

Feeding of carob (Ceratonia siliqua) to sheep infected with gastrointestinal nematodes reduces faecal egg counts and worm fecundity

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1 2 Feeding of carob (Ceratonia siliqua) to sheep infected with gastrointestinal nematodes 3 reduces faecal egg counts and worm fecundity 4 Katerina Saratsi^{a,b}, Hervé Hoste^{c,d}, Nikolaos Voutzourakis^a, Nikolaos Tzanidakis^a, 5 Alexandros Stefanakis^a, Stig Milan Thamsborg^e, Irene Mueller-Harvey^f, S., Ioannis 6 Hadjigeorgiou^{b+}, Smaragda Sotiraki^{a+}, 7 8 ^aVeterinary Research Institute, Hellenic Agricultural Organization-Demeter, Thermi 9 10 57001, Thessaloniki, Greece 11 ^bDepartment of Nutritional Physiology and Feeding, Faculty of Animal Science, 12 Agricultural University of Athens, 11855 Athens, Greece 13 ^cINRAE, UMR 1225 IHAP, 23 Chemin des Capelles, Toulouse F-31076, France 14 ^dUniversité de Toulouse, ENVT, 23 Chemin des Capelles, Toulouse F-31076, France 15 ^eDepartment of Veterinary and Animal Sciences, Faculty of Health and Medical 16 Sciences, University of Copenhagen, Copenhagen, Denmark. 17 ^fSchool of Agriculture, Policy and Development, University of Reading, P.O. Box 18 236,1 Earley Gate, Reading RG6 6AT, UK. 19 20 *Corresponding author: Saratsi Katerina, e-mail address: saratsikaterina@yahoo.gr 21 ⁺authors equally supervised the work 22 23 24 25

Abstract

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The present study explored the anthelmintic effects of condensed tannins (CT) in carob (Ceratonia siliqua) pods fed to sheep against gastrointestinal nematodes. Three independent in vivo trials tested whether i) carob pod (CaBP)-containing feed had an anthelmintic effect and if yes, which was the optimal concentration in the diet; ii) whether this effect could be attributed to tannins through the polyethylene glycol (PEG) test and iii) whether there were any synergistic effects when combined with another tannin-containing feed (e.g. sainfoin). In all trials 6-month old nematodenaive lambs, experimentally infected with both Haemonchus contortus and Trichostrongylus colubriformis, were used. Faecal egg counts (FEC) were performed regularly and at the end of each trial adult worm counts (AWC) and female worm fecundity were recorded. In trial 1, 35 lambs (five groups of seven lambs) were fed different CaBP concentrations ranging from 0% to 12% w/w. FEC declined up to 39.2% only in the group fed with 12%CaBP, while a declining trend (P<0.06) was demonstrated for the AWC of T. colubriformis, which was associated with the increasing concentration of CaBP in feed. Female worm fecundity was reduced in groups fed CaBP for both parasites, however this was only significant for H. contortus (P<0.001), in a dose dependent manner. In trial 2, four groups of six infected lambs each were used, which received the carob diets CaBP or CaBP+PEG, and the tannin-free diets with or without PEG (C or C+PEG). Results showed that FEC of Groups C, C+PEG, and CaBP+PEG were comparable throughout the trial, while the group receiving only CaBP showed lower FEC from DAY 25 onwards. AWC showed a reduction (67.7%) only for *H. contortus* (P<0.03). Reversal of the anthelmintic effect of CaBP after PEG administration suggested that CT contributed to the anthelmintic action. However, no effect of CaBP was observed on T.

colubriformis AWC and on female worm fecundity for both species. Finally, for trial
3 four groups of six lambs each received a diet based on CaBP, sainfoin (S) or a
combination (CaBP+S) and were compared to a control (C) diet of lucerne. On DAY
37 FEC values in groups CaBP+S and S tended to be lower compared to the two other
groups (C, CaBP), while for AWCs no significant differences were observed for both
parasites. The fecundity of <i>H. contortus</i> and <i>T. colubriformis</i> demonstrated significant
differences between the treated and control groups, with lower values in the animals
receiving CaBP+S. Overall, the results supported the hypothesis that carob had an
anthelmintic effect due to its CT, but there was no clear indication of a synergistic
effect with sainfoin.
Keywords: Carob, Sainfoin, Haemonchus contortus, Trichostrongylus colubriformis,
gastrointestinal nematodes, sheep, feed additives

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1. Introduction

Infections by gastrointestinal nematodes (GIN) affect both health and welfare of grazing ruminants, causing anorexia, impaired digestion and nutrient absorption with related production losses, diarrhoea, anaemia and even death (Perry and Randolph, 1999; Stear et al., 2007; Hoste et al., 2016). Currently, the control of these parasites relies on repeated dosing with commercial anthelmintic drugs. However, the development of anthelmintic resistance in worm populations against one or multiple classes of anthelmintic has become a serious problem in several regions of the world, making it increasingly difficult to control parasitic infections (Kaplan, 2020). At the same time the increasing concerns of consumers about the presence of drug residues in foods and in the environment (McKellar, 1997) have stimulated the search for alternative or complementary solutions (Hoste and Torres-Acosta, 2011) within the context of organic farming and sustainable agriculture (Charlier et al., 2018). Such alternatives include the use of bioactive plants with anthelmintic properties. Many results indicate that such resources, because of the presence of plant secondary metabolites (PSMs), might help to interfere with the biology of key-stages during nematode cycle and to reduce the consequences of GIN infections in grazing ruminants. Particular attention has been given to plants containing condensed tannins (CT) and some related polyphenols (see reviews by Terrill et al., 2012; Hoste et al., 2015, 2016). Based on previous studies, the need to explore new plant resources to develop non-drug-based strategies for the integrated control of nematode parasites in grazing ruminants has recently become a research priority in livestock production, especially in ruminant breeding as also reviewed by Morgan et al, 2020.

Small ruminants (sheep and goats) are a major component of the dairy sector in the Mediterranean basin (Hadjigeorgiou et al., 2005). Sheep and goat production often occupy marginal lands that are unsuitable for crop production but are rich in local plants, such as rangeland vegetation, which can be exploited by animals as a feed resource (Frutos et al., 2008; Méndez-Ortiz et al., 2018). Many rangeland plants also contain PSMs, such as tannins (Papachristou et al., 2005) and several in vitro and in vivo studies have evaluated their anthelmintic effects against GINs of small ruminants (Manolaraki et al., 2010; Moreno-Gonzalo et al., 2012, 2013a,b, 2014; Arroyo-Lopez et al., 2014; Silva Soares et al., 2018). Overall, CTs have been shown to directly or indirectly interfere with the life cycle of several GINs and, therefore, CT-containing plants, which also include many legumes, are proving to be beneficial nutritional resources. However, a high degree of variability with respect to their anthelmintic activity has also been recorded. Besides the total tannin concentration in ruminant diets, recent studies have demonstrated that CT molecular composition or structural characteristics can also affect anthelmintic activity (Mueller-Harvey et al., 2019). Carob (Ceratonia siliqua) and sainfoin (Onobrychis viciifolia) are both resources of the Fabaceae family and contain CTs. Carob is a leguminous tree that is widely cultivated in the Mediterranean area. It is an important species both for economic and environmental reasons (Batlle and Tous, 1997). Carob pods (fruits) are mostly used in the food industry; pulp accounts for 90% by pod weight and seeds for 10%. They contain high sugar (48–56%), but low protein (3–4%) and lipid concentrations (0.4– 0.8%) (Marakis, 1996; Batlle and Tous, 1997). Moreover, ripe carob pods contain high concentrations of CTs (16-20% w/w DM) (Bravo et al., 1994; Batlle and Tous,

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1997). This has been debated by Priolo et al. (2000, 2002) who claimed that the pods

have low content of CTs, but with exceptionally high biological activity. Silanikove et

al. (2006) have demonstrated that the yield of CTs is considerably affected by the extraction method applied (from 5.0% with acidic methanol to 17.2% with urea-buffer solution), suggesting that carob pods are a rich source of CTs. The high CT concentration in by-products from carob pod processing justifies researching its value as a feed additive with possible effect against GIN species. Sainfoin, which can be found especially in southern parts of Europe, has been the subject of renewed interest because of its beneficial effects in the context of agroecology (Hayot Carbonero et al., 2011), its beneficial impact on ruminant production and the environment and its potential antiparasitic effects on small ruminants (Manolaraki et al., 2010; Hoste et al., 2015; Saratsis et al., 2016; Mueller-Harvey et. al., 2019). In vitro studies have shown that sainfoin extracts have a dosedependent effect against different GIN species (Brunet et al., 2007; Manolaraki et al., 2010; Novobilsky et al., 2013). Moreover, in vivo anthelmintic effects have also been described in sheep and/or goats fed with sainfoin; i.e. 42-68% reduction in parasitic egg excretion, which was associated with a 17.6% decrease in female worm fecundity and a 45% decrease in worm numbers for *Haemonchus contortus* (Arroyo-Lopez et al., 2014).

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The present study, therefore, sought to explore the anthelmintic effects of feeding regimes employing two CT-containing plant resources that may be relevant for Mediterranean conditions. These were offered to lambs either alone or in combination to evaluate their efficacies against two GIN species (*H. contortus* and *Trichostrongylus colubriformis*). The specific objectives were to explore whether a) the anthelmintic effect of carob in the feed is dose dependent (Trial 1), b) this anthelmintic effect is associated with tannins by using polyethylene glycol (PEG) as a

tannin-inhibitor (Trial 2), and c) there are any synergistic effects between carob and sainfoin feeds (Trial 3).

2. Materials and Methods

2.1. Stabling and animals

The experiments were carried out at the Asomaton Research Station of HAO Demeter on the island of Crete, Greece. The animal barn was of an open-sided shed type, with straw bedding. During the whole study the animals were kept indoors with each group in a separate pen of approximately 10 m² and 10 m² open yard. The study included three trials with lambs belonging to the local "Sfakion" breed. In order to achieve uniformity of the experimental animals, all lambs included were female, 6-month-old with a comparable body weight (BW), which was within the normal BW range of the breed (22-30 kg) at the specific age. The lambs were raised indoors under helminth-free conditions. Fourteen days before the start of each trial, they were drenched with albendazole at the higher commercially recommended dose (ALBENDAZOLE Drench, PROVET, 7.5 mg/kg) and they tested negative by faecal egg counts at the start of each trial. No anthelmintic resistance was previously recorded for this specific flock.

2.3. Infective larvae

Third-stage infective larvae of H. contortus and T. colubriformis strains, susceptible to all classes of anthelmintic drugs, had been cultured from faeces of monospecifically infected donor sheep. Larvae were recovered using the Baermann technique and then stored for 1-2 months at 4° C until use.

2.4. Tannin-containing plant resources

174 Carob pods (after removal of the seeds) were locally purchased and offered as crushed 175 flour meal incorporated in the concentrate feed supplement. Sainfoin pellets (Perly 176 cultivar, 3rd cut) were provided by Multifolia (Viapres le Petit 10380, France) as part 177 of the Research project CARES.

2.5. Tannin concentration and composition

Tannin concentrations and compositions were determined in triplicate using two different assays, i.e. the acetone-butanol-HCl assay and the thiolytic degradation with benzyl mercaptan. Both techniques were applied in order to ensure a comprehensive analysis since it was previously demonstrated that, depending on the types of CTs, the acetone-HCl-butanol assay can give higher CT concentrations than the thiolysis assay. The acetone-HCl-butanol assay was carried out as previously described by Grabber et al. (2013) and Desrues et al. (2017).

The thiolysis reaction was carried out with benzyl mercaptan (Gea et al., 2011; Ropiak et al., 2016), the reaction products were identified by HPLC-MS analysis (Williams et al., 2014; Desrues et al., 2017) and quantified based on peak areas at 280 nm (Gea et al., 2011; Ropiak et al., 2016). This provided information on CT concentration (g CT/100 g DW), CT size (in terms of mean degree of polymerisation, mDP), molar percentages of prodelphinidins (PD) and procyanidins (PC) within CTs, and molar percentages of *cis*- vs *trans*- flavan-3-ol subunits (Ropiak et al., 2016).

2.6. Experimental design

All diets offered to the animals during the experimental period (with or without the tannin sources) were formulated to meet the nutrient requirements of the animals (NRC, 2007) and the total rations were always iso-nitrogenous and iso-energetic as

well as balanced for crude fibre, Ca, P and Ca/P ratio (Suppl Table 1). Animals had access to clean water at all times. The animals' appetite was assessed and feed consumption (as feed offered minus refusals) was recorded on a daily basis by the farm manager.

2.6.1. *Trial* 1

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- To determine the anthelmintic effect of carob pod meal and to define the optimal concentration in a sheep ration, a subset of 35 lambs were randomly allocated to 5
- 204 groups (n=7 lambs/diet) (Table 1).
- 205 Carob meal (CaBP) was offered as feed supplement, at increasing rates of 0%, 3%,
- 206 6%, 9% and 12% (g CaBP/100g DM) of the total ration. The highest proportion, of
- 207 carob meal contributed to concentrate feed was set to 12% (due to its poor energy and
- 208 protein contents) in order to enable formulating a ration, which could cover the
- 209 nutritional requirements of lambs.
- Feeding the experimental diets started 2 weeks prior (D14) to experimental infection
- with nematode larvae (D0) in order for the animals to adapt to the feed.
- 212 On DAY 0, all lambs in groups (i) to (v) were infected with a single dose of 12.000
- 213 3rd stage larvae (L3) of *H. contortus* and 12.000 L3 of *T. colubriformis*. At the end of
- 214 the experimental period (D49), all lambs were euthanised by injection of a massive
- 215 dose of pentobarbital (Dolethal®).

2.6.2. Trial 2

- Four groups of 6 lambs were included in a two-factorial trial (diet and PEG-addition).
- 218 Two groups were offered CaBP as feed supplement at the rate of 12% in the total
- 219 ration, and two groups remained on standard diet (Table 1). Half of the lambs in each
- 220 diet group were offered PEG (Polyethylene Glycol 4000, Fisher Scientific USA)

221 orally (60 g/lamb diluted in 200 ml water) on a daily basis after being allocated into 222 groups. 223 On D0, all lambs were experimentally infected with 8.000 L3 of H. contortus and 224 16.000 L3 of T. colubriformis. On D21, after parasite infection was confirmed by positive faecal examination, the animals were allocated into 4 groups of 6 lambs each, 225 226 according to the experimental diets. On D37 they were euthanised as described above. 227 **2.6.3.** *Trial 3* 228 To determine the possible synergistic anthelmintic effects between 2 CT-containing 229 resources namely carob (C. siliqua) and sainfoin (O. viciifolia), 4 groups of 6 lambs 230 were included in a two-factorial design (Table 1). 231 On D-14 each group of lambs received the allocated diet, containing i) carob meal 232 (CaBP) alone ii) sainfoin (S) pellets; iii) a combination of carob meal and sainfoin 233 pellets (CaBP+S) while iv) a control group (C), received an isoproteic diet based on 234 lucerne. Carob was offered as a feed supplement at the rate of 12% in the total ration. 235 Sainfoin was offered as pellets representing 35% of the total ration. On D0 all lambs 236 were infected with a single dose of 12.000 L3 of H. contortus and 12.000 L3 of T. 237 colubriformis. At the end of the experimental period (D37), all lambs were euthanised 238 as previously described. 239 2.7. Pathophysiological parameters 240 Individual blood samples were collected once weekly (from D0 to D49) during Trial 1 241 and once every two weeks (from D0 to D28) during Trial 3, by jugular venipuncture

into heparinized tubes (BD Vacutainer®, UK) to determine the packed cell volume

(PCV), as an indicator of anaemia, according to the micro-haematocrit method. In

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Trial 2 due to its short duration, the recording of PCV values was not included in the design.

2.8. Parasitological parameters

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(10%) of the total volume.

Individual faecal samples were collected weekly directly from the rectum, during the 1st and 3rd trial, and twice weekly during the 2nd trial in order to determine faecal egg counts (FEC) using a modified McMaster technique (Roepstorff and Nansen, 1998). FEC data were expressed as eggs per gram of faeces (EPG). At necropsy, the abomasa and the first 12 meters of small intestine were separated, ligated, rapidly removed and immediately processed to collect the adult worms from the luminal contents. For the intra-mucosal larvae, pepsin digestion was applied both on the abomasum and intestinal mucosa (MAFF, 1986). After 4h incubation at 37°C the larvae were collected. After storage in 10% alcohol, worm counts were performed according to a 10% aliquot technique (MAFF, 1986). Morphological identification of worm stages, sex and species were conducted using standard procedures (MAFF, 1986). The fecundity of female worms was measured on 10 worms per lamb. For T. colubriformis, eggs were counted directly in utero after clearing in 85% lactic acid solution. All egg counts were performed under a microscope set at 10 times magnification (total 100 ×). For H. contortus, the fecundity was determined using the method described by Kloostermann et al. (1978). Briefly, the worms were soaked for 5 min in a large volume of distilled water, before being placed individually in microtubes with 1000 µl of 0.125% hypochlorite concentration solution and kept at room temperature for 20 minutes. Treatment resulted in female worms disintegrating thus enabling the direct counting of eggs under a stereo-microscope using an aliquot

2.9. Statistical analyses

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The data of FEC and adult worm counts (AWC) were log10(x+1) transformed prior to analysis. For the FEC values, comparison of all groups was first performed using an analysis of variance (ANOVA) with time as repeated measurement. Then, the comparison of results to the control values were carried out date by date, using oneway ANOVA completed by the post-hoc Bonferroni test for pairwise comparisons. Group means of AWC were compared by one-way ANOVA (Trial 1) or two-way ANOVA (Trial 2: CaBP +/- and PEG +/-; Trial 3: CaBP +/- and sainfoin +/-). Regarding the fecundity of female worms, the Shapiro-Wilk Test of normality, which is more appropriate for small sample sizes, was used. In cases where the data deviated significantly (P<0.05) from a normal distribution (Trial 1 and 3 for both parasite species and Trial 2 for T. colubriformis) the appropriate test to check the difference of fecundity between the groups, which is the non-parametric test of Kruskal-Wallis, was used. Where the dependent variable was normally distributed (P>0.05) the parametric test of one-way ANOVA (H. contortus of Trial 2) was used. Additionally, for the Trial 1, the model of linear regression was used, in order to be investigated if there was a negative correlation between the variables "percentages of carob" and "fecundity of female worms" for both parasite species (H. contortus and T. colubriformis). Finally, the Tukey HSD test was used for data of trial 3, in order to investigate statistically significant differences between groups.

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2.10. Ethical considerations

292 The study was carried out in compliance with the national animal welfare regulations.

All statistical analyses were performed using the SyStat SPSS 9.0 Software.

293 All trials took place in a Research Station of the Veterinary Research Institute. The

experimental protocol was approved by the responsible institutional committee (VRI Committee for Approval of Experimental protocols as appointed at 26/5/2014, Decision nr 972). Euthanasia was performed in a humane manner according to EU regulations.

3. Results

The CT concentrations and compositions are presented in Table 2. The HBA assay yielded similar CT concentrations for both plant materials, whereas the thiolysis assay generated lower CT concentrations for the sainfoin pellets. The thiolysis assay revealed that: i) both carob and sainfoin CTs consisted mainly of prodelphinidins, 96.7 and 74.7 mole percentages, respectively; ii) carob CTs were highly galloylated (i.e. 41.1% of flavan-3-ol subunits are galloylated), but sainfoin CTs did not contain any esterified galloyl groups; iii) carob CTs were characterised by a relatively high average molecular weight (mDP = 31.1), whereas sainfoin CTs had an mDP value of 11.5.

3.1. Trial 1

- 311 The results of Trial 1 are shown in Table 3 and Figure 1
- The analyses of FEC, based on the ANOVA on Repeated Measures from D21 to D49, showed an overall non-significant difference between groups, but significant difference over time (between days of sampling). Meanwhile, the date-by-date ANOVA of FEC showed no significant differences between groups, whatever the date, as well as no dose effect. Reduction in FEC, up to 39.2% on DAY 49 as compared to controls, was observed only for the group fed with the highest concentration of carob meal.

- For *H. contortus*, the AWC declined in the groups receiving the highest concentration
- of carob meal but this effect was not statistically significant (P= 0.964). In contrast,
- 321 there was a declining trend (P<0.06) for the numbers of T. colubriformis with
- increasing carob concentration.
- 323 The fecundity values showed significant differences (15.6%-59.3% lower than
- 324 0%CaBP respectively from the lowest to the highest CaBP concentration) between
- groups for *H. contortus* demonstrating a dose dependent effect (P<0.05).
- 326 The Box plot (Figure 1b) for *H. contortus* fecundity suggests that worms from the
- 327 0%CaBP group tended to be more fecund than other CaBP groups and there may be
- 328 some degree of fecundity discrepancy between CaBP groups. This trend was
- 329 confirmed with the non-parametric test of Kruskal-Wallis, which showed that there
- were statistically significant differences in fecundity between the groups (P<0.001).
- 331 More specifically, fecundity was statistically significantly greater for 0% CaBP group
- than the other CaBP groups. On the other hand, regarding *T. colubriformis* fecundity,
- 333 there was no statistically significant difference between the groups (P=0.128).
- 334 However, the model of linear regression, which was implemented and was
- statistically significant (P<0.05), showed a negative correlation between the variables
- "group" and "fecundity" for both parasite species.
- No GIN larvae were recovered after pepsin digestion.
- Mean PCV values (±SD) for groups 0%CaBP, 3%CaBP, 6%CaBP, 9%CaBP, and
- 339 12%CaBP on the last day of the trial were 25.29 (± 5.96), 23.00 (± 5.72), 21.00
- (± 6.32) , 23.00 (± 5.89) and 24.00 (± 5.00) respectively. No significant differences were
- 341 found between the groups in PCV.
- 342 Average daily gain (ADG) as calculated for the whole trial duration for 0%CaBP,
- 343 3%CaBP, 6%CaBP, 9%CaBP and 12%CaBP groups was (mean±s.d.) 69.2 g (±31.0),

 $61.5(\pm 36.1)$, $68.7(\pm 33.0)$, $74.8(\pm 37.5)$ and $64.4(\pm 32.9)$ g respectively, which yielded 344 345 no significant differences between the groups.

3.2.Trial 2

346 347 The results of Trial 2 are presented in Table 4 and Figure 2. 348 The Repeated Measurements Analyses of FEC showed an overall statistical difference 349 (P<0.001) between the 4 groups. The date-by-date ANOVA of FEC indicated that differences were most prominent on DAY 29 (significant statistical differences, 350 351 P<0.02) and then on DAY 33 (trend, P<0.07). Specifically, the values of the C, 352 C+PEG, CaBP+PEG groups were comparable throughout the trial, while the group 353 receiving only carob (CaBP) showed consistently lower FEC starting from DAY 25 354 until the last day of the experiment. It was evident that the effect of carob on FEC was 355 nullified by PEG. 356 Results on AWC, showed reduction only for *H. contortus* (P<0.03) resulting in an 357 overall statistical difference between the 4 groups, since the lowest worm counts were 358 found for the CaBP group. Especially, for *H. contortus*, a reduction of approximately 359 65% was observed in the carob group compared to the control. The AWC in the 360 CaBP+PEG group were similar to the other 2 control groups showing no reduction in 361 worm population. On the other hand, no effect of carob was observed on T. 362 colubriformis worm counts. 363 No effect of carob on female fecundity was also observed, irrespective of the parasite 364 species. Both control and carob groups showed comparable levels of female fecundity 365 for the two parasite species. The Box plot in Figure 2b showed that the range of 366 fecundity of H. contortus for CaBP group was greater than for C, C+PEG and CaBP+PEG groups and the interquartile range (middle 50% of the records) was lower 367 368 on the fecundity scale in the CaBP group than in the other groups.

- No GIN larvae were recovered after pepsin digestion.
- 370 The average daily gain (ADG) of lambs as calculated for the whole trial duration for
- 371 (C), (C+PEG), (CaBP) and (CaBP+PEG) groups was $51.8(\pm 30.1)$ ($\pm s.d.$), $69.8(\pm 19.9)$,
- $60.8(\pm 29.3)$ and $40.5(\pm 25.6)$ g, respectively, which resulted in no significant
- differences between the groups.

374 **3.3.Trial 3**

- 375 The results of Trial 3 are shown in Table 5 and Figure 3.
- 376 The FEC values of all experimental groups remained at very low levels up to DAY
- 377 21. The overall repeated analyses based on 3 dates of the patent phase (DAY 21,
- 378 DAY 28, DAY 37) showed a trend for differences (P<0.07) between groups. The
- 379 results of the date-by-date ANOVA test did not show difference on DAY 21 and on
- DAY 28, while on DAY 37, the values of FEC in groups CaBP+S and S tended to be
- reduced (P< 0.06) compared to the two other groups. When compared to the control
- 382 values of FEC, the reductions in the 3 treated groups ranged from 44.6% to
- approximately 86 %. These differences were mainly found for the sainfoin group (S)
- and carob+sainfoin (CaBP+S) groups. As regards the AWCs, no significant
- differences were observed neither in the number of *H. contortus* and *T. colubriformis*.
- No GIN larvae were recovered after pepsin digestion.
- 387 The non-parametric test of Kruskal-Wallis showed that there were statistically
- significant differences in fecundity between the groups (P<0.001). Specifically, the C
- group presented the highest fecundity values, while the CaBP+S group presented the
- lowest ones for both parasite species. Tukey HSD test for *H. contortus* showed that
- 391 the C group differed significantly from CaBP, S and CaBP+S, while for T.
- 392 colubriformis fecundity for CaBP group was also statistically different from CaBP+S
- 393 (Figure 3b).

- When exploring the pathophysiological parameters (i.e. PCV), the analysis of
- 395 variance on repeated measures and also the date by date ANOVA did not show
- significant differences between the groups. Specific values for mean PCV (±SD) on
- 397 DAY 28 of the respective groups C, CaBP, S and CaBP+S were 31.67 (±3.39), 33.00
- 398 (± 4.86) , 31.33 (± 3.61) and 30.50 (± 4.37) .
- 399 The average daily gain (ADG) as calculated for the whole trial duration for (C),
- 400 (CaBP), (S) and (CaBP+S) groups was (mean±s.d.) 122.5(±38.1), 88.2(±39.2),
- 401 $104.6(\pm 11.9)$ and $124.8(\pm 39.7)$ g, respectively and there were no significant
- 402 differences between the groups.

4. Discussion

- 404 The literature contains several in vitro and in vivo studies, conducted on small
- ruminants, which evaluated the anthelmintic effect of tannin-containing plants. Such
- 406 studies first examined temperate forage legumes fed through grazing, as hay, silage or
- 407 pellets. Examples are sainfoin (Hoste et al., 2016; Legendre et al., 2018; Mueller-
- 408 Harvey et al., 2019), sericea lespedeza (Lespedeza cuneata) (Burke et al., 2012a,b;
- 409 Mechineni et al., 2014; Kommuru et al., 2014, 2015), and sulla (Hedysarum
- 410 *coronarium*) (Niezen et al., 1995, 2002). More recently, there has been also a growing
- 411 interest in tannin-containing by-products from the food industry as illustrated by
- 412 studies with hazelnut peels (Corylus avellana fruits) (Desrues et al., 2012; Girard et
- al., 2013), carob pods (Manolaraki et al., 2010; Arroyo-Lopez et al., 2014) and
- 414 browse plants such as *Pistacia lentiscus* (Landau et al., 2010; Manolaraki et al.,
- 415 2010), Ouercus coccifera (Manolaraki et al., 2010) and Salix spp (Mupevo et al.,
- 416 2011).

In the current study, we further explored the *in vivo* anthelmintic effects of carob pod meal since it represents a common feed resource in the Mediterranean region and there was some previous evidence of its anthelmintic (Arroyo-Lopez et al., 2014) and anticoccidial (Saratsis et al., 2016; Legendre et al., 2018) properties. In order to develop a practical implementation tool for carob as dietary intervention, we wanted to identify a) the optimal carob concentration in the feed for bioactivity, b) whether CTs contributed to such an activity and c) whether there were any synergistic effects with other plant sources with different types of CTs (i.e. sainfoin). For all 3 trials a balanced and palatable ration was specifically designed for all animals. This aimed to achieve similar production indexes in all groups and ensured that any observed differences in the effects of parasitism would not stem from quantitative differences in the dietary composition but rather from differences in the bioactive CTs (Coop and Kyriazakis, 1999; Athanasiadou et al., 2008; Hoste et al., 2015). The parasites that served as models for this study (*H. contortus* and *T. colubriformis*) are the most pathogenic and/or prevalent GIN species in European sheep and goats (Charlier et al., 2018). These experiments allowed us to investigate carob-pods efficacy against nematodes in the different anatomical location within the gut, as location can affect the exposure of worms to different CT concentrations (Desrues et al., 2017; Quijada et al., 2018). Results of Trial 1 showed decreases in the mean values of FEC and AWC only in the group fed with the highest concentration of CaBP in the concentrate feed, although not significant. However, fecundity values showed a negative correlation to CaBP concentration in the feed indicating a dose-dependent fecundity suppression effect. The results suggest that carob used in feed at 12% has a potential anthelmintic effect

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and this effect is due mainly to the reduction of female worm fecundity (predominantly in H. contortus) and to a lesser extent to the reduction of establishment and development of the worms. Since H. contortus produce a remarkably high daily egg output compared to T. colubriformis (Besier et al., 2016), we suggest that the reduction in FEC seen in this trial can be attributed to the effect the carob diet had against *H. contortus*. Overall, the results of this trial suggest that the higher the concentration of carob in the ration the higher the anthelmintic activity; this effect that was more evident for *H. contortus*. Unfortunately, there are limitations to the quantity of carob pod meal that can be included in a well balanced ration since carob pods contain high sugar but low protein and lipid concentrations (Priolo et al., 1998; Karabulut et al., 2006). During Trial 2, the main results i) confirmed that CaBP reduced FEC in lambs, as these reductions compared to control values ranged from 20% to 45%, ii) that these reductions in FEC seemed to be mainly due to the lower numbers from the highly prolific H. contortus species and not from T. colubriformis, and that there were no effects on female fecundity of both species and iii) that the anthelmintic effect of CaBP may be attributed to CTs, because a restoration to control values for FEC and Haemonchus worm numbers was observed in the CaBP + PEG group. PEG is a nonnutritive synthetic polymer that is capable of binding and deactivating CTs; it has been used in many animal nutrition studies to increase the intake of CT-containing feeds and to improve protein absorption (Silanikove et al., 1996; Bermingham et al., 2001; Theodoridou et al., 2012). This ability has also been used to test (Brunet et al., 2007, 2008; Debela et al., 2012; Brito et al., 2018) whether any observed in vivo

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anthelmintic activity was linked to the presence of CTs.

Finally, the aim of Trial 3 was to investigate two hypotheses: firstly, that carob CTs generate a stronger anthelmintic effect than sainfoin CTs and secondly, that synergistic effects could be achieved by combining carob with sainfoin. The rationale for these hypotheses is based on the fact that carob and sainfoin contain different types of CTs and that these could target different stages of the GIN life cycle. Carob CTs are highly galloylated prodelphinidins, whereas sainfoin CTs are non-galloylated prodelphinidins. Previous studies found two structural features in CTs that enhance anthelmintic activity in vitro: i) prodelphinidin CTs are more potent than procyanidin CTs and ii) galloylation increases the anthelmintic effect of CTs (Hoste et al., 2016; Kommuru et al., 2014, 2015). Therefore, carob CTs, which have a high prodelphinidin/procyanidin ratio (96.7% prodelphinidins/3.3% procyanidins) and are also highly galloylated (i.e. 41.1% of the flavan-3-ol subunits are galloylated) should produce a stronger anthelmintic effect than sainfoin, as sainfoin CTs have less prodelphinidins (74.8%) and no galloyl groups (N.B. % stands for mole percent within CT molecules; Table 2). There are several important reasons that could explain why the results from Trial 3 did not support either of these hypotheses. Firstly, sainfoin - but not carob - was fed in a pelleted form, while it has been demonstrated previously that the pelleting process has a marked effect on CTs in terms of their analysis (Mueller-Harvey et al., 2019). Table 2 shows that the CT concentrations in sainfoin pellets differed considerably between the two assays (6.5 and 1.7 g CT/100 g DW) in contrast to the carob meal data (5.8 and 7.2 g CT/100g DW). However, we currently do not know whether the pelleting process enhances the anthelmintic activity of CTs or not. Secondly, up to now most attempts to unravel links between CT structural features and anthelmintic effects have employed in vitro assays. Therefore, in vivo feeding trials such as the

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present ones are vital to test the laboratory data. It may turn out that the esterified galloyl groups are not stable in the digestive tract and that the prodelphinidins in carob and sainfoin were the active CTs. Therefore, preliminary conclusions from the Trial 3 data could be that galloylation is unlikely to enhance anthelmintic activity in vivo in terms of H. contortus fecundity or total worm counts and that pelleting of CT-plants might lead to lower FEC. These indications will, however, need rigorous testing in the future. The nutritional and/or anthelmintic properties of sainfoin fed as direct grazing, silage, hay or pellets have been evaluated in both sheep and goats, with promising anthelmintic results when used either alone (Paolini et al., 2005; Heckendorn et al., 2006; Ríos-de Alvarez et al., 2008; Gaudin et al., 2016) or in combination with other CT sources (Girard et al., 2013). Previous results have demonstrated that sainfoin consumption under different forms of preservation can reduce FEC and also reduce female worm fecundity of *H. contortus* (Manolaraki et al., 2010; Arroyo-Lopez et al., 2014) or T. colubriformis (Manolaraki et al., 2010); however, in other studies a lack of effect has been observed (Heckendorn et al., 2006). The issue of the variable results has also been addressed in several reviews (Hoste et al., 2015; Hoste and Niderkorn, 2019). To summarise, the main results of trial 3 for FECs were i) a confirmation of significant reductions of FEC due to the consumption of both CaBP and sainfoin pellets; ii) a temporal increase in the anthelmintic effects of sainfoin but not for CaBP, and iii) no synergistic effects of the combination CaBP + sainfoin. In addition, it would appear that these results can largely be explained by significant effects on female fecundity of both species, but there were only limited effects on the worm populations. No significant effects on AWC were observed for any of the species. On

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the other hand, although the differences were not significant, the percentage of reduction compared to the controls (Group C) for *H. contortus* worm numbers were respectively, for Groups CaBP 35.5%, S 62.1% and CaBP+S 53.5%.

In conclusion, the results of these three trials, which focussed on carob pod meal alone or in combination, raised future research questions regarding what causes the differences in results when different CT-containing resources are used and what is required for a more rational use of CT-containing resources as nutraceutical feeds under farm conditions and in different production systems (Hoste et al., 2015).

Our results confirmed that i) the consumption of CT containing resources can modulate the biology of GINs; ii) that CT were involved in the anthelmintic effects of carob and iii) the concentration in the diet influenced the anthelmintic effects as previously shown in other *in vivo* studies with sericea lespedeza (Shaik et al., 2004, 2006) or sainfoin (Brunet et al., 2007) and iv) different mechanisms appeared to affect the worm population and could explain the reduction of FECs: either a reduced fecundity of female adult worms (see Trial 1 and 3) and /or a reduction of the number of worms (see Trial 2).

The data of these 3 studies also illustrated that results depended on the type of nematode species (abomasal or intestinal species) and/or on the nature of CT resources (in our case carob vs sainfoin) and on the CTs. As stated by Quijada (2015) and Desrues et al. (2016) the quantitative and qualitative differences in CTs appear to influence the anthelmintic activity on the different species of parasitic nematodes.

Our results suggest that, when worm populations are exposed to CTs in the gastrointestinal tract, upon their ingestion by the host the most evident effect recorded is the reduction of female fecundity. Particularly for *H. contortus*, it appears that

fecundity is only affected when the worms are exposed to CTs during maturation (Trial 1 and 3) and not when they are already mature adults (Trial 2). On the other hand, when CaBP was consumed for two weeks by animals in which adult worm populations were already established and patent, the main finding was a significant decrease in *H. contortus* worm counts. Moreover, the current study adds further support to the observation that most of the CT effect is related to abomasal parasite – and not so much to the small intestinal parasite. This is possibly due to the higher CT concentration in the abomasum compared with the rumen and intestines, along with higher prodelphinidin percentage as already shown in studies on the cattle abomasal parasite *Ostertagia ostertagi* (Desrues et al., 2017).

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1 2 Feeding of carob (Ceratonia siliqua) to sheep infected with gastrointestinal nematodes 3 reduces faecal egg counts and worm fecundity 4 Katerina Saratsi^{a,b}, Hervé Hoste^{c,d}, Nikolaos Voutzourakis^a, Nikolaos Tzanidakis^a, 5 Alexandros Stefanakis^a, Stig Milan Thamsborg^e, Irene Mueller-Harvey^f, S., Ioannis 6 Hadjigeorgiou^{b+}, Smaragda Sotiraki^{a+}, 7 8 9 ^aVeterinary Research Institute, Hellenic Agricultural Organization-Demeter, Thermi 10 57001, Thessaloniki, Greece ^bDepartment of Nutritional Physiology and Feeding, Faculty of Animal Science, 12 Agricultural University of Athens, 11855 Athens, Greece 13 ^cINRAE, UMR 1225 IHAP, 23 Chemin des Capelles, Toulouse F-31076, France 14 ^dUniversité de Toulouse, ENVT, 23 Chemin des Capelles, Toulouse F-31076, France 15 ^eDepartment of Veterinary and Animal Sciences, Faculty of Health and Medical 16 Sciences, University of Copenhagen, Copenhagen, Denmark. 17 ^fSchool of Agriculture, Policy and Development, University of Reading, P.O. Box 18 236,1 Earley Gate, Reading RG6 6AT, UK. 19 20 *Corresponding author: Saratsi Katerina, e-mail address: saratsikaterina@yahoo.gr ⁺authors equally supervised the work 22 23 24 25

26 Abstract

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The present study explored the anthelmintic effects of condensed tannins (CT) in carob (Ceratonia siliqua) pods fed to sheep against gastrointestinal nematodes. Three independent in vivo trials tested whether i) carob pod (CaBP)-containing feed had an anthelmintic effect and if yes, which was the optimal concentration in the diet; ii) whether this effect could be attributed to tannins through the polyethylene glycol (PEG) test and iii) whether there were any synergistic effects when combined with another tannin-containing feed (e.g. sainfoin). In all trials 6-month old nematodenaive lambs, experimentally infected with both Haemonchus contortus and Trichostrongylus colubriformis, were used. Faecal egg counts (FEC) were performed regularly and at the end of each trial adult worm counts (AWC) and female worm fecundity were recorded. In trial 1, 35 lambs (five groups of seven lambs) were fed different CaBP concentrations ranging from 0% to 12% w/w. FEC declined up to 39.2% only in the group fed with 12%CaBP, while a declining trend (P<0.06) was demonstrated for the AWC of T. colubriformis, which was associated with the increasing concentration of CaBP in feed. Female worm fecundity was reduced in groups fed CaBP for both parasites, however this was only significant for H. contortus (P<0.001), in a dose dependent manner. In trial 2, four groups of six infected lambs each were used, which received the carob diets CaBP or CaBP+PEG, and the tannin-free diets with or without PEG (C or C+PEG). Results showed that FEC of Groups C, C+PEG, and CaBP+PEG were comparable throughout the trial, while the group receiving only CaBP showed lower FEC from DAY 25 onwards. AWC showed a reduction (67.7%) only for *H. contortus* (P<0.03). Reversal of the anthelmintic effect of CaBP after PEG administration suggested that CT contributed to the anthelmintic action. However, no effect of CaBP was observed on T.

colubriformis AWC and on female worm fecundity for both species. Finally, for trial
3 four groups of six lambs each received a diet based on CaBP, sainfoin (S) or a
combination (CaBP+S) and were compared to a control (C) diet of lucerne. On DAY
37 FEC values in groups CaBP+S and S tended to be lower compared to the two other
groups (C, CaBP), while for AWCs no significant differences were observed for both
parasites. The fecundity of <i>H. contortus</i> and <i>T. colubriformis</i> demonstrated significant
differences between the treated and control groups, with lower values in the animals
receiving CaBP+S. Overall, the results supported the hypothesis that carob had an
anthelmintic effect due to its CT, but there was no clear indication of a synergistic
effect with sainfoin.
Keywords: Carob, Sainfoin, Haemonchus contortus, Trichostrongylus colubriformis,
gastrointestinal nematodes, sheep, feed additives

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1. Introduction

Infections by gastrointestinal nematodes (GIN) affect both health and welfare of grazing ruminants, causing anorexia, impaired digestion and nutrient absorption with related production losses, diarrhoea, anaemia and even death (Perry and Randolph, 1999; Stear et al., 2007; Hoste et al., 2016). Currently, the control of these parasites relies on repeated dosing with commercial anthelmintic drugs. However, the development of anthelmintic resistance in worm populations against one or multiple classes of anthelmintic has become a serious problem in several regions of the world, making it increasingly difficult to control parasitic infections (Kaplan, 2020). At the same time the increasing concerns of consumers about the presence of drug residues in foods and in the environment (McKellar, 1997) have stimulated the search for alternative or complementary solutions (Hoste and Torres-Acosta, 2011) within the context of organic farming and sustainable agriculture (Charlier et al., 2018). Such alternatives include the use of bioactive plants with anthelmintic properties. Many results indicate that such resources, because of the presence of plant secondary metabolites (PSMs), might help to interfere with the biology of key-stages during nematode cycle and to reduce the consequences of GIN infections in grazing ruminants. Particular attention has been given to plants containing condensed tannins (CT) and some related polyphenols (see reviews by Terrill et al., 2012; Hoste et al., 2015, 2016). Based on previous studies, the need to explore new plant resources to develop non-drug-based strategies for the integrated control of nematode parasites in grazing ruminants has recently become a research priority in livestock production, especially in ruminant breeding as also reviewed by Morgan et al, 2020.

Small ruminants (sheep and goats) are a major component of the dairy sector in the Mediterranean basin (Hadjigeorgiou et al., 2005). Sheep and goat production often occupy marginal lands that are unsuitable for crop production but are rich in local plants, such as rangeland vegetation, which can be exploited by animals as a feed resource (Frutos et al., 2008; Méndez-Ortiz et al., 2018). Many rangeland plants also contain PSMs, such as tannins (Papachristou et al., 2005) and several in vitro and in vivo studies have evaluated their anthelmintic effects against GINs of small ruminants (Manolaraki et al., 2010; Moreno-Gonzalo et al., 2012, 2013a,b, 2014; Arroyo-Lopez et al., 2014; Silva Soares et al., 2018). Overall, CTs have been shown to directly or indirectly interfere with the life cycle of several GINs and, therefore, CT-containing plants, which also include many legumes, are proving to be beneficial nutritional resources. However, a high degree of variability with respect to their anthelmintic activity has also been recorded. Besides the total tannin concentration in ruminant diets, recent studies have demonstrated that CT molecular composition or structural characteristics can also affect anthelmintic activity (Mueller-Harvey et al., 2019). Carob (Ceratonia siliqua) and sainfoin (Onobrychis viciifolia) are both resources of the Fabaceae family and contain CTs. Carob is a leguminous tree that is widely cultivated in the Mediterranean area. It is an important species both for economic and environmental reasons (Batlle and Tous, 1997). Carob pods (fruits) are mostly used in the food industry; pulp accounts for 90% by pod weight and seeds for 10%. They contain high sugar (48–56%), but low protein (3–4%) and lipid concentrations (0.4– 0.8%) (Marakis, 1996; Batlle and Tous, 1997). Moreover, ripe carob pods contain high concentrations of CTs (16-20% w/w DM) (Bravo et al., 1994; Batlle and Tous,

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1997). This has been debated by Priolo et al. (2000, 2002) who claimed that the pods

have low content of CTs, but with exceptionally high biological activity. Silanikove et

al. (2006) have demonstrated that the yield of CTs is considerably affected by the extraction method applied (from 5.0% with acidic methanol to 17.2% with urea-buffer solution), suggesting that carob pods are a rich source of CTs. The high CT concentration in by-products from carob pod processing justifies researching its value as a feed additive with possible effect against GIN species. Sainfoin, which can be found especially in southern parts of Europe, has been the subject of renewed interest because of its beneficial effects in the context of agroecology (Hayot Carbonero et al., 2011), its beneficial impact on ruminant production and the environment and its potential antiparasitic effects on small ruminants (Manolaraki et al., 2010; Hoste et al., 2015; Saratsis et al., 2016; Mueller-Harvey et. al., 2019). In vitro studies have shown that sainfoin extracts have a dosedependent effect against different GIN species (Brunet et al., 2007; Manolaraki et al., 2010; Novobilsky et al., 2013). Moreover, in vivo anthelmintic effects have also been described in sheep and/or goats fed with sainfoin; i.e. 42-68% reduction in parasitic egg excretion, which was associated with a 17.6% decrease in female worm fecundity and a 45% decrease in worm numbers for *Haemonchus contortus* (Arroyo-Lopez et al., 2014).

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The present study, therefore, sought to explore the anthelmintic effects of feeding regimes employing two CT-containing plant resources that may be relevant for Mediterranean conditions. These were offered to lambs either alone or in combination to evaluate their efficacies against two GIN species (*H. contortus* and *Trichostrongylus colubriformis*). The specific objectives were to explore whether a) the anthelmintic effect of carob in the feed is dose dependent (Trial 1), b) this anthelmintic effect is associated with tannins by using polyethylene glycol (PEG) as a

tannin-inhibitor (Trial 2), and c) there are any synergistic effects between carob and sainfoin feeds (Trial 3).

2. Materials and Methods

2.1. Stabling and animals

The experiments were carried out at the Asomaton Research Station of HAO Demeter on the island of Crete, Greece. The animal barn was of an open-sided shed type, with straw bedding. During the whole study the animals were kept indoors with each group in a separate pen of approximately 10 m² and 10 m² open yard. The study included three trials with lambs belonging to the local "Sfakion" breed. In order to achieve uniformity of the experimental animals, all lambs included were female, 6-month-old with a comparable body weight (BW), which was within the normal BW range of the breed (22-30 kg) at the specific age. The lambs were raised indoors under helminth-free conditions. Fourteen days before the start of each trial, they were drenched with albendazole at the higher commercially recommended dose (ALBENDAZOLE Drench, PROVET, 7.5 mg/kg) and they tested negative by faecal egg counts at the start of each trial. No anthelmintic resistance was previously recorded for this specific flock.

2.3. Infective larvae

Third-stage infective larvae of H. contortus and T. colubriformis strains, susceptible to all classes of anthelmintic drugs, had been cultured from faeces of monospecifically infected donor sheep. Larvae were recovered using the Baermann technique and then stored for 1-2 months at 4° C until use.

2.4. Tannin-containing plant resources

174 Carob pods (after removal of the seeds) were locally purchased and offered as crushed 175 flour meal incorporated in the concentrate feed supplement. Sainfoin pellets (Perly 176 cultivar, 3rd cut) were provided by Multifolia (Viapres le Petit 10380, France) as part 177 of the Research project CARES.

2.5. Tannin concentration and composition

Tannin concentrations and compositions were determined in triplicate using two different assays, i.e. the acetone-butanol-HCl assay and the thiolytic degradation with benzyl mercaptan. Both techniques were applied in order to ensure a comprehensive analysis since it was previously demonstrated that, depending on the types of CTs, the acetone-HCl-butanol assay can give higher CT concentrations than the thiolysis assay.

The acetone-HCl-butanol assay was carried out as previously described by Grabber et al. (2013) and Desrues et al. (2017).

The thiolysis reaction was carried out with benzyl mercaptan (Gea et al., 2011;

Ropiak et al., 2016), the reaction products were identified by HPLC-MS analysis (Williams et al., 2014; Desrues et al., 2017) and quantified based on peak areas at 280 nm (Gea et al., 2011; Ropiak et al., 2016). This provided information on CT concentration (g CT/100 g DW), CT size (in terms of mean degree of polymerisation, mDP), molar percentages of prodelphinidins (PD) and procyanidins (PC) within CTs, and molar percentages of *cis*- vs *trans*- flavan-3-ol subunits (Ropiak et al., 2016).

2.6. Experimental design

All diets offered to the animals during the experimental period (with or without the tannin sources) were formulated to meet the nutrient requirements of the animals (NRC, 2007) and the total rations were always iso-nitrogenous and iso-energetic as

well as balanced for crude fibre, Ca, P and Ca/P ratio (Suppl Table 1). Animals had access to clean water at all times. The animals' appetite was assessed and feed consumption (as feed offered minus refusals) was recorded on a daily basis by the farm manager.

2.6.1. *Trial* 1

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- To determine the anthelmintic effect of carob pod meal and to define the optimal concentration in a sheep ration, a subset of 35 lambs were randomly allocated to 5
- 204 groups (n=7 lambs/diet) (Table 1).
- 205 Carob meal (CaBP) was offered as feed supplement, at increasing rates of 0%, 3%,
- 206 6%, 9% and 12% (g CaBP/100g DM) of the total ration. The highest proportion, of
- 207 carob meal contributed to concentrate feed was set to 12% (due to its poor energy and
- 208 protein contents) in order to enable formulating a ration, which could cover the
- 209 nutritional requirements of lambs.
- Feeding the experimental diets started 2 weeks prior (D14) to experimental infection
- with nematode larvae (D0) in order for the animals to adapt to the feed.
- 212 On DAY 0, all lambs in groups (i) to (v) were infected with a single dose of 12.000
- 213 3rd stage larvae (L3) of *H. contortus* and 12.000 L3 of *T. colubriformis*. At the end of
- 214 the experimental period (D49), all lambs were euthanised by injection of a massive
- 215 dose of pentobarbital (Dolethal®).

2.6.2. Trial 2

- Four groups of 6 lambs were included in a two-factorial trial (diet and PEG-addition).
- 218 Two groups were offered CaBP as feed supplement at the rate of 12% in the total
- 219 ration, and two groups remained on standard diet (Table 1). Half of the lambs in each
- 220 diet group were offered PEG (Polyethylene Glycol 4000, Fisher Scientific USA)

221 orally (60 g/lamb diluted in 200 ml water) on a daily basis after being allocated into 222 groups. 223 On D0, all lambs were experimentally infected with 8.000 L3 of H. contortus and 224 16.000 L3 of T. colubriformis. On D21, after parasite infection was confirmed by positive faecal examination, the animals were allocated into 4 groups of 6 lambs each, 225 226 according to the experimental diets. On D37 they were euthanised as described above. 227 **2.6.3.** *Trial 3* 228 To determine the possible synergistic anthelmintic effects between 2 CT-containing 229 resources namely carob (C. siliqua) and sainfoin (O. viciifolia), 4 groups of 6 lambs 230 were included in a two-factorial design (Table 1). 231 On D-14 each group of lambs received the allocated diet, containing i) carob meal 232 (CaBP) alone ii) sainfoin (S) pellets; iii) a combination of carob meal and sainfoin 233 pellets (CaBP+S) while iv) a control group (C), received an isoproteic diet based on 234 lucerne. Carob was offered as a feed supplement at the rate of 12% in the total ration. 235 Sainfoin was offered as pellets representing 35% of the total ration. On D0 all lambs 236 were infected with a single dose of 12.000 L3 of H. contortus and 12.000 L3 of T. 237 colubriformis. At the end of the experimental period (D37), all lambs were euthanised 238 as previously described. 239 2.7. Pathophysiological parameters 240 Individual blood samples were collected once weekly (from D0 to D49) during Trial 1 241 and once every two weeks (from D0 to D28) during Trial 3, by jugular venipuncture

into heparinized tubes (BD Vacutainer®, UK) to determine the packed cell volume

(PCV), as an indicator of anaemia, according to the micro-haematocrit method. In

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Trial 2 due to its short duration, the recording of PCV values was not included in the design.

2.8. Parasitological parameters

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(10%) of the total volume.

Individual faecal samples were collected weekly directly from the rectum, during the 1st and 3rd trial, and twice weekly during the 2nd trial in order to determine faecal egg counts (FEC) using a modified McMaster technique (Roepstorff and Nansen, 1998). FEC data were expressed as eggs per gram of faeces (EPG). At necropsy, the abomasa and the first 12 meters of small intestine were separated, ligated, rapidly removed and immediately processed to collect the adult worms from the luminal contents. For the intra-mucosal larvae, pepsin digestion was applied both on the abomasum and intestinal mucosa (MAFF, 1986). After 4h incubation at 37°C the larvae were collected. After storage in 10% alcohol, worm counts were performed according to a 10% aliquot technique (MAFF, 1986). Morphological identification of worm stages, sex and species were conducted using standard procedures (MAFF, 1986). The fecundity of female worms was measured on 10 worms per lamb. For T. colubriformis, eggs were counted directly in utero after clearing in 85% lactic acid solution. All egg counts were performed under a microscope set at 10 times magnification (total 100 ×). For H. contortus, the fecundity was determined using the method described by Kloostermann et al. (1978). Briefly, the worms were soaked for 5 min in a large volume of distilled water, before being placed individually in microtubes with 1000 µl of 0.125% hypochlorite concentration solution and kept at room temperature for 20 minutes. Treatment resulted in female worms disintegrating thus enabling the direct counting of eggs under a stereo-microscope using an aliquot

2.9. Statistical analyses

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The data of FEC and adult worm counts (AWC) were log10(x+1) transformed prior to analysis. For the FEC values, comparison of all groups was first performed using an analysis of variance (ANOVA) with time as repeated measurement. Then, the comparison of results to the control values were carried out date by date, using oneway ANOVA completed by the post-hoc Bonferroni test for pairwise comparisons. Group means of AWC were compared by one-way ANOVA (Trial 1) or two-way ANOVA (Trial 2: CaBP +/- and PEG +/-; Trial 3: CaBP +/- and sainfoin +/-). Regarding the fecundity of female worms, the Shapiro-Wilk Test of normality, which is more appropriate for small sample sizes, was used. In cases where the data deviated significantly (P<0.05) from a normal distribution (Trial 1 and 3 for both parasite species and Trial 2 for T. colubriformis) the appropriate test to check the difference of fecundity between the groups, which is the non-parametric test of Kruskal-Wallis, was used. Where the dependent variable was normally distributed (P>0.05) the parametric test of one-way ANOVA (H. contortus of Trial 2) was used. Additionally, for the Trial 1, the model of linear regression was used, in order to be investigated if there was a negative correlation between the variables "percentages of carob" and "fecundity of female worms" for both parasite species (H. contortus and T. colubriformis). Finally, the Tukey HSD test was used for data of trial 3, in order to investigate statistically significant differences between groups.

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2.10. Ethical considerations

292 The study was carried out in compliance with the national animal welfare regulations.

All statistical analyses were performed using the SyStat SPSS 9.0 Software.

293 All trials took place in a Research Station of the Veterinary Research Institute. The

experimental protocol was approved by the responsible institutional committee (VRI Committee for Approval of Experimental protocols as appointed at 26/5/2014, Decision nr 972). Euthanasia was performed in a humane manner according to EU regulations.

3. Results

The CT concentrations and compositions are presented in Table 2. The HBA assay yielded similar CT concentrations for both plant materials, whereas the thiolysis assay generated lower CT concentrations for the sainfoin pellets. The thiolysis assay revealed that: i) both carob and sainfoin CTs consisted mainly of prodelphinidins, 96.7 and 74.7 mole percentages, respectively; ii) carob CTs were highly galloylated (i.e. 41.1% of flavan-3-ol subunits are galloylated), but sainfoin CTs did not contain any esterified galloyl groups; iii) carob CTs were characterised by a relatively high average molecular weight (mDP = 31.1), whereas sainfoin CTs had an mDP value of 11.5.

3.1. Trial 1

- 311 The results of Trial 1 are shown in Table 3 and Figure 1
- The analyses of FEC, based on the ANOVA on Repeated Measures from D21 to D49, showed an overall non-significant difference between groups, but significant difference over time (between days of sampling). Meanwhile, the date-by-date ANOVA of FEC showed no significant differences between groups, whatever the date, as well as no dose effect. Reduction in FEC, up to 39.2% on DAY 49 as compared to controls, was observed only for the group fed with the highest concentration of carob meal.

- For *H. contortus*, the AWC declined in the groups receiving the highest concentration
- of carob meal but this effect was not statistically significant (P= 0.964). In contrast,
- 321 there was a declining trend (P<0.06) for the numbers of T. colubriformis with
- increasing carob concentration.
- 323 The fecundity values showed significant differences (15.6%-59.3% lower than
- 324 0%CaBP respectively from the lowest to the highest CaBP concentration) between
- groups for *H. contortus* demonstrating a dose dependent effect (P<0.05).
- 326 The Box plot (Figure 1b) for *H. contortus* fecundity suggests that worms from the
- 327 0%CaBP group tended to be more fecund than other CaBP groups and there may be
- 328 some degree of fecundity discrepancy between CaBP groups. This trend was
- 329 confirmed with the non-parametric test of Kruskal-Wallis, which showed that there
- were statistically significant differences in fecundity between the groups (P<0.001).
- 331 More specifically, fecundity was statistically significantly greater for 0% CaBP group
- than the other CaBP groups. On the other hand, regarding *T. colubriformis* fecundity,
- 333 there was no statistically significant difference between the groups (P=0.128).
- 334 However, the model of linear regression, which was implemented and was
- statistically significant (P<0.05), showed a negative correlation between the variables
- "group" and "fecundity" for both parasite species.
- No GIN larvae were recovered after pepsin digestion.
- Mean PCV values (±SD) for groups 0%CaBP, 3%CaBP, 6%CaBP, 9%CaBP, and
- 339 12%CaBP on the last day of the trial were 25.29 (± 5.96), 23.00 (± 5.72), 21.00
- (± 6.32) , 23.00 (± 5.89) and 24.00 (± 5.00) respectively. No significant differences were
- 341 found between the groups in PCV.
- 342 Average daily gain (ADG) as calculated for the whole trial duration for 0%CaBP,
- 343 3%CaBP, 6%CaBP, 9%CaBP and 12%CaBP groups was (mean±s.d.) 69.2 g (±31.0),

 $61.5(\pm 36.1)$, $68.7(\pm 33.0)$, $74.8(\pm 37.5)$ and $64.4(\pm 32.9)$ g respectively, which yielded 344 345 no significant differences between the groups.

3.2.Trial 2

346 347 The results of Trial 2 are presented in Table 4 and Figure 2. 348 The Repeated Measurements Analyses of FEC showed an overall statistical difference 349 (P<0.001) between the 4 groups. The date-by-date ANOVA of FEC indicated that differences were most prominent on DAY 29 (significant statistical differences, 350 351 P<0.02) and then on DAY 33 (trend, P<0.07). Specifically, the values of the C, 352 C+PEG, CaBP+PEG groups were comparable throughout the trial, while the group 353 receiving only carob (CaBP) showed consistently lower FEC starting from DAY 25 354 until the last day of the experiment. It was evident that the effect of carob on FEC was 355 nullified by PEG. 356 Results on AWC, showed reduction only for *H. contortus* (P<0.03) resulting in an 357 overall statistical difference between the 4 groups, since the lowest worm counts were 358 found for the CaBP group. Especially, for *H. contortus*, a reduction of approximately 359 65% was observed in the carob group compared to the control. The AWC in the 360 CaBP+PEG group were similar to the other 2 control groups showing no reduction in 361 worm population. On the other hand, no effect of carob was observed on T. 362 colubriformis worm counts. 363 No effect of carob on female fecundity was also observed, irrespective of the parasite 364 species. Both control and carob groups showed comparable levels of female fecundity 365 for the two parasite species. The Box plot in Figure 2b showed that the range of 366 fecundity of H. contortus for CaBP group was greater than for C, C+PEG and CaBP+PEG groups and the interquartile range (middle 50% of the records) was lower 367 368 on the fecundity scale in the CaBP group than in the other groups.

- No GIN larvae were recovered after pepsin digestion.
- 370 The average daily gain (ADG) of lambs as calculated for the whole trial duration for
- 371 (C), (C+PEG), (CaBP) and (CaBP+PEG) groups was $51.8(\pm 30.1)$ ($\pm s.d.$), $69.8(\pm 19.9)$,
- $60.8(\pm 29.3)$ and $40.5(\pm 25.6)$ g, respectively, which resulted in no significant
- differences between the groups.

374 **3.3.Trial 3**

- 375 The results of Trial 3 are shown in Table 5 and Figure 3.
- 376 The FEC values of all experimental groups remained at very low levels up to DAY
- 377 21. The overall repeated analyses based on 3 dates of the patent phase (DAY 21,
- 378 DAY 28, DAY 37) showed a trend for differences (P<0.07) between groups. The
- 379 results of the date-by-date ANOVA test did not show difference on DAY 21 and on
- DAY 28, while on DAY 37, the values of FEC in groups CaBP+S and S tended to be
- reduced (P< 0.06) compared to the two other groups. When compared to the control
- 382 values of FEC, the reductions in the 3 treated groups ranged from 44.6% to
- approximately 86 %. These differences were mainly found for the sainfoin group (S)
- and carob+sainfoin (CaBP+S) groups. As regards the AWCs, no significant
- differences were observed neither in the number of *H. contortus* and *T. colubriformis*.
- No GIN larvae were recovered after pepsin digestion.
- 387 The non-parametric test of Kruskal-Wallis showed that there were statistically
- significant differences in fecundity between the groups (P<0.001). Specifically, the C
- group presented the highest fecundity values, while the CaBP+S group presented the
- lowest ones for both parasite species. Tukey HSD test for *H. contortus* showed that
- 391 the C group differed significantly from CaBP, S and CaBP+S, while for T.
- 392 colubriformis fecundity for CaBP group was also statistically different from CaBP+S
- 393 (Figure 3b).

- When exploring the pathophysiological parameters (i.e. PCV), the analysis of
- 395 variance on repeated measures and also the date by date ANOVA did not show
- significant differences between the groups. Specific values for mean PCV (±SD) on
- 397 DAY 28 of the respective groups C, CaBP, S and CaBP+S were 31.67 (±3.39), 33.00
- 398 (± 4.86) , 31.33 (± 3.61) and 30.50 (± 4.37) .
- 399 The average daily gain (ADG) as calculated for the whole trial duration for (C),
- 400 (CaBP), (S) and (CaBP+S) groups was (mean±s.d.) 122.5(±38.1), 88.2(±39.2),
- 401 $104.6(\pm 11.9)$ and $124.8(\pm 39.7)$ g, respectively and there were no significant
- 402 differences between the groups.

4. Discussion

- 404 The literature contains several in vitro and in vivo studies, conducted on small
- ruminants, which evaluated the anthelmintic effect of tannin-containing plants. Such
- 406 studies first examined temperate forage legumes fed through grazing, as hay, silage or
- 407 pellets. Examples are sainfoin (Hoste et al., 2016; Legendre et al., 2018; Mueller-
- 408 Harvey et al., 2019), sericea lespedeza (Lespedeza cuneata) (Burke et al., 2012a,b;
- 409 Mechineni et al., 2014; Kommuru et al., 2014, 2015), and sulla (Hedysarum
- 410 *coronarium*) (Niezen et al., 1995, 2002). More recently, there has been also a growing
- 411 interest in tannin-containing by-products from the food industry as illustrated by
- 412 studies with hazelnut peels (Corylus avellana fruits) (Desrues et al., 2012; Girard et
- al., 2013), carob pods (Manolaraki et al., 2010; Arroyo-Lopez et al., 2014) and
- 414 browse plants such as *Pistacia lentiscus* (Landau et al., 2010; Manolaraki et al.,
- 415 2010), Ouercus coccifera (Manolaraki et al., 2010) and Salix spp (Mupevo et al.,
- 416 2011).

In the current study, we further explored the *in vivo* anthelmintic effects of carob pod meal since it represents a common feed resource in the Mediterranean region and there was some previous evidence of its anthelmintic (Arroyo-Lopez et al., 2014) and anticoccidial (Saratsis et al., 2016; Legendre et al., 2018) properties. In order to develop a practical implementation tool for carob as dietary intervention, we wanted to identify a) the optimal carob concentration in the feed for bioactivity, b) whether CTs contributed to such an activity and c) whether there were any synergistic effects with other plant sources with different types of CTs (i.e. sainfoin). For all 3 trials a balanced and palatable ration was specifically designed for all animals. This aimed to achieve similar production indexes in all groups and ensured that any observed differences in the effects of parasitism would not stem from quantitative differences in the dietary composition but rather from differences in the bioactive CTs (Coop and Kyriazakis, 1999; Athanasiadou et al., 2008; Hoste et al., 2015). The parasites that served as models for this study (*H. contortus* and *T. colubriformis*) are the most pathogenic and/or prevalent GIN species in European sheep and goats (Charlier et al., 2018). These experiments allowed us to investigate carob-pods efficacy against nematodes in the different anatomical location within the gut, as location can affect the exposure of worms to different CT concentrations (Desrues et al., 2017; Quijada et al., 2018). Results of Trial 1 showed decreases in the mean values of FEC and AWC only in the group fed with the highest concentration of CaBP in the concentrate feed, although not significant. However, fecundity values showed a negative correlation to CaBP concentration in the feed indicating a dose-dependent fecundity suppression effect. The results suggest that carob used in feed at 12% has a potential anthelmintic effect

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and this effect is due mainly to the reduction of female worm fecundity (predominantly in H. contortus) and to a lesser extent to the reduction of establishment and development of the worms. Since H. contortus produce a remarkably high daily egg output compared to T. colubriformis (Besier et al., 2016), we suggest that the reduction in FEC seen in this trial can be attributed to the effect the carob diet had against *H. contortus*. Overall, the results of this trial suggest that the higher the concentration of carob in the ration the higher the anthelmintic activity; this effect that was more evident for *H. contortus*. Unfortunately, there are limitations to the quantity of carob pod meal that can be included in a well balanced ration since carob pods contain high sugar but low protein and lipid concentrations (Priolo et al., 1998; Karabulut et al., 2006). During Trial 2, the main results i) confirmed that CaBP reduced FEC in lambs, as these reductions compared to control values ranged from 20% to 45%, ii) that these reductions in FEC seemed to be mainly due to the lower numbers from the highly prolific H. contortus species and not from T. colubriformis, and that there were no effects on female fecundity of both species and iii) that the anthelmintic effect of CaBP may be attributed to CTs, because a restoration to control values for FEC and Haemonchus worm numbers was observed in the CaBP + PEG group. PEG is a nonnutritive synthetic polymer that is capable of binding and deactivating CTs; it has been used in many animal nutrition studies to increase the intake of CT-containing feeds and to improve protein absorption (Silanikove et al., 1996; Bermingham et al., 2001; Theodoridou et al., 2012). This ability has also been used to test (Brunet et al., 2007, 2008; Debela et al., 2012; Brito et al., 2018) whether any observed in vivo

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anthelmintic activity was linked to the presence of CTs.

Finally, the aim of Trial 3 was to investigate two hypotheses: firstly, that carob CTs generate a stronger anthelmintic effect than sainfoin CTs and secondly, that synergistic effects could be achieved by combining carob with sainfoin. The rationale for these hypotheses is based on the fact that carob and sainfoin contain different types of CTs and that these could target different stages of the GIN life cycle. Carob CTs are highly galloylated prodelphinidins, whereas sainfoin CTs are non-galloylated prodelphinidins. Previous studies found two structural features in CTs that enhance anthelmintic activity in vitro: i) prodelphinidin CTs are more potent than procyanidin CTs and ii) galloylation increases the anthelmintic effect of CTs (Hoste et al., 2016; Kommuru et al., 2014, 2015). Therefore, carob CTs, which have a high prodelphinidin/procyanidin ratio (96.7% prodelphinidins/3.3% procyanidins) and are also highly galloylated (i.e. 41.1% of the flavan-3-ol subunits are galloylated) should produce a stronger anthelmintic effect than sainfoin, as sainfoin CTs have less prodelphinidins (74.8%) and no galloyl groups (N.B. % stands for mole percent within CT molecules; Table 2). There are several important reasons that could explain why the results from Trial 3 did not support either of these hypotheses. Firstly, sainfoin - but not carob - was fed in a pelleted form, while it has been demonstrated previously that the pelleting process has a marked effect on CTs in terms of their analysis (Mueller-Harvey et al., 2019). Table 2 shows that the CT concentrations in sainfoin pellets differed considerably between the two assays (6.5 and 1.7 g CT/100 g DW) in contrast to the carob meal data (5.8 and 7.2 g CT/100g DW). However, we currently do not know whether the pelleting process enhances the anthelmintic activity of CTs or not. Secondly, up to now most attempts to unravel links between CT structural features and anthelmintic effects have employed in vitro assays. Therefore, in vivo feeding trials such as the

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present ones are vital to test the laboratory data. It may turn out that the esterified galloyl groups are not stable in the digestive tract and that the prodelphinidins in carob and sainfoin were the active CTs. Therefore, preliminary conclusions from the Trial 3 data could be that galloylation is unlikely to enhance anthelmintic activity in vivo in terms of H. contortus fecundity or total worm counts and that pelleting of CT-plants might lead to lower FEC. These indications will, however, need rigorous testing in the future. The nutritional and/or anthelmintic properties of sainfoin fed as direct grazing, silage, hay or pellets have been evaluated in both sheep and goats, with promising anthelmintic results when used either alone (Paolini et al., 2005; Heckendorn et al., 2006; Ríos-de Alvarez et al., 2008; Gaudin et al., 2016) or in combination with other CT sources (Girard et al., 2013). Previous results have demonstrated that sainfoin consumption under different forms of preservation can reduce FEC and also reduce female worm fecundity of *H. contortus* (Manolaraki et al., 2010; Arroyo-Lopez et al., 2014) or T. colubriformis (Manolaraki et al., 2010); however, in other studies a lack of effect has been observed (Heckendorn et al., 2006). The issue of the variable results has also been addressed in several reviews (Hoste et al., 2015; Hoste and Niderkorn, 2019). To summarise, the main results of trial 3 for FECs were i) a confirmation of significant reductions of FEC due to the consumption of both CaBP and sainfoin pellets; ii) a temporal increase in the anthelmintic effects of sainfoin but not for CaBP, and iii) no synergistic effects of the combination CaBP + sainfoin. In addition, it would appear that these results can largely be explained by significant effects on female fecundity of both species, but there were only limited effects on the worm populations. No significant effects on AWC were observed for any of the species. On

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the other hand, although the differences were not significant, the percentage of reduction compared to the controls (Group C) for *H. contortus* worm numbers were respectively, for Groups CaBP 35.5%, S 62.1% and CaBP+S 53.5%.

In conclusion, the results of these three trials, which focussed on carob pod meal alone or in combination, raised future research questions regarding what causes the differences in results when different CT-containing resources are used and what is required for a more rational use of CT-containing resources as nutraceutical feeds under farm conditions and in different production systems (Hoste et al., 2015).

Our results confirmed that i) the consumption of CT containing resources can modulate the biology of GINs; ii) that CT were involved in the anthelmintic effects of carob and iii) the concentration in the diet influenced the anthelmintic effects as previously shown in other *in vivo* studies with sericea lespedeza (Shaik et al., 2004, 2006) or sainfoin (Brunet et al., 2007) and iv) different mechanisms appeared to affect the worm population and could explain the reduction of FECs: either a reduced fecundity of female adult worms (see Trial 1 and 3) and /or a reduction of the number of worms (see Trial 2).

The data of these 3 studies also illustrated that results depended on the type of nematode species (abomasal or intestinal species) and/or on the nature of CT resources (in our case carob vs sainfoin) and on the CTs. As stated by Quijada (2015) and Desrues et al. (2016) the quantitative and qualitative differences in CTs appear to influence the anthelmintic activity on the different species of parasitic nematodes.

Our results suggest that, when worm populations are exposed to CTs in the gastrointestinal tract, upon their ingestion by the host the most evident effect recorded is the reduction of female fecundity. Particularly for *H. contortus*, it appears that

fecundity is only affected when the worms are exposed to CTs during maturation (Trial 1 and 3) and not when they are already mature adults (Trial 2). On the other hand, when CaBP was consumed for two weeks by animals in which adult worm populations were already established and patent, the main finding was a significant decrease in *H. contortus* worm counts. Moreover, the current study adds further support to the observation that most of the CT effect is related to abomasal parasite – and not so much to the small intestinal parasite. This is possibly due to the higher CT concentration in the abomasum compared with the rumen and intestines, along with higher prodelphinidin percentage as already shown in studies on the cattle abomasal parasite *Ostertagia ostertagi* (Desrues et al., 2017).

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Table 1: Experimental design (CaBP=Carob; CaBP+PEG=Carob+PEG; C=Control; C+PEG=Control+PEG; CaBP+S=Carob+Sainfoin; S=Sainfoin)

Trial	Groups	Lambs/ group	Mean BW at start ±s.d. (kg)	Day feeding started	Infection Day	Inoculation dose (L3)	Day trial ended
	CaBP 0%		25.8±1.1			12,000 H.	
	CaBP 3%		26.2 ± 3.7		0	contortus	
1	CaBP 6%	7	25.2 ± 2.6	-14		&	49
	CaBP 9%		27.1±3.5			12,000 T. colubriformis	
	CaBP 12%		26.4±2.8				
	С	6	26.4±2.7			8,000 L3 <i>H</i> .	
2	C+PEG		26.5±2.6	21	0	contortus	27
2	CaBP (12%)		26.3 ± 2.4	21	0	&	37
	CaBP+PEG		27.0±1.7			16,000 L3 T. colubriformis	
	С		27.0±3.1			12,000 L3 <i>H</i> .	
2	CaBP (12%)	6	27.1±2.5	1 /	0	contortus	27
3	CaBP+S		27.1±2.9	-14	0	&	37
	S		26.8±3.2			12,000 L3 <i>T.</i> colubriformis	

Table 2. Condensed tannin concentrations (expressed as g CT/100 g DW) measured either with the acetone-HCl-butanol or the thiolysis assays as well as tannin compositions in the two different feeds [abbreviations: % refers to molar percentages of galloylation, prodelphinidins (PD), procyanidins (PC), *cis*- or *trans*- flavan-3-ol subunits; mean degree of polymerisation (mDP)].

ND: non detected

	% galloylation	PD/PC	Tannins (acetone- HCl/butanol)	Tannins (thiolysis)	mDP	cis/trans-flavan-3-ols
Carob meal	41.1 (±0.6)	96.7/3.3 (±0.1)	5.84 (±0.2)	7.20 (±0.0)	31.2 (±0.1)	45.9/54.1 (±0.0)
Sainfoin pellets ^a	ND	74.8/25.2 (±0.5)	6.50 (±0.3)	1.70 (±0.1)	11.5 (±0.3)	85.3/14.7 (±0.1)

^aThe same sainfoin pellets were used in another study (Quijada et al., 2018) and the data are reported here for comparison purposes.

Table 3.

Trial 1: Effect of diet regimes containing different concentration of Carob (CaBP) on adult worms recovered at necropsy in the different experimental lamb groups. Adult worm counts (AWC) shown as arithmetic mean of adult worms (and SD in brackets) per group fed different amount of Carob pod meal (CaBP = Carob).

Treatment		H. contortus			T. colubriformis	
Group	Female	Male	Total	Female	Male	Total
00/ CaDD	2,777	2,331	5,109	2,063	1,097	3,160
0% CaBP	$(\pm 1,579)$	$(\pm 1,382)$	$(\pm 2,802)$	(± 534)	(± 471)	(±944)
3% CaBP	2,789	2,006	4,794	2,493	1,163	3,656
376 Cabr	$(\pm 1,606)$	$(\pm 1,117)$	$(\pm 2,709)$	(± 986)	(± 568)	$(\pm 1,461)$
6% CaBP	3,584	2,570	6,154	2,514	903	3,417
0 / 0 Cabi	$(\pm 1,570)$	$(\pm 1,178)$	$(\pm 2,595)$	(±497)	(± 676)	$(\pm 1,073)$
9% CaBP	3,029	2,799	5,827	1,910	633	2,543
970 Cabi	$(\pm 1,385)$	$(\pm 1,415)$	$(\pm 2,692)$	$(\pm 1,043)$	(± 427)	$(\pm 1,418)$
12% CaBP	2,160	2,039	4,199	1,550	944	2,494
12 /0 Cabr	$(\pm 1,362)$	$(\pm 1,212)$	$(\pm 2,422)$	(± 801)	(± 630)	$(\pm 1,416)$

Table 4. Trial 2: Effect of PEG intake on adult worms recovered at necropsy in the different groups of lambs fed with carob rich diet (Groups: CaBP (Carob) and CaBP+PEG (Carob+PEG)) or not (Groups: C (Control), C+PEG (Control+PEG)). Adult worm counts (AWC) shown as arithmetic mean of adult worms (female, male, total) (and SD in brackets) per group.

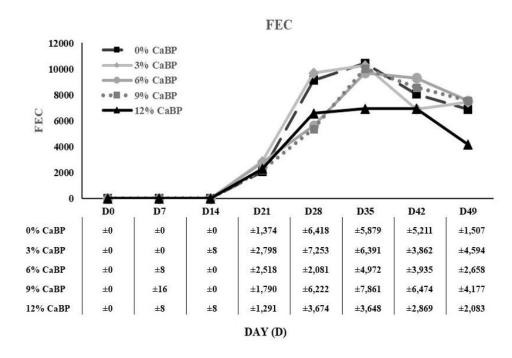
Treatment		H. contortus		T. colubriformis			
Group	Female	Male	Total	Female	Male	Total	
C	897	584	1,480 ^a	5,783	4,382	10,166	
C	(± 736)	(±461)	$(\pm 1,194)$	$(\pm 2,104)$	$(\pm 1,529)$	$(\pm 3,599)$	
C - DEC	1,002	710	1,712 ^a	6,028	4,713	10,742	
C+PEG	(± 323)	(± 279)	(± 593)	$(\pm 2,740)$	$(\pm 2,067)$	$(\pm 4,776)$	
CaBP	288	243	532 ^b	5,397	4,882	10,279	
СавР	(± 220)	(± 199)	(±399)	$(\pm 2,280)$	$(\pm 2,171)$	$(\pm 4,439)$	
CaBP+PEG	701	617	1,318 ^a	5,751	5,303	11,054	
Cadr+PEG	(± 250)	(± 178)	(± 424)	$(\pm 2,387)$	$(\pm 1,743)$	$(\pm 4,091)$	

Table 5. Trial 3: Effect of diet regimes containing different condensed tannin diets on adult worms recovered at necropsy in the different experimental lamb groups C (Control), CaBP (Carob), S (Sainfoin pellets), CaBP+S (Carob+Sainfoin pellets). Adult worm counts (AWC) shown as arithmetic mean of adult worms (female, male, total) (and SD in brackets) per group.

Treatment		H. contortus		T. colubriformis			
Group	Female	Male	Total	Female	Male	Total	
C	3,482	3,302	6,783	1,370	1,082	2,452	
C	$(\pm 3,170)$	$(\pm 2,797)$	$(\pm 5,924)$	(±443)	(± 438)	(± 842)	
CaBP	2,087	2,288	4,375	1,140	1,052	2,192	
Cabr	$(\pm 2,339)$	$(\pm 2,562)$	$(\pm 4,788)$	(± 373)	(± 259)	(±617)	
S	1,315	1,255	2,570	1,735	1,453	3,188	
3	$(\pm 1,594)$	$(\pm 1,707)$	$(\pm 3,286)$	$(\pm 1,414)$	$(\pm 1,029)$	$(\pm 2,406)$	
CaBP+S	1,470	1,685	3,155	1,088	863	1,952	
Cadr+3	$(\pm 1,257)$	$(\pm 1,393)$	$(\pm 2,607)$	$(\pm 1,198)$	(±961)	$(\pm 2,156)$	

Figure 1. Trial 1: Effect of diet regimes containing different amounts of Carob pod meal (CaBP) on A) faecal egg counts (FEC) on Day 0 to 49) (SD in table below) and B) box-plots for female worm fecundity (95% confidence interval) in the different experimental lambs for *Haemonchus contortus* and *Trichostrongylus colubriformis*.

A.



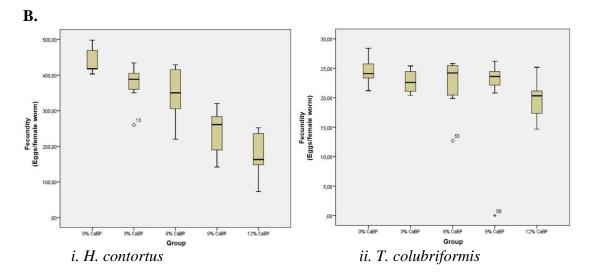
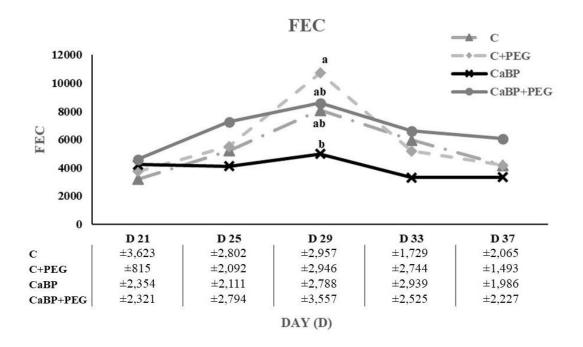


Figure 2. Trial 2: Effect of PEG intake on A) faecal egg counts (FEC) on Day 21 to 37 (SD in table below) and B) box-plots for female worm fecundity (95% confidence interval) for *Haemonchus contortus* and *Trichostrongylus colubriformis* in the different experimental lambs fed with carob pods meal at 12% (CaBP and CaBP+PEG) or served as Controls (C and C+PEG).



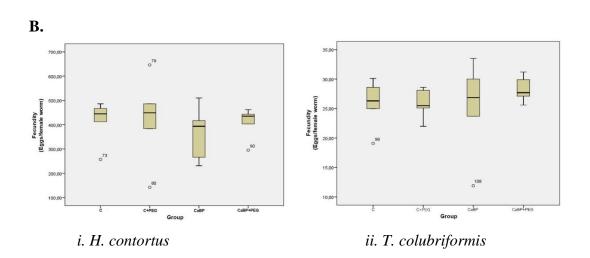
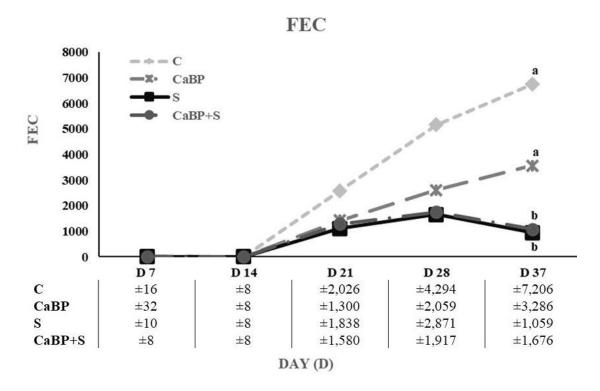
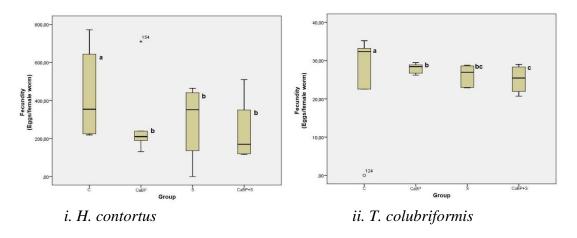


Figure 3. Trial 3: Effect of diet regimes containing different condensed tannin diets on A) faecal egg counts (FEC) on Day 7 to 37 (SD in table below) and B) box-plots for female worm fecundity (95% confidence interval), in the different experimental lambs groups C (Control), CaBP (Carob), S (Sainfoin pellets), CaBP+S (Carob+Sainfoin pellets) for *Haemonchus contortus* and *Trichostrongylus colubriformis*.

A.



B.



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*Conflict of Interest

Declaration of interests
\boxtimes The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.
☐The authors declare the following financial interests/personal relationships which may be considered as potential competing interests:

Author Contribution Statement

SARATSI K was a PhD student and the paper is part of her PhD Thesis, **SOTIRAKI S**, **HADJIGEORGIOU I**, and **HOSTE H** were the Supervisors of her Thesis including the current work. **VOUTZOURAKIS N** and **STEFANAKIS A** are specialist in animal nutrition and supporting the preparation of ration and acquiring the samples, data interpretation and writing the paper. **TZANIDAKIS N**, is a veterinarian supporting with laboratory techniques, **THAMSBORG SM**, is a senior scientists coordinator of CARES project who supervised the trials and supported data analyses, interpretation and writing the paper and **MUELLER-HARVEY I** is a senior researcher expert in tannin analyses who supported in chemical analysis of the feeds and data interpretation and writing the paper.

According to CRediT:

SARATSI K: Conceptualization, Visualization, Investigation, Resources, Formal Analyses, Writing- Original draft preparation, Reviewing and Editing, **SOTIRAKI S, HADJIGEORGIOU I, HOSTE H** and **THAMSBORG SM:** Conceptualization, Supervision, Writing, Reviewing and Editing. **VOUTZOURAKIS N, TZANIDAKIS N and STEFANAKIS A** Investigation, Resources, Reviewing and Editing, **MUELLER-HARVEY I:** Methodology, Investigation, Reviewing and Editing