experiment (on the 56th day), the rumen fluid of lambs was sampled via the stomach tube before morning feeding and at 3 h after that for measuring protozoa and ammonia nitrogen, respectively. The rumen pH (pH meter, WTW, Germany) was recorded immediately after sampling (at 3 h after the morning feeding). The ruminal
fluid was filtered with four layers of cheesecloth and acidified with an equal proportion of 0.2 M hydrochloric acid and used to measure ammonia nitrogen using the phenol-hypochlorite method (Spectrophotometer, Bio-Rad, Libra S22, Cambridge, England), (Broderick and Kang 1980). In order to stabilize protozoa, an equal amount of ruminal fluid was mixed with 37% formaldehyde solution (diluted 50:50 with distilled water), and protozoa were counted by the Neubauer counting chamber (Dehority 2003) using an optical microscope (NIS-Elements F 3.0, Japan).

On the 65th day, to evaluate the effect of experimental diets on blood parameters, the blood samples were taken from the jugular vein in tubes containing EDTA-Ca 10%, about 3 h after the morning meal. Blood samples were centrifuged at 3000 rpm for 15 min (Hermel 236 HK, Germany), and the plasma was separated. Blood parameters were measured using an auto-analyzer (Mindray BS200, Guangzhou, China).

The lambs were weighed before the experiment and then every 15 days before morning feeding with 14–16 h of fasting, to study growth and fattening performance. Considering the amount of feed consumed and weight changes, feed conversion ratio, feed efficiency, and average daily gain and final weight were calculated (Eynipour et al. 2019).

On the last day of the experiment, the lambs were slaughtered after about 12 h of starvation, and then different carcass parts were weighed (Fisher and de Boer 1994).

### Table 1

| Feed ingredients and chemical composition of the experimental diets fed to lambs | Treatments | Percentage of siris replacement with alfalfa (siris in the whole ration) |
|---|---|---|
| Feed ingredients (%DM) | Control (without siris) | 50% siris | 75% siris |
| Alfalfa hay | 30 | 15 | 7.5 |
| Siris | 0 | 15 | 22.5 |
| Wheat straw | 20 | 20 | 20 |
| Barley grain | 20 | 20 | 20 |
| Corn grain | 3.5 | 3.5 | 3.5 |
| Canola meal | 5 | 5 | 5 |
| Wheat bran | 20 | 20 | 20 |
| Salt | 0.5 | 0.5 | 0.5 |
| Mineral + vitamin premixa | 1 | 1 | 1 |

**Chemical composition**

- Dry matter (%) -
- Ash (%) -
- Crude protein (%) -
- NDF (%) -
- ADF (%) -
- Tannin (%)b -
- ME (Mcal/kg)c -
- NEI (Mcal/kg)d -
- Percentage of digestibility of organic matter (DOM)e -
- Apparent digestibility of organic matter (IVOMAD) (%)f -
- Short-chain fatty acids (SCFA) (mmol/200 mg DM)g -

**Notes:**

aPremix contained (per kg): vitamin A, 500,000 IU/mg; vitamin D3, 100,000 IU/mg; vitamin E, 100 mg/kg; Ca, 180 g/kg; P, 60,000 mg/kg; Na, 60,000 mg/kg; Mg, 19,000 mg/kg; Zn, 3000 mg/kg; Fe, 3000 mg/kg; Mn, 19,000 mg/kg; Cu, 300 mg/kg; Co, 100 mg/kg; Sc, 1 mg/kg; I, 10 mg/kg; antioxidant, 400 mg/kg; carrier, up to 1000 g.

bTotal tannin measured for siris in the present experiment was 3.66%.

cME (MJ/kg DM) = 0.04 + 0.1639GP + 0.0079CP + 0.0239EE (mixed feed).

dNEL (Mcal/kg) = 0.04 + 0.1195GP + 0.0051CP + 0.0152EE (mixed feed).

 eDOM (%) = 0.9042 GP + 0.0492CP + 0.0387CA + 16.49 (n = 85, r2 = 0.93) (forage).

fIVOMAD (%) = 14.88 + 0.8893GP + 0.0448CP + 0.0651 ash.

gSCFA (mmol/200 mg DM) = 0.0222 GP − 0.00425.
After slaughtering the lambs, the color of the order muscle, located between ribs 12 and 13, was measured according to the L* (brightness), a* (redness), and b* (yellowness) systems using a colorimeter (Konica, model CR 400, Japan), with three replicates for each sample. The chromaticity and hue were calculated according to the colorimetric indexes (AMSA 2012).

Data were statistically analyzed using the GLM procedure of SAS (version 9.4). The means were compared by Duncan's multiple range test at the error level of 0.05 for significant effects.

\[ Y_{ij} = \mu + T_i + \epsilon_{ij} \]

In this model, \( Y_{ij} \) is the observed value, \( \mu \) is the population mean, \( T_i \) is the effect of the ith treatment, and \( \epsilon_{ij} \) is the residue error.

**Results**

**Nutrient intake and digestibility**

The results of the experiment showed (Table 2) that the effect of experimental treatments on the dry matter, organic matter, and protein intake was not significant (\( P > 0.05 \)). However, it had a significant effect on NDF and ADF intake (\( P < 0.05 \)).

The effect of experimental diets on the apparent digestibility of dry matter, organic matter, ADF, and crude protein was not significant (\( P > 0.05 \)). However, replacing alfalfa with siris in the diet caused to increase in the digestibility of NDF (\( P < 0.05 \)) and was the highest in 75% replacement treatment.

**Rumen fermentation parameters and protozoa population**

The results showed that the effect of experimental treatments on ammonia nitrogen concentration and ruminal pH was not significant (\( P > 0.05 \)), but numerically the concentration of ammonia nitrogen in diets containing siris was lower than its concentration in the control diet (Table 3).

Table 3 showed that experimental diets had no significant effect on the total population of ruminal protozoa (\( P > 0.05 \)). Numerically, the highest populations of holotrichs, cellulolytic, and total were observed in 75% replacement of siris.

### Table 2 Nutrients intake (g/d) and digestibility (%) in lambs fed diets containing of siris replaced with alfalfa hay

| Item                  | Treatment (percentage of siris replacement with alfalfa (% in the whole ration)) | SEM | P-value |
|-----------------------|---------------------------------------------------------------------------------|-----|---------|
|                       | Control (without siris)  50 (15)  75 (22.5)                                  |     |         |
| Dry matter            | 1439.89  1546.64  1428.08                                                   | 56.787 | 0.2877  |
| Organic matter        | 1303.05  1406.97  1299.51                                                   | 51.531 | 0.2679  |
| NDF                   | 761.24\(^b\)  940.01\(^a\)  957.67\(^a\)                                  | 32.290 | 0.0005  |
| ADF                   | 360.97\(^b\)  375.22\(^a\)  326.02\(^a\)                                  | 14.029 | 0.0158  |
| Crude protein         | 200.80  209.66  200.36                                                   | 11.468 | 0.8152  |
| Digestibility, %      | 60.83  59.88  60.86                                                   | 4.056  | 0.4338  |
| Dry matter            | 63.33  62.76  64.34                                                   | 3.595  | 0.4433  |
| Organic matter        | 52.32\(^b\)  56.59\(^ab\)  58.89\(^a\)                                  | 2.304  | 0.0031  |
| NDF                   | 36.06  39.58  37.99                                                   | 3.609  | 0.3666  |
| ADF                   | 74.51  69.50  73.09                                                   | 2.305  | 0.3311  |

### Table 3 Rumen fermentation parameters and protozoa population (× 10\(^4\)) in lambs fed experimental diets containing siris replaced with alfalfa hay

| Item                   | Treatment (percentage of siris replacement with alfalfa (% in the whole ration)) | SEM | P-value |
|------------------------|---------------------------------------------------------------------------------|-----|---------|
|                       | Control (without siris)  50 (15)  75 (22.5)                                  |     |         |
| Ammonia nitrogen, mg/100 mL | 19.56  18.09  18.08                                                   | 1.714 | 0.7839  |
| pH                     | 6.71  6.81  6.85                                                   | 0.090 | 0.5613  |
| Protozoa population    |                                                                  |     |         |
| Holotrichia spp.       | 6.25  5.57  7.37                                                   | 1.126 | 0.5301  |
| Entidiniomorph spp.    | 1.12  0.57  0.62                                                   | 0.272 | 0.3052  |
| Cellulolytic\(^a\)     | 3.25  3.57  3.87                                                   | 0.748 | 0.8412  |
| Whole Protozoa population | 10.62  9.71  11.87                                                 | 1.0527 | 0.3637 |

\(^a\)Polyplastron, Epidinium, and Eudiplodinium
Blood metabolites

Concentrations of glucose, cholesterol, urea nitrogen, LDL, HDL, triglyceride, aspartate aminotransferase, alkaline phosphatase, aspartate transferase, and blood creatine were not affected by experimental treatments ($P > 0.05$) (Table 4).

Feed consumption and growth performance

Table 5 showed that feed consumption of different periods was not affected by experimental diets ($P > 0.05$). Performance characteristics of lambs, such as weight changes, average daily gain, and feed conversion ratio, were not affected by experimental diets (Table 6).

Carcass traits

Carcass traits were not affected by experimental treatments (Table 7).

Meat color

The results showed that experimental diets had no significant effect on the $L^*$ (lightness), $a^*$ (redness), $b^*$ (yellowness), $c^*$ (chroma), and hue of the longissimus muscle of the fattening lambs (Table 8).

Discussion

Nutrient intake and digestibility

The experimental treatments had significant effect on NDF and ADF intake, which the higher NDF intake in diets containing siris may be related to the higher NDF content of these diets (Table 1), and the higher NDF of siris in compared to alfalfa (55.59 vs. 50.90) (Babadi et al. 2017). According to the results of the present experiment, when substituting different amounts (0, 50, and 75%) of siris instead of alfalfa in the diet of Najdi goats, the percentage of NDF and ADF and their intake from diets containing siris was higher than that of the control. In addition, the diet containing 75% siris had the highest intake and digestion (Babadi et al. 2018). Compared to the control diet (no siris and containing 100% alfalfa) and a diet containing 50% siris, replacement of 75% siris instead of alfalfa in the diet of Najdi goats had the highest nutrient intake, and dry matter digestibility (Babadi et al. 2018).

Agreeing with the results of the present experiment, in comparison to the control diet (no siris), in the Najdi goats fed 50 and 75% siris instead of alfalfa, NDF digestibility increased, and a diet containing 75% replacement had the

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Table 4  Concentrations of blood parameters (mg/100 mL) in lambs fed diets containing siris replaced with alfalfa hay

| Item                        | Treatment (percentage of siris replacement with alfalfa (% in the whole ration)) |
|-----------------------------|--------------------------------------------------------------------------------|
|                             | Control (without siris) | 50 (15) | 75 (22.5) | SEM | $P$-value |
| Glucose                     | 80.62                    | 76.50   | 81.62     | 4.0132 | 0.6386    |
| Blood urea nitrogen (BUN)   | 30.87                    | 35.37   | 37.00     | 2.013  | 0.1075    |
| Cholesterol                 | 66.75                    | 59.37   | 62.12     | 4.976  | 0.5790    |
| Triglyceride                | 16.25                    | 13.60   | 16.01     | 1.597  | 0.4444    |
| Aspartate transaminase (AST or SGOT) | 84.50 | 90.50   | 85.37     | 6.074  | 0.7551    |
| Alanine aminotransferase (ALT or SGPT) | 23.37 | 29.00   | 25.37     | 2.355  | 0.2535    |
| Alkaline phosphatase (ALP)  | 38.14                    | 25.00   | 23.50     | 4.950  | 0.0944    |
| Creatinine                  | 0.71                     | 0.65    | 0.69      | 0.036  | 0.4782    |
| HDL                         | 30.00                    | 23.00   | 23.86     | 3.211  | 0.2658    |
| LDL                         | 13.14                    | 12.37   | 13.62     | 0.995  | 0.4504    |

SEM standard error of means

Table 5  Feed intake (g/day) of fattening lambs fed diets containing siris replaced with alfalfa hay

| Item                        | Treatment (percentage of siris replacement with alfalfa (% in the whole ration)) |
|-----------------------------|--------------------------------------------------------------------------------|
|                             | Control (without siris) | 50 (15) | 75 (22.5) | SEM | $P$-value |
| Days 0–15                   | 1074.75                  | 1220.27 | 1135.49   | 73.740 | 0.3909    |
| Days 16–30                  | 1230.31                  | 1406.72 | 1292.16   | 72.652 | 0.2423    |
| Days 31–45                  | 1380.35                  | 1527.39 | 1392.85   | 67.895 | 0.2590    |
| Days 46–60                  | 1468.19                  | 1554.40 | 1439.49   | 55.395 | 0.3311    |
| Days 0–60                   | 1288.37                  | 1427.19 | 1314.99   | 62.957 | 0.2760    |

SEM standard error of means
highest NDF digestibility (Babadi et al. 2018). Contrary to the results of the present experiment, when substituting 25, 50, 75, and 100% siris leaves instead of alfalfa in diets of sheep, in vitro dry matter digestibility was significantly increased (Yousefi et al. 2014). However, the in vitro digestibility of NDF was not affected by the experimental treatments, which may be due to the use of leaves in their experiments compared to the complete branch or foliage (leaves, pods, and seeds) in the present experiment (Yousefi et al. 2014, 2017). On the other hand, replacing 25, 50, 75, and 100% siris flowers in diets of sheep instead of alfalfa, the in vitro digestibility of dry matter and NDF significantly increased (Yousefi et al. 2014). Agreeing with the results of the present experiment, 50 and 75% replacement of the pods of siris containing seeds instead of alfalfa forage, there was no difference in in vitro dry matter digestibility of diets (Yousefi et al. 2014). The substitution of 50 and 100% Subabel (siris family) instead of alfalfa in the diets of cattle and buffaloes had no significant effect on the in vitro digestibility of dry matter, NDF, and ADF (Shahriari et al. 2017). However, in comparison to the control (without siris), up to 50% replacement resulted in numerically increased digestibility of these nutrients and decreased them when 100% of the replacement (Shahriari et al. 2017). Feeding oak fruit kernels (such as siris containing tannins, the total tannin measured for siris in the present experiment was 3.66%) to fattening goat kids did not affect the digestibility of dry matter, organic matter, ADF, and crude protein of experimental diets (Hoseinpour-mohammadabadi and Chaji 2019), which was in agreement with the results of the present experiment.

The reason for increasing the digestibility of nutrients, especially NDF, by using siris in the diets, may be related to the lower ADF and the higher ME and protein of siris (Table 1, Babadi et al. 2017, 2018), and the higher digestibility of siris NDF in comparison to the alfalfa (Babadi et al. 2017).

**Rumen fermentation parameters and protozoa population**

The normal pH range varies from 6 to 7, depending on the type of ration. In the present experiment, the rumen pH was in the optimal range for the activity of microorganisms (Table 3). Consistent with the present results, tannin-containing feedstuff sources did not affect ruminal pH (Bhatta et al. 2007). The pH may remain constant due to no change in the protozoa population (Table 3) or fatty acid concentration, especially the propionic acid. Contrary to the results of the present experiment, the use of some tannin-containing feeds, including oak (Maldar et al. 2010), olive pomace (Yanez Ruiz et al. 2004), and siris foliage (Babadi et al. 2018), caused to decrease ruminal pH. Because protozoa have stabilizing properties in the rumen, due to the rapid digestion and storage of starch by ciliated protozoa (Hristov et al. 2001), part of the decrease in pH has been attributed to a decrease in the rumen protozoa population with the consumption of tannin-containing feed. Perhaps the reason for the difference in their results with the present experiment is the higher concentration of tannin in oak (8 to 10%) (Maldar et al. 2010) compared to siris (3.66%), which did not affect the protozoa population (Table 3).

| Item                     | Treatment (percentage of siris replacement with alfalfa (% in the whole ration)) | Control (without siris) | 50 (15) | 75 (22.5) | SEM  | P-value |
|--------------------------|---------------------------------------------------------------------------------|-------------------------|---------|-----------|------|---------|
| Initial weight, kg       |                                                                                 | 28.06                   | 29.92   | 28.49     | 1.998| 0.7898  |
| Weight day 15, kg        |                                                                                 | 30.60                   | 32.30   | 30.90     | 1.987| 0.8134  |
| Weight day 30, kg        |                                                                                 | 33.32                   | 34.82   | 33.52     | 2.038| 0.8534  |
| Weight day 45, kg        |                                                                                 | 35.95                   | 37.05   | 37.50     | 1.950| 0.8739  |
| Final weight (day 60), kg|                                                                                 | 38.57                   | 39.45   | 38.27     | 1.973| 0.9091  |
| Average daily gain, kg/day (day 0–15) |                                                                              | 0.16                    | 0.19    | 0.17     | 0.016| 0.5487  |
| Average daily gain, kg/day (day 16–30) |                                                                              | 0.19                    | 0.16    | 0.17     | 0.019| 0.6624  |
| Average daily gain, kg/day (day 31–45) |                                                                              | 0.17                    | 0.16    | 0.14     | 0.018| 0.3308  |
| Average daily gain, kg/day (day 46–60) |                                                                              | 0.19                    | 0.15    | 0.19     | 0.015| 0.1474  |
| Average daily gain, kg/day (day 60) |                                                                                | 0.16                    | 0.17    | 0.17     | 0.017| 0.8370  |
| Feed conversion ratio (day 0–15) |                                                                                | 6.49                    | 7.81    | 7.11     | 0.610| 0.3262  |
| Feed conversion ratio (day 16–30) |                                                                                | 6.94                    | 8.87    | 7.59     | 0.723| 0.1804  |
| Feed conversion ratio (day 31–45) |                                                                                | 7.81                    | 7.83    | 8.78     | 0.716| 0.5564  |
| Feed conversion ratio (day 46–60) |                                                                                | 8.57                    | 10.19   | 8.62     | 0.703| 0.2069  |
| Feed conversion ratio (day 60) |                                                                                | 7.47                    | 9.05    | 8.14     | 0.505| 0.1103  |

*SEM* standard error of means
Decreased rumen protein degradation due to tannin binding to feed protein (Yanez Ruiz et al. 2004) and reduced growth and activity of proteolytic bacteria (Min et al. 2005) resulted in reducing the ammonia nitrogen production in the rumen. Therefore, the tannin in siris may be the cause of ammonia nitrogen reduction; however, its low concentration in the diet does not make a significant difference (Frutos et al. 2002). Protozoa have proteolytic and deamination activities that lead to the production of ammonia in the rumen (Williams and Coleman 1991); therefore, due to no differences in protozoa populations among experimental treatments (Table 3), the lack of differences in ammonia nitrogen concentration was expected. Agreeing to the results of the present experiment, consumption of oak leaves (such as siris containing tannins) did not cause a significant decrease in ruminal ammonia nitrogen concentration (Yildiz et al. 2005).

The feeding of 50% and 75% of siris foliage instead of alfalfa to Najdi goat significantly reduced concentration ammonia nitrogen, and 75% replacement had the lowest concentration of ammonia nitrogen, which is contrary to the results of the present experiment; however, in the present experiment, a numerical decrease in ammonia nitrogen was observed in diets containing siris. Compared to alfalfa, feeding siris foliage to goats significantly decreased ruminal pH, and ammonia nitrogen concentration, due to the presence of tannins in siris (Babadi et al. 2017). The tannin in siris protects the feed protein against ruminal degradation, therefore, the tannin in siris may be the cause of ammonia nitrogen reduction; however, its low concentration in the diet does not make a significant difference (Frutos et al. 2002). Protozoa have proteolytic and deamination activities that lead to the production of ammonia in the rumen (Williams and Coleman 1991); therefore, due to no differences in protozoa populations among experimental treatments (Table 3), the lack of differences in ammonia nitrogen concentration was expected. Agreeing to the results of the present experiment, consumption of oak leaves (such as siris containing tannins) did not cause a significant decrease in ruminal ammonia nitrogen concentration (Yildiz et al. 2005).

### Table 7 Average weight and size of certain parts of lamb carcass fed diets containing siris replaced with alfalfa hay

| Item                          | Treatment (percentage of siris replacement with alfalfa (% in the whole ration)) | Control (without siris) | 50 (15) | 75 (22.5) | SEM | P-value |
|-------------------------------|-----------------------------------------------------------------------------|------------------------|--------|-----------|-----|---------|
| Live weight, kg               |                                                                             | 38.70                  | 40.70  | 38.53     | 1.453 | 0.5673  |
| Skin, g                       |                                                                             | 4900<sup>a</sup>       | 4400<sup>b</sup>      | 4400<sup>b</sup> | 129.099 | 0.0034  |
| Leg of lamb, g                |                                                                             | 825                    | 817.50 | 862.50    | 42.180 | 0.7441  |
| Liver of sheep, g             |                                                                             | 527.50                 | 567.50 | 619.50    | 32.978 | 0.2859  |
| Lung, g                       |                                                                             | 432.50                 | 455      | 440      | 44.558 | 0.9374  |
| Lean heart, g                 |                                                                             | 112.50                 | 132.50  | 125      | 10.206 | 0.4704  |
| Fat around the heart, g       |                                                                             | 62                     | 35      | 35       | 11.637 | 0.2981  |
| Kidneys, g                    |                                                                             | 102.50                 | 102.50  | 100      | 5.401  | 0.9326  |
| Fat around the kidneys, g     |                                                                             | 64                     | 50      | 52       | 5.916  | 0.3305  |
| Digestive system, g           |                                                                             | 9050                   | 10,100  | 9100     | 429.146 | 0.2924  |
| Fat around the gastrointestinal tract, g |                                                                 | 207.50                | 260      | 250      | 74.624 | 0.8751  |
| Thigh, g                      |                                                                             | 2570                   | 2560    | 2750     | 94.207 | 0.3946  |
| Wristband, g                  |                                                                             | 1527.50                | 1477.50 | 1587/50  | 57.355 | 0.4874  |
| Neck, g                       |                                                                             | 1410                   | 1597/50 | 1707/50  | 87.058 | 0.1934  |
| Order and fillet, g           |                                                                             | 1710                   | 1607/50 | 1657/50  | 226.743 | 0.9510  |
| The spleen, g                 |                                                                             | 60                     | 100     | 70       | 10.408 | 0.1424  |
| Testicles, g                  |                                                                             | 197.50                 | 222.50  | 220      | 20.716 | 0.6790  |
| Flank, g                      |                                                                             | 540                    | 1052.50 | 535      | 273.179 | 0.4176  |
| Head, g                       |                                                                             | 2312.50                | 2287.50 | 2210     | 144.669 | 0.8776  |
| Large colon, g                |                                                                             | 207.50                 | 420     | 317.50   | 135.723 | 0.5981  |
| Empty colon, g                |                                                                             | 154.50                 | 240     | 199.50   | 66.396  | 0.6933  |
| Small intestine full, g       |                                                                             | 2550                   | 2550    | 2175     | 136.168 | 0.2272  |
| Full rumen, g                 |                                                                             | 5938.50                | 5950    | 6250     | 383.641 | 0.8202  |
| Empty rumen, g                |                                                                             | 1132.50                | 1395    | 800      | 314.421 | 0.4943  |
| Visceral fat, g               |                                                                             | 80                     | 67.50   | 80       | 16.137  | 0.8288  |
| Tail, g                       |                                                                             | 1925                   | 3027.50 | 2060     | 245.938 | 0.0898  |
| Horn, g                       |                                                                             | 182.50                 | 285     | 335.00   | 38.106  | 0.1364  |
| Gear, g                       |                                                                             | 650                    | 820     | 1485     | 80.572  | 0.5122  |
| Half carcass weight, g        |                                                                             | 8600                   | 6922.50 | 7000     | 824.016 | 0.3878  |

*SEM* standard error of means

Means in the same row with different superscript letters are significantly different ($P < 0.05$)
Table 8 Characterization of order muscle tissue in lambs fed diets containing siris replaced with alfalfa hay

| Item       | Treatment (percentage of siris replacement with alfalfa (% in the whole ration)) | Control (without siris) | 50 (15) | 75 (22.5) | SEM | P-value |
|------------|---------------------------------------------------------------------------------|-------------------------|---------|-----------|-----|---------|
| L*         | 27.27                                                                           | 27.10                   | 25.20   | 0.655     | 0.1877 |
| A*         | 8.37                                                                            | 8.54                    | 7.39    | 0.469     | 0.3120 |
| B*         | 2.71                                                                            | 2.99                    | 2.82    | 0.327     | 0.8342 |
| Chroma*    | 8.80                                                                            | 9.05                    | 7.92    | 0.520     | 0.3879 |
| Hueb       | 1.52                                                                            | 1.52                    | 1.52    | 0.004     | 0.4635 |

L*: lightness; a*: redness; b*: yellowness;
*aChroma: chromaticity (A*2 + B*2)1/2
bHue = arc tangent B/A × 57.29
SEM standard error of means

consequently decreasing ammonia nitrogen concentration (Frutos et al. 2002). Disagreeing with the results of the present experiment, by replacing the 25, 50, 75, and 100% leaves, flowers, and pods (containing seeds) of siris instead of alfalfa in sheep diets, the pH of the culture medium were significantly reduced (Yousef et al. 2014). However, agreeing to the results of the present experiment, the ammonia nitrogen concentration of treatments with 50% replacement leaves and pods and both 50 and 75% levels of flowers was not affected by the replacement of alfalfa by siris (Yousefi et al. 2014). In contrary to the results of the present experiment, the ammonia nitrogen concentration was significantly reduced, when replacing alfalfa with 75% of leaves or pods of siris. The probable reason for these differences could be the use of leaves, flowers, and pods separately in their experiment compared to the foliage (leaf, pod, and seed) in the present experiment, because the chemical composition of these components is different from each other and foliage (Babadi et al. 2018; Yousefi et al. 2014, 2017).

It was shown that using siris (Babadi et al. 2018) and oak leaves (Maldar et al. 2010) in the ration of goats reduced the population of rumen protozoa. However, in an experiment, feeding goats with oak kernel did not affect the population of protozoa (Hoseinpour-mohammadabadi and Chaji 2019); this was attributed to the presence of the high amount of starch in the oak kernel because starch, like protein, binds to tannins and reduces its harmful effect. Tannins have an adverse effect on proteolytic bacteria, which are a feed for the protozoa (McSweeney et al. 2001). On the other hand, increasing the amount of tannin in ration decreased the ruminal pH that it can result in a reduction of the protozoa population (Dehority 2003), which is contrary to the results of the present experiment because in the present experiment, the pH was not affected by tannin of siris (Table 3). One of the reasons for not observing adverse effects for tannins of siris could be its low amount (3.66% of siris dry matter) in the whole diet and no reduction of ruminal pH (Dehority 2003). On the other hand, saponin in siris (669.40 mg/kg compared to 80.27 mg/kg for alfalfa) may also harm the population of protozoa (Babadi et al. 2017). The most significant effect of saponin is on the rumen protozoa (Wallace et al. 2002). Saponins cause cell lysis by binding to sterols in the protozoa membrane. However, because of the lack of this sterol in the bacterial membrane, saponins cannot bind to bacteria. Although many studies have shown a decrease in the population of protozoa in the presence of saponins, some have shown that its anti-protozoa effect is unstable and transient; maybe this was a reason for no adverse effects of siris, which contains saponins (Wallace et al. 2002). On the other hand, the suitable pH for protozoa in the solid part of the rumen contents is 6.7, which in the present experiment, despite the use of siris (contains tannins), the pH was in the appropriate range for their activity (Table 3).

Blood metabolites

It was shown that in comparison with tannin-free diets, feeding diets containing tannins in goats did not affect blood glucose (Merkel et al. 2001), total protein, serum albumin (Ben Salem et al. 2002), cholesterol, and blood urea nitrogen (Babadi et al. 2018), which is consistent with the results of the present experiment. Saponin (669.40 mg/kg vs. 80.27 mg/kg for in siris and alfalfa, respectively) can affect membrane permeability and reduce blood cholesterol (Hu et al. 2005), in which in the present experiment, there has been a numerical reduction; this may be due to a low concentration of saponin in the diet (Hu et al. 2005).

In the liver, blood urea nitrogen (BUN) is synthesized from absorbed ruminal ammonia nitrogen so that the concentration of BUN has a positive correlation with the ruminal ammonia nitrogen. Therefore, the lack of change in the BUN is because of no change in ruminal ammonia nitrogen (Hosoda et al. 2005). No change in the concentration of hepatic factors such as aspartate aminotransferase, aspartate transamine, and creatine indicates no damage to the parenchymal tissue of the liver (Al-Shanti et al. 2013).

Feed consumption

In an experiment, the consumption of siris was higher than that of alfalfa (Babadi et al. 2017). Using 50% and 75% of siris foliage instead of alfalfa forage in the diet of Najdi goats caused a linear increase in daily feed consumption (Babadi et al. 2018). It was shown that the use of 15% of siris leaves in the sheep diet as a supplement to seasonal dry grasses led to an increase in their consumption (Kennedy et al. 2002). This study showed that siris could be used as a
source of nitrogen effectively and economically to improve the use of fiber during the dry season (Kennedy et al. 2002). Supplementation of ammoniated sugarcane bagasse with siris leaves or wheat bran led to an increase in goat feed intake, which was the most increase in the diet containing siris leaves (Balgees et al. 2009). Overall, the use of siris in several experiments increased feed intake, which was used as a supplement to low-quality roughages, which are typically poor in nutrients, especially protein. Perhaps the lack of effect on feed intake in the present experiment is the sufficient nitrogen richness of the diet and the closer similarity of the alfalfa protein concentration with siris.

**Growth performance and carcass traits**

Animal growth rate and carcass traits depend on feed intake and dietary nutrient availability, and the efficiency of lambs in converting feed to live weight depends on various factors such as breed, sex, age, and nutritional levels (Thompson et al. 1987). Therefore, considering the level of feed consumption, the same nutrition, and race, age, and sex, the insignificant differences in most characteristics between treatments were reasonable and expected. Due to the higher price of alfalfa forage and its need for suitable land, abundant water, and special reservation conditions, no difference in the results of treatments is a valuable and positive achievement for the present experiment. Therefore, according to the results of the present experiment, the nutritional value of siris foliage is comparable to alfalfa forage, and it can be used in the diet of ruminants. It is recommended that siris nutritional value be studied as an alternative to alfalfa in other ruminants, especially dairy livestock and their offspring such as lambs, goat kids, and suckling calves, male and female calves, or heifers.

**Conclusion**

The results of the present experiment showed the use of the siris foliage in the diet of fattening lambs as a substitute with part of alfalfa not only did not hurt digestion, fattening performance, ruminal and blood parameters, but also in some cases improved them. Due to the high price of alfalfa forage and its need for suitable land, abundant water, and special reservation conditions, no difference in the results of treatments is a valuable and positive achievement for the present experiment. Therefore, according to the results of the present experiment, the nutritional value of siris foliage is comparable to alfalfa forage, and it can be used in the diet of ruminants. It is recommended that siris nutritional value be studied as an alternative to alfalfa in other ruminants, especially dairy livestock and their offspring such as lambs, goat kids, and suckling calves, male and female calves, or heifers.

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**Author contribution** M conceived and designed the research. M, H, and O conducted the experiments. M and H analyzed the data. M and H wrote the manuscript. All authors read and approved the manuscript.

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**Declarations**

**Ethical approval** The manuscript does not contain clinical studies or patient data.

**Consent to participate** This research did not involve human subjects, or human transplantation studies, and no organs/tissues were obtained from the prisoners.

**Conflict of interest** The authors declare no competing interests.

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