

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

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| 2  | Feeding of carob (Ceratonia siliqua) to sheep infected with gastrointestinal nematodes                                                 |
| 3  | reduces faecal egg counts and worm fecundity                                                                                           |
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## 26 Abstract

27 The present study explored the anthelmintic effects of condensed tannins (CT) in 28 carob (Ceratonia siliqua) pods fed to sheep against gastrointestinal nematodes. Three 29 independent in vivo trials tested whether i) carob pod (CaBP)-containing feed had an 30 anthelmintic effect and if yes, which was the optimal concentration in the diet; ii) 31 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 32 33 another tannin-containing feed (e.g. sainfoin). In all trials 6-month old nematode-34 naive lambs, experimentally infected with both Haemonchus contortus and 35 Trichostrongylus colubriformis, were used. Faecal egg counts (FEC) were performed 36 regularly and at the end of each trial adult worm counts (AWC) and female worm 37 fecundity were recorded. In trial 1, 35 lambs (five groups of seven lambs) were fed 38 different CaBP concentrations ranging from 0% to 12% w/w. FEC declined up to 39 39.2% only in the group fed with 12%CaBP, while a declining trend (P<0.06) was 40 demonstrated for the AWC of T. colubriformis, which was associated with the 41 increasing concentration of CaBP in feed. Female worm fecundity was reduced in 42 groups fed CaBP for both parasites, however this was only significant for H. 43 contortus (P<0.001), in a dose dependent manner. In trial 2, four groups of six 44 infected lambs each were used, which received the carob diets CaBP or CaBP+PEG, 45 and the tannin-free diets with or without PEG (C or C+PEG). Results showed that 46 FEC of Groups C, C+PEG, and CaBP+PEG were comparable throughout the trial, 47 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 48 49 anthelmintic effect of CaBP after PEG administration suggested that CT contributed to the anthelmintic action. However, no effect of CaBP was observed on T. 50

| 51 | colubriformis AWC and on female worm fecundity for both species. Finally, for trial    |
|----|----------------------------------------------------------------------------------------|
| 52 | 3 four groups of six lambs each received a diet based on CaBP, sainfoin (S) or a       |
| 53 | combination (CaBP+S) and were compared to a control (C) diet of lucerne. On DAY        |
| 54 | 37 FEC values in groups CaBP+S and S tended to be lower compared to the two other      |
| 55 | groups (C, CaBP), while for AWCs no significant differences were observed for both     |
| 56 | parasites. The fecundity of H. contortus and T. colubriformis demonstrated significant |
| 57 | differences between the treated and control groups, with lower values in the animals   |
| 58 | receiving CaBP+S. Overall, the results supported the hypothesis that carob had an      |
| 59 | anthelmintic effect due to its CT, but there was no clear indication of a synergistic  |
| 60 | effect with sainfoin.                                                                  |
| 61 |                                                                                        |
| 62 | Keywords: Carob, Sainfoin, Haemonchus contortus, Trichostrongylus colubriformis,       |
| 63 | gastrointestinal nematodes, sheep, feed additives                                      |
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## 77

## 78 **1. Introduction**

79 Infections by gastrointestinal nematodes (GIN) affect both health and welfare of 80 grazing ruminants, causing anorexia, impaired digestion and nutrient absorption with 81 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 82 83 relies on repeated dosing with commercial anthelmintic drugs. However, the 84 development of anthelmintic resistance in worm populations against one or multiple 85 classes of anthelmintic has become a serious problem in several regions of the world, 86 making it increasingly difficult to control parasitic infections (Kaplan, 2020). At the 87 same time the increasing concerns of consumers about the presence of drug residues 88 in foods and in the environment (McKellar, 1997) have stimulated the search for 89 alternative or complementary solutions (Hoste and Torres-Acosta, 2011) within the 90 context of organic farming and sustainable agriculture (Charlier et al., 2018).

91 Such alternatives include the use of bioactive plants with anthelmintic properties. 92 Many results indicate that such resources, because of the presence of plant secondary 93 metabolites (PSMs), might help to interfere with the biology of key-stages during 94 nematode cycle and to reduce the consequences of GIN infections in grazing 95 ruminants. Particular attention has been given to plants containing condensed tannins 96 (CT) and some related polyphenols (see reviews by Terrill et al., 2012; Hoste et al., 97 2015, 2016). Based on previous studies, the need to explore new plant resources to 98 develop non-drug-based strategies for the integrated control of nematode parasites in grazing ruminants has recently become a research priority in livestock production, 99 100 especially in ruminant breeding as also reviewed by Morgan et al, 2020.

101 Small ruminants (sheep and goats) are a major component of the dairy sector in the Mediterranean basin (Hadijgeorgiou et al., 2005). Sheep and goat production often 102 103 occupy marginal lands that are unsuitable for crop production but are rich in local 104 plants, such as rangeland vegetation, which can be exploited by animals as a feed 105 resource (Frutos et al., 2008; Méndez-Ortiz et al., 2018). Many rangeland plants also 106 contain PSMs, such as tannins (Papachristou et al., 2005) and several in vitro and in 107 vivo studies have evaluated their anthelmintic effects against GINs of small ruminants 108 (Manolaraki et al., 2010; Moreno-Gonzalo et al., 2012, 2013a,b, 2014; Arroyo-Lopez 109 et al., 2014; Silva Soares et al., 2018). Overall, CTs have been shown to directly or 110 indirectly interfere with the life cycle of several GINs and, therefore, CT-containing 111 plants, which also include many legumes, are proving to be beneficial nutritional 112 resources. However, a high degree of variability with respect to their anthelmintic 113 activity has also been recorded. Besides the total tannin concentration in ruminant 114 diets, recent studies have demonstrated that CT molecular composition or structural 115 characteristics can also affect anthelmintic activity (Mueller-Harvey et al., 2019).

116 Carob (Ceratonia siliqua) and sainfoin (Onobrychis viciifolia) are both resources of 117 the Fabaceae family and contain CTs. Carob is a leguminous tree that is widely 118 cultivated in the Mediterranean area. It is an important species both for economic and 119 environmental reasons (Batlle and Tous, 1997). Carob pods (fruits) are mostly used in 120 the food industry; pulp accounts for 90% by pod weight and seeds for 10%. They 121 contain high sugar (48–56%), but low protein (3–4%) and lipid concentrations (0.4– 122 0.8%) (Marakis, 1996; Batlle and Tous, 1997). Moreover, ripe carob pods contain 123 high concentrations of CTs (16-20% w/w DM) (Bravo et al., 1994; Batlle and Tous, 124 1997). This has been debated by Priolo et al. (2000, 2002) who claimed that the pods 125 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.

131 Sainfoin, which can be found especially in southern parts of Europe, has been the 132 subject of renewed interest because of its beneficial effects in the context of 133 agroecology (Hayot Carbonero et al., 2011), its beneficial impact on ruminant 134 production and the environment and its potential antiparasitic effects on small 135 ruminants (Manolaraki et al., 2010; Hoste et al., 2015; Saratsis et al., 2016; Mueller-136 Harvey et. al., 2019). In vitro studies have shown that sainfoin extracts have a dose-137 dependent effect against different GIN species (Brunet et al., 2007; Manolaraki et al., 138 2010; Novobilsky et al., 2013). Moreover, in vivo anthelmintic effects have also been 139 described in sheep and/or goats fed with sainfoin; i.e. 42-68% reduction in parasitic 140 egg excretion, which was associated with a 17.6% decrease in female worm fecundity 141 and a 45% decrease in worm numbers for Haemonchus contortus (Arroyo-Lopez et 142 al., 2014).

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 andsainfoin feeds (Trial 3).

## 152 **2. Materials and Methods**

## 153 **2.1. Stabling and animals**

154 The experiments were carried out at the Asomaton Research Station of HAO Demeter 155 on the island of Crete, Greece. The animal barn was of an open-sided shed type, with 156 straw bedding. During the whole study the animals were kept indoors with each group in a separate pen of approximately  $10 \text{ m}^2$  and  $10 \text{ m}^2$  open yard. The study included 157 158 three trials with lambs belonging to the local "Sfakion" breed. In order to achieve 159 uniformity of the experimental animals, all lambs included were female, 6-month-old 160 with a comparable body weight (BW), which was within the normal BW range of the 161 breed (22-30 kg) at the specific age. The lambs were raised indoors under helminthfree conditions. Fourteen days before the start of each trial, they were drenched with 162 163 albendazole at the higher commercially recommended dose (ALBENDAZOLE 164 Drench, PROVET, 7.5 mg/kg) and they tested negative by faecal egg counts at the 165 start of each trial. No anthelmintic resistance was previously recorded for this specific 166 flock.

## 167 **2.3. Infective larvae**

168 Third-stage infective larvae of *H. contortus* and *T. colubriformis* strains, susceptible 169 to all classes of anthelmintic drugs, had been cultured from faeces of mono-170 specifically infected donor sheep. Larvae were recovered using the Baermann 171 technique and then stored for 1-2 months at 4°C until use.

172

## 173 **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, 3<sup>rd</sup> cut) were provided by Multifolia (Viapres le Petit 10380, France) as part

177 of the Research project CARES.

## 178 **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.

184 The acetone-HCl-butanol assay was carried out as previously described by Grabber et185 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).

## 193 **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.

201 2.6.1. Trial 1

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 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 infectionwith 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 3<sup>rd</sup> stage larvae (L3) of *H. contortus* and 12.000 L3 of *T. colubriformis*. At the end of

the experimental period (D49), all lambs were euthanised by injection of a massivedose of pentobarbital (Dolethal®).

216 **2.6.2.** *Trial 2* 

217 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 242 into heparinized tubes (BD Vacutainer®, UK) to determine the packed cell volume 243 (PCV), as an indicator of anaemia, according to the micro-haematocrit method. In

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

## 246 **2.8. Parasitological parameters**

Individual faecal samples were collected weekly directly from the rectum, during the
1<sup>st</sup> and 3<sup>rd</sup> trial, and twice weekly during the 2<sup>nd</sup> 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).

251 At necropsy, the abomasa and the first 12 meters of small intestine were separated, 252 ligated, rapidly removed and immediately processed to collect the adult worms from 253 the luminal contents. For the intra-mucosal larvae, pepsin digestion was applied both 254 on the abomasum and intestinal mucosa (MAFF, 1986). After 4h incubation at 37°C 255 the larvae were collected. After storage in 10% alcohol, worm counts were performed 256 according to a 10% aliquot technique (MAFF, 1986). Morphological identification of 257 worm stages, sex and species were conducted using standard procedures (MAFF, 258 1986).

259 The fecundity of female worms was measured on 10 worms per lamb. For T. 260 colubriformis, eggs were counted directly in utero after clearing in 85% lactic acid 261 solution. All egg counts were performed under a microscope set at 10 times 262 magnification (total 100  $\times$ ). For *H. contortus*, the fecundity was determined using the 263 method described by Kloostermann et al. (1978). Briefly, the worms were soaked for 264 5 min in a large volume of distilled water, before being placed individually in 265 microtubes with 1000  $\mu$ l of 0.125% hypochlorite concentration solution and kept at 266 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 267 268 (10%) of the total volume.

## 269 **2.9. Statistical analyses**

270 The data of FEC and adult worm counts (AWC) were log10(x+1) transformed prior to 271 analysis. For the FEC values, comparison of all groups was first performed using an 272 analysis of variance (ANOVA) with time as repeated measurement. Then, the 273 comparison of results to the control values were carried out date by date, using one-274 way ANOVA completed by the post-hoc Bonferroni test for pairwise comparisons. 275 Group means of AWC were compared by one-way ANOVA (Trial 1) or two-way 276 ANOVA (Trial 2: CaBP +/- and PEG +/-; Trial 3: CaBP +/- and sainfoin +/-). 277 Regarding the fecundity of female worms, the Shapiro-Wilk Test of normality, which 278 is more appropriate for small sample sizes, was used. In cases where the data deviated 279 significantly (P<0.05) from a normal distribution (Trial 1 and 3 for both parasite 280 species and Trial 2 for *T. colubriformis*) the appropriate test to check the difference of 281 fecundity between the groups, which is the non-parametric test of Kruskal-Wallis, 282 was used. Where the dependent variable was normally distributed (P>0.05) the 283 parametric test of one-way ANOVA (H. contortus of Trial 2) was used. Additionally, 284 for the Trial 1, the model of linear regression was used, in order to be investigated if 285 there was a negative correlation between the variables "percentages of carob" and 286 "fecundity of female worms" for both parasite species (H. contortus and T. 287 colubriformis). Finally, the Tukey HSD test was used for data of trial 3, in order to 288 investigate statistically significant differences between groups.

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

290

## 291 **2.10. Ethical considerations**

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

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.

- 298
- 299 **3. Results**

300 The CT concentrations and compositions are presented in Table 2. The HBA assay 301 yielded similar CT concentrations for both plant materials, whereas the thiolysis assay 302 generated lower CT concentrations for the sainfoin pellets. The thiolysis assay 303 revealed that: i) both carob and sainfoin CTs consisted mainly of prodelphinidins, 304 96.7 and 74.7 mole percentages, respectively; ii) carob CTs were highly galloylated 305 (i.e. 41.1% of flavan-3-ol subunits are galloylated), but sainfoin CTs did not contain 306 any esterified galloyl groups; iii) carob CTs were characterised by a relatively high 307 average molecular weight (mDP = 31.1), whereas sainfoin CTs had an mDP value of 308 11.5.

309

## 310 **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, 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 325 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 330 were statistically significant differences in fecundity between the groups (P<0.001). 331 More specifically, fecundity was statistically significantly greater for 0% CaBP group 332 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). However, the model of linear regression, which was implemented and was 334 335 statistically significant (P<0.05), showed a negative correlation between the variables 336 "group" and "fecundity" for both parasite species.

337 No GIN larvae were recovered after pepsin digestion.

Mean PCV values ( $\pm$ SD) for groups 0%CaBP, 3%CaBP, 6%CaBP, 9%CaBP, and 12%CaBP on the last day of the trial were 25.29 ( $\pm$ 5.96), 23.00 ( $\pm$ 5.72), 21.00 ( $\pm$ 6.32), 23.00 ( $\pm$ 5.89) and 24.00 ( $\pm$ 5.00) respectively. No significant differences were 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 no significant differences between the groups.

346 **3.2.Trial 2** 

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.

Results on AWC, showed reduction only for *H. contortus* (P<0.03) resulting in an overall statistical difference between the 4 groups, since the lowest worm counts were found for the CaBP group. Especially, for *H. contortus*, a reduction of approximately 65% was observed in the carob group compared to the control. The AWC in the CaBP+PEG group were similar to the other 2 control groups showing no reduction in worm population. On the other hand, no effect of carob was observed on *T. colubriformis* worm counts.

No effect of carob on female fecundity was also observed, irrespective of the parasite species. Both control and carob groups showed comparable levels of female fecundity for the two parasite species. The Box plot in Figure 2b showed that the range of 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 on the fecundity scale in the CaBP group than in the other groups. 369 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(±30.1) (±s.d.), 69.8(±19.9),

372  $60.8(\pm 29.3)$  and  $40.5(\pm 25.6)$  g, respectively, which resulted in no significant

differences between the groups.

374 3.3.Trial 3

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 380 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 381 382 values of FEC, the reductions in the 3 treated groups ranged from 44.6% to 383 approximately 86 %. These differences were mainly found for the sainfoin group (S) 384 and carob+sainfoin (CaBP+S) groups. As regards the AWCs, no significant 385 differences were observed neither in the number of *H. contortus* and *T. colubriformis*. 386 No GIN larvae were recovered after pepsin digestion.

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 the C group differed significantly from CaBP, S and CaBP+S, while for *T. colubriformis* fecundity for CaBP group was also statistically different from CaBP+S (Figure 3b). When exploring the pathophysiological parameters (i.e. PCV), the analysis of variance on repeated measures and also the date by date ANOVA did not show significant differences between the groups. Specific values for mean PCV ( $\pm$ SD) on DAY 28 of the respective groups C, CaBP, S and CaBP+S were 31.67 ( $\pm$ 3.39), 33.00 ( $\pm$ 4.86), 31.33 ( $\pm$ 3.61) and 30.50 ( $\pm$ 4.37).

The average daily gain (ADG) as calculated for the whole trial duration for (C), (CaBP), (S) and (CaBP+S) groups was (mean $\pm$ s.d.) 122.5( $\pm$ 38.1), 88.2( $\pm$ 39.2), 104.6( $\pm$ 11.9) and 124.8( $\pm$ 39.7) g, respectively and there were no significant differences between the groups.

## 403 **4. Discussion**

404 The literature contains several in vitro and in vivo studies, conducted on small 405 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 413 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 (Mupeyo et al., 416 2011).

417 In the current study, we further explored the *in vivo* anthelmintic effects of carob pod 418 meal since it represents a common feed resource in the Mediterranean region and 419 there was some previous evidence of its anthelmintic (Arroyo-Lopez et al., 2014) and 420 anticoccidial (Saratsis et al., 2016; Legendre et al., 2018) properties. In order to 421 develop a practical implementation tool for carob as dietary intervention, we wanted 422 to identify a) the optimal carob concentration in the feed for bioactivity, b) whether 423 CTs contributed to such an activity and c) whether there were any synergistic effects 424 with other plant sources with different types of CTs (i.e. sainfoin). For all 3 trials a 425 balanced and palatable ration was specifically designed for all animals. This aimed to 426 achieve similar production indexes in all groups and ensured that any observed 427 differences in the effects of parasitism would not stem from quantitative differences in 428 the dietary composition but rather from differences in the bioactive CTs (Coop and 429 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 441 and this effect is due mainly to the reduction of female worm fecundity 442 (predominantly in H. contortus) and to a lesser extent to the reduction of establishment and development of the worms. Since H. contortus produce a 443 444 remarkably high daily egg output compared to T. colubriformis (Besier et al., 2016), 445 we suggest that the reduction in FEC seen in this trial can be attributed to the effect 446 the carob diet had against H. contortus. Overall, the results of this trial suggest that 447 the higher the concentration of carob in the ration the higher the anthelmintic activity; 448 this effect that was more evident for *H. contortus*. Unfortunately, there are limitations 449 to the quantity of carob pod meal that can be included in a well balanced ration since 450 carob pods contain high sugar but low protein and lipid concentrations (Priolo et al., 451 1998; Karabulut et al., 2006).

452 During Trial 2, the main results i) confirmed that CaBP reduced FEC in lambs, as 453 these reductions compared to control values ranged from 20% to 45%, ii) that these 454 reductions in FEC seemed to be mainly due to the lower numbers from the highly 455 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 456 457 CaBP may be attributed to CTs, because a restoration to control values for FEC and 458 Haemonchus worm numbers was observed in the CaBP + PEG group. PEG is a non-459 nutritive synthetic polymer that is capable of binding and deactivating CTs; it has 460 been used in many animal nutrition studies to increase the intake of CT-containing 461 feeds and to improve protein absorption (Silanikove et al., 1996; Bermingham et al., 462 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 463 464 anthelmintic activity was linked to the presence of CTs.

465 Finally, the aim of Trial 3 was to investigate two hypotheses: firstly, that carob CTs 466 generate a stronger anthelmintic effect than sainfoin CTs and secondly, that 467 synergistic effects could be achieved by combining carob with sainfoin. The rationale 468 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 469 470 CTs are highly galloylated prodelphinidins, whereas sainfoin CTs are non-galloylated 471 prodelphinidins. Previous studies found two structural features in CTs that enhance 472 anthelmintic activity in vitro: i) prodelphinidin CTs are more potent than procyanidin 473 CTs and ii) galloylation increases the anthelmintic effect of CTs (Hoste et al., 2016; 474 Kommuru et al., 2014, 2015). Therefore, carob CTs, which have a high 475 prodelphinidin/procyanidin ratio (96.7% prodelphinidins/3.3% procyanidins) and are 476 also highly galloylated (i.e. 41.1% of the flavan-3-ol subunits are galloylated) should 477 produce a stronger anthelmintic effect than sainfoin, as sainfoin CTs have less 478 prodelphinidins (74.8%) and no galloyl groups (N.B. % stands for mole percent 479 within CT molecules; Table 2).

480 There are several important reasons that could explain why the results from Trial 3 481 did not support either of these hypotheses. Firstly, sainfoin - but not carob - was fed in 482 a pelleted form, while it has been demonstrated previously that the pelleting process 483 has a marked effect on CTs in terms of their analysis (Mueller-Harvey et al., 2019). 484 Table 2 shows that the CT concentrations in sainfoin pellets differed considerably 485 between the two assays (6.5 and 1.7 g CT/100 g DW) in contrast to the carob meal 486 data (5.8 and 7.2 g CT/100g DW). However, we currently do not know whether the 487 pelleting process enhances the anthelmintic activity of CTs or not. Secondly, up to 488 now most attempts to unravel links between CT structural features and anthelmintic 489 effects have employed in vitro assays. Therefore, in vivo feeding trials such as the 490 present ones are vital to test the laboratory data. It may turn out that the esterified 491 galloyl groups are not stable in the digestive tract and that the prodelphinidins in 492 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.

497 The nutritional and/or anthelmintic properties of sainfoin fed as direct grazing, silage, 498 hay or pellets have been evaluated in both sheep and goats, with promising 499 anthelmintic results when used either alone (Paolini et al., 2005; Heckendorn et al., 500 2006; Ríos-de Alvarez et al., 2008; Gaudin et al., 2016) or in combination with other 501 CT sources (Girard et al., 2013). Previous results have demonstrated that sainfoin 502 consumption under different forms of preservation can reduce FEC and also reduce 503 female worm fecundity of H. contortus (Manolaraki et al., 2010; Arroyo-Lopez et al., 504 2014) or T. colubriformis (Manolaraki et al., 2010); however, in other studies a lack 505 of effect has been observed (Heckendorn et al., 2006). The issue of the variable 506 results has also been addressed in several reviews (Hoste et al., 2015; Hoste and 507 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

515 the other hand, although the differences were not significant, the percentage of 516 reduction compared to the controls (Group C) for *H. contortus* worm numbers were 517 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).

523 Our results confirmed that i) the consumption of CT containing resources can 524 modulate the biology of GINs; ii) that CT were involved in the anthelmintic effects of 525 carob and iii) the concentration in the diet influenced the anthelmintic effects as 526 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 527 528 the worm population and could explain the reduction of FECs: either a reduced 529 fecundity of female adult worms (see Trial 1 and 3) and /or a reduction of the number 530 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.

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537 Our results suggest that, when worm populations are exposed to CTs in the 538 gastrointestinal tract, upon their ingestion by the host the most evident effect recorded 539 is the reduction of female fecundity. Particularly for *H. contortus*, it appears that 540 fecundity is only affected when the worms are exposed to CTs during maturation 541 (Trial 1 and 3) and not when they are already mature adults (Trial 2). On the other 542 hand, when CaBP was consumed for two weeks by animals in which adult worm 543 populations were already established and patent, the main finding was a significant 544 decrease in H. contortus worm counts. Moreover, the current study adds further 545 support to the observation that most of the CT effect is related to abomasal parasite -546 and not so much to the small intestinal parasite. This is possibly due to the higher CT 547 concentration in the abomasum compared with the rumen and intestines, along with 548 higher prodelphinidin percentage as already shown in studies on the cattle abomasal 549 parasite Ostertagia ostertagi (Desrues et al., 2017).

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| 2  | Feeding of carob (Ceratonia siliqua) to sheep infected with gastrointestinal nematodes                                                 |
| 3  | reduces faecal egg counts and worm fecundity                                                                                           |
| 4  |                                                                                                                                        |
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#### 26 Abstract

27 The present study explored the anthelmintic effects of condensed tannins (CT) in 28 carob (Ceratonia siliqua) pods fed to sheep against gastrointestinal nematodes. Three 29 independent in vivo trials tested whether i) carob pod (CaBP)-containing feed had an 30 anthelmintic effect and if yes, which was the optimal concentration in the diet; ii) 31 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 32 33 another tannin-containing feed (e.g. sainfoin). In all trials 6-month old nematode-34 naive lambs, experimentally infected with both Haemonchus contortus and 35 Trichostrongylus colubriformis, were used. Faecal egg counts (FEC) were performed 36 regularly and at the end of each trial adult worm counts (AWC) and female worm 37 fecundity were recorded. In trial 1, 35 lambs (five groups of seven lambs) were fed 38 different CaBP concentrations ranging from 0% to 12% w/w. FEC declined up to 39 39.2% only in the group fed with 12%CaBP, while a declining trend (P<0.06) was 40 demonstrated for the AWC of T. colubriformis, which was associated with the 41 increasing concentration of CaBP in feed. Female worm fecundity was reduced in 42 groups fed CaBP for both parasites, however this was only significant for H. 43 contortus (P<0.001), in a dose dependent manner. In trial 2, four groups of six 44 infected lambs each were used, which received the carob diets CaBP or CaBP+PEG, 45 and the tannin-free diets with or without PEG (C or C+PEG). Results showed that 46 FEC of Groups C, C+PEG, and CaBP+PEG were comparable throughout the trial, 47 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 48 49 anthelmintic effect of CaBP after PEG administration suggested that CT contributed to the anthelmintic action. However, no effect of CaBP was observed on T. 50

| 51 | colubriformis AWC and on female worm fecundity for both species. Finally, for trial    |
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| 52 | 3 four groups of six lambs each received a diet based on CaBP, sainfoin (S) or a       |
| 53 | combination (CaBP+S) and were compared to a control (C) diet of lucerne. On DAY        |
| 54 | 37 FEC values in groups CaBP+S and S tended to be lower compared to the two other      |
| 55 | groups (C, CaBP), while for AWCs no significant differences were observed for both     |
| 56 | parasites. The fecundity of H. contortus and T. colubriformis demonstrated significant |
| 57 | differences between the treated and control groups, with lower values in the animals   |
| 58 | receiving CaBP+S. Overall, the results supported the hypothesis that carob had an      |
| 59 | anthelmintic effect due to its CT, but there was no clear indication of a synergistic  |
| 60 | effect with sainfoin.                                                                  |
| 61 |                                                                                        |
| 62 | Keywords: Carob, Sainfoin, Haemonchus contortus, Trichostrongylus colubriformis,       |
| 63 | gastrointestinal nematodes, sheep, feed additives                                      |
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# 78 **1. Introduction**

79 Infections by gastrointestinal nematodes (GIN) affect both health and welfare of 80 grazing ruminants, causing anorexia, impaired digestion and nutrient absorption with 81 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 82 83 relies on repeated dosing with commercial anthelmintic drugs. However, the 84 development of anthelmintic resistance in worm populations against one or multiple 85 classes of anthelmintic has become a serious problem in several regions of the world, 86 making it increasingly difficult to control parasitic infections (Kaplan, 2020). At the 87 same time the increasing concerns of consumers about the presence of drug residues 88 in foods and in the environment (McKellar, 1997) have stimulated the search for 89 alternative or complementary solutions (Hoste and Torres-Acosta, 2011) within the 90 context of organic farming and sustainable agriculture (Charlier et al., 2018).

91 Such alternatives include the use of bioactive plants with anthelmintic properties. 92 Many results indicate that such resources, because of the presence of plant secondary 93 metabolites (PSMs), might help to interfere with the biology of key-stages during 94 nematode cycle and to reduce the consequences of GIN infections in grazing 95 ruminants. Particular attention has been given to plants containing condensed tannins 96 (CT) and some related polyphenols (see reviews by Terrill et al., 2012; Hoste et al., 97 2015, 2016). Based on previous studies, the need to explore new plant resources to 98 develop non-drug-based strategies for the integrated control of nematode parasites in grazing ruminants has recently become a research priority in livestock production, 99 100 especially in ruminant breeding as also reviewed by Morgan et al, 2020.

101 Small ruminants (sheep and goats) are a major component of the dairy sector in the Mediterranean basin (Hadijgeorgiou et al., 2005). Sheep and goat production often 102 103 occupy marginal lands that are unsuitable for crop production but are rich in local 104 plants, such as rangeland vegetation, which can be exploited by animals as a feed 105 resource (Frutos et al., 2008; Méndez-Ortiz et al., 2018). Many rangeland plants also 106 contain PSMs, such as tannins (Papachristou et al., 2005) and several in vitro and in 107 vivo studies have evaluated their anthelmintic effects against GINs of small ruminants 108 (Manolaraki et al., 2010; Moreno-Gonzalo et al., 2012, 2013a,b, 2014; Arroyo-Lopez 109 et al., 2014; Silva Soares et al., 2018). Overall, CTs have been shown to directly or 110 indirectly interfere with the life cycle of several GINs and, therefore, CT-containing 111 plants, which also include many legumes, are proving to be beneficial nutritional 112 resources. However, a high degree of variability with respect to their anthelmintic 113 activity has also been recorded. Besides the total tannin concentration in ruminant 114 diets, recent studies have demonstrated that CT molecular composition or structural 115 characteristics can also affect anthelmintic activity (Mueller-Harvey et al., 2019).

116 Carob (Ceratonia siliqua) and sainfoin (Onobrychis viciifolia) are both resources of 117 the Fabaceae family and contain CTs. Carob is a leguminous tree that is widely 118 cultivated in the Mediterranean area. It is an important species both for economic and 119 environmental reasons (Batlle and Tous, 1997). Carob pods (fruits) are mostly used in 120 the food industry; pulp accounts for 90% by pod weight and seeds for 10%. They 121 contain high sugar (48–56%), but low protein (3–4%) and lipid concentrations (0.4– 122 0.8%) (Marakis, 1996; Batlle and Tous, 1997). Moreover, ripe carob pods contain 123 high concentrations of CTs (16-20% w/w DM) (Bravo et al., 1994; Batlle and Tous, 124 1997). This has been debated by Priolo et al. (2000, 2002) who claimed that the pods 125 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.

131 Sainfoin, which can be found especially in southern parts of Europe, has been the 132 subject of renewed interest because of its beneficial effects in the context of 133 agroecology (Hayot Carbonero et al., 2011), its beneficial impact on ruminant 134 production and the environment and its potential antiparasitic effects on small 135 ruminants (Manolaraki et al., 2010; Hoste et al., 2015; Saratsis et al., 2016; Mueller-136 Harvey et. al., 2019). In vitro studies have shown that sainfoin extracts have a dose-137 dependent effect against different GIN species (Brunet et al., 2007; Manolaraki et al., 138 2010; Novobilsky et al., 2013). Moreover, in vivo anthelmintic effects have also been 139 described in sheep and/or goats fed with sainfoin; i.e. 42-68% reduction in parasitic 140 egg excretion, which was associated with a 17.6% decrease in female worm fecundity 141 and a 45% decrease in worm numbers for Haemonchus contortus (Arroyo-Lopez et 142 al., 2014).

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 andsainfoin feeds (Trial 3).

### 152 **2. Materials and Methods**

## 153 **2.1. Stabling and animals**

154 The experiments were carried out at the Asomaton Research Station of HAO Demeter 155 on the island of Crete, Greece. The animal barn was of an open-sided shed type, with 156 straw bedding. During the whole study the animals were kept indoors with each group in a separate pen of approximately  $10 \text{ m}^2$  and  $10 \text{ m}^2$  open yard. The study included 157 158 three trials with lambs belonging to the local "Sfakion" breed. In order to achieve 159 uniformity of the experimental animals, all lambs included were female, 6-month-old 160 with a comparable body weight (BW), which was within the normal BW range of the 161 breed (22-30 kg) at the specific age. The lambs were raised indoors under helminthfree conditions. Fourteen days before the start of each trial, they were drenched with 162 163 albendazole at the higher commercially recommended dose (ALBENDAZOLE 164 Drench, PROVET, 7.5 mg/kg) and they tested negative by faecal egg counts at the 165 start of each trial. No anthelmintic resistance was previously recorded for this specific 166 flock.

#### 167 **2.3. Infective larvae**

168 Third-stage infective larvae of *H. contortus* and *T. colubriformis* strains, susceptible 169 to all classes of anthelmintic drugs, had been cultured from faeces of mono-170 specifically infected donor sheep. Larvae were recovered using the Baermann 171 technique and then stored for 1-2 months at 4°C until use.

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#### 173 **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, 3<sup>rd</sup> cut) were provided by Multifolia (Viapres le Petit 10380, France) as part

177 of the Research project CARES.

### 178 **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.

184 The acetone-HCl-butanol assay was carried out as previously described by Grabber et185 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).

# 193 **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.

201 2.6.1. Trial 1

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 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 infectionwith 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 3<sup>rd</sup> stage larvae (L3) of *H. contortus* and 12.000 L3 of *T. colubriformis*. At the end of

the experimental period (D49), all lambs were euthanised by injection of a massivedose of pentobarbital (Dolethal®).

216 **2.6.2.** *Trial 2* 

217 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 242 into heparinized tubes (BD Vacutainer®, UK) to determine the packed cell volume 243 (PCV), as an indicator of anaemia, according to the micro-haematocrit method. In

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

#### 246 **2.8. Parasitological parameters**

Individual faecal samples were collected weekly directly from the rectum, during the
1<sup>st</sup> and 3<sup>rd</sup> trial, and twice weekly during the 2<sup>nd</sup> 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).

251 At necropsy, the abomasa and the first 12 meters of small intestine were separated, 252 ligated, rapidly removed and immediately processed to collect the adult worms from 253 the luminal contents. For the intra-mucosal larvae, pepsin digestion was applied both 254 on the abomasum and intestinal mucosa (MAFF, 1986). After 4h incubation at 37°C 255 the larvae were collected. After storage in 10% alcohol, worm counts were performed 256 according to a 10% aliquot technique (MAFF, 1986). Morphological identification of 257 worm stages, sex and species were conducted using standard procedures (MAFF, 258 1986).

259 The fecundity of female worms was measured on 10 worms per lamb. For T. 260 colubriformis, eggs were counted directly in utero after clearing in 85% lactic acid 261 solution. All egg counts were performed under a microscope set at 10 times 262 magnification (total 100  $\times$ ). For *H. contortus*, the fecundity was determined using the 263 method described by Kloostermann et al. (1978). Briefly, the worms were soaked for 264 5 min in a large volume of distilled water, before being placed individually in 265 microtubes with 1000  $\mu$ l of 0.125% hypochlorite concentration solution and kept at 266 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 267 268 (10%) of the total volume.

#### 269 **2.9. Statistical analyses**

270 The data of FEC and adult worm counts (AWC) were log10(x+1) transformed prior to 271 analysis. For the FEC values, comparison of all groups was first performed using an 272 analysis of variance (ANOVA) with time as repeated measurement. Then, the 273 comparison of results to the control values were carried out date by date, using one-274 way ANOVA completed by the post-hoc Bonferroni test for pairwise comparisons. 275 Group means of AWC were compared by one-way ANOVA (Trial 1) or two-way 276 ANOVA (Trial 2: CaBP +/- and PEG +/-; Trial 3: CaBP +/- and sainfoin +/-). 277 Regarding the fecundity of female worms, the Shapiro-Wilk Test of normality, which 278 is more appropriate for small sample sizes, was used. In cases where the data deviated 279 significantly (P<0.05) from a normal distribution (Trial 1 and 3 for both parasite 280 species and Trial 2 for *T. colubriformis*) the appropriate test to check the difference of 281 fecundity between the groups, which is the non-parametric test of Kruskal-Wallis, 282 was used. Where the dependent variable was normally distributed (P>0.05) the 283 parametric test of one-way ANOVA (H. contortus of Trial 2) was used. Additionally, 284 for the Trial 1, the model of linear regression was used, in order to be investigated if 285 there was a negative correlation between the variables "percentages of carob" and 286 "fecundity of female worms" for both parasite species (H. contortus and T. 287 colubriformis). Finally, the Tukey HSD test was used for data of trial 3, in order to 288 investigate statistically significant differences between groups.

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

290

# 291 **2.10. Ethical considerations**

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

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.

- 298
- 299 **3. Results**

300 The CT concentrations and compositions are presented in Table 2. The HBA assay 301 yielded similar CT concentrations for both plant materials, whereas the thiolysis assay 302 generated lower CT concentrations for the sainfoin pellets. The thiolysis assay 303 revealed that: i) both carob and sainfoin CTs consisted mainly of prodelphinidins, 304 96.7 and 74.7 mole percentages, respectively; ii) carob CTs were highly galloylated 305 (i.e. 41.1% of flavan-3-ol subunits are galloylated), but sainfoin CTs did not contain 306 any esterified galloyl groups; iii) carob CTs were characterised by a relatively high 307 average molecular weight (mDP = 31.1), whereas sainfoin CTs had an mDP value of 308 11.5.

309

# 310 **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, 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 325 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 330 were statistically significant differences in fecundity between the groups (P<0.001). 331 More specifically, fecundity was statistically significantly greater for 0% CaBP group 332 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). However, the model of linear regression, which was implemented and was 334 335 statistically significant (P<0.05), showed a negative correlation between the variables 336 "group" and "fecundity" for both parasite species.

337 No GIN larvae were recovered after pepsin digestion.

Mean PCV values ( $\pm$ SD) for groups 0%CaBP, 3%CaBP, 6%CaBP, 9%CaBP, and 12%CaBP on the last day of the trial were 25.29 ( $\pm$ 5.96), 23.00 ( $\pm$ 5.72), 21.00 ( $\pm$ 6.32), 23.00 ( $\pm$ 5.89) and 24.00 ( $\pm$ 5.00) respectively. No significant differences were 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 no significant differences between the groups.

346 **3.2.Trial 2** 

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.

Results on AWC, showed reduction only for *H. contortus* (P<0.03) resulting in an overall statistical difference between the 4 groups, since the lowest worm counts were found for the CaBP group. Especially, for *H. contortus*, a reduction of approximately 65% was observed in the carob group compared to the control. The AWC in the CaBP+PEG group were similar to the other 2 control groups showing no reduction in worm population. On the other hand, no effect of carob was observed on *T. colubriformis* worm counts.

No effect of carob on female fecundity was also observed, irrespective of the parasite species. Both control and carob groups showed comparable levels of female fecundity for the two parasite species. The Box plot in Figure 2b showed that the range of 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 on the fecundity scale in the CaBP group than in the other groups. 369 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(±30.1) (±s.d.), 69.8(±19.9),

372  $60.8(\pm 29.3)$  and  $40.5(\pm 25.6)$  g, respectively, which resulted in no significant

differences between the groups.

374 3.3.Trial 3

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 380 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 381 382 values of FEC, the reductions in the 3 treated groups ranged from 44.6% to 383 approximately 86 %. These differences were mainly found for the sainfoin group (S) 384 and carob+sainfoin (CaBP+S) groups. As regards the AWCs, no significant 385 differences were observed neither in the number of *H. contortus* and *T. colubriformis*. 386 No GIN larvae were recovered after pepsin digestion.

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 the C group differed significantly from CaBP, S and CaBP+S, while for *T. colubriformis* fecundity for CaBP group was also statistically different from CaBP+S (Figure 3b). When exploring the pathophysiological parameters (i.e. PCV), the analysis of variance on repeated measures and also the date by date ANOVA did not show significant differences between the groups. Specific values for mean PCV ( $\pm$ SD) on DAY 28 of the respective groups C, CaBP, S and CaBP+S were 31.67 ( $\pm$ 3.39), 33.00 ( $\pm$ 4.86), 31.33 ( $\pm$ 3.61) and 30.50 ( $\pm$ 4.37).

The average daily gain (ADG) as calculated for the whole trial duration for (C), (CaBP), (S) and (CaBP+S) groups was (mean $\pm$ s.d.) 122.5( $\pm$ 38.1), 88.2( $\pm$ 39.2), 104.6( $\pm$ 11.9) and 124.8( $\pm$ 39.7) g, respectively and there were no significant differences between the groups.

# 403 **4. Discussion**

404 The literature contains several in vitro and in vivo studies, conducted on small 405 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 413 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 (Mupeyo et al., 416 2011).

417 In the current study, we further explored the *in vivo* anthelmintic effects of carob pod 418 meal since it represents a common feed resource in the Mediterranean region and 419 there was some previous evidence of its anthelmintic (Arroyo-Lopez et al., 2014) and 420 anticoccidial (Saratsis et al., 2016; Legendre et al., 2018) properties. In order to 421 develop a practical implementation tool for carob as dietary intervention, we wanted 422 to identify a) the optimal carob concentration in the feed for bioactivity, b) whether 423 CTs contributed to such an activity and c) whether there were any synergistic effects 424 with other plant sources with different types of CTs (i.e. sainfoin). For all 3 trials a 425 balanced and palatable ration was specifically designed for all animals. This aimed to 426 achieve similar production indexes in all groups and ensured that any observed 427 differences in the effects of parasitism would not stem from quantitative differences in 428 the dietary composition but rather from differences in the bioactive CTs (Coop and 429 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 441 and this effect is due mainly to the reduction of female worm fecundity 442 (predominantly in H. contortus) and to a lesser extent to the reduction of establishment and development of the worms. Since H. contortus produce a 443 444 remarkably high daily egg output compared to T. colubriformis (Besier et al., 2016), 445 we suggest that the reduction in FEC seen in this trial can be attributed to the effect 446 the carob diet had against H. contortus. Overall, the results of this trial suggest that 447 the higher the concentration of carob in the ration the higher the anthelmintic activity; 448 this effect that was more evident for *H. contortus*. Unfortunately, there are limitations 449 to the quantity of carob pod meal that can be included in a well balanced ration since 450 carob pods contain high sugar but low protein and lipid concentrations (Priolo et al., 451 1998; Karabulut et al., 2006).

452 During Trial 2, the main results i) confirmed that CaBP reduced FEC in lambs, as 453 these reductions compared to control values ranged from 20% to 45%, ii) that these 454 reductions in FEC seemed to be mainly due to the lower numbers from the highly 455 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 456 457 CaBP may be attributed to CTs, because a restoration to control values for FEC and 458 Haemonchus worm numbers was observed in the CaBP + PEG group. PEG is a non-459 nutritive synthetic polymer that is capable of binding and deactivating CTs; it has 460 been used in many animal nutrition studies to increase the intake of CT-containing 461 feeds and to improve protein absorption (Silanikove et al., 1996; Bermingham et al., 462 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 463 464 anthelmintic activity was linked to the presence of CTs.

465 Finally, the aim of Trial 3 was to investigate two hypotheses: firstly, that carob CTs 466 generate a stronger anthelmintic effect than sainfoin CTs and secondly, that 467 synergistic effects could be achieved by combining carob with sainfoin. The rationale 468 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 469 470 CTs are highly galloylated prodelphinidins, whereas sainfoin CTs are non-galloylated 471 prodelphinidins. Previous studies found two structural features in CTs that enhance 472 anthelmintic activity in vitro: i) prodelphinidin CTs are more potent than procyanidin 473 CTs and ii) galloylation increases the anthelmintic effect of CTs (Hoste et al., 2016; 474 Kommuru et al., 2014, 2015). Therefore, carob CTs, which have a high 475 prodelphinidin/procyanidin ratio (96.7% prodelphinidins/3.3% procyanidins) and are 476 also highly galloylated (i.e. 41.1% of the flavan-3-ol subunits are galloylated) should 477 produce a stronger anthelmintic effect than sainfoin, as sainfoin CTs have less 478 prodelphinidins (74.8%) and no galloyl groups (N.B. % stands for mole percent 479 within CT molecules; Table 2).

480 There are several important reasons that could explain why the results from Trial 3 481 did not support either of these hypotheses. Firstly, sainfoin - but not carob - was fed in 482 a pelleted form, while it has been demonstrated previously that the pelleting process 483 has a marked effect on CTs in terms of their analysis (Mueller-Harvey et al., 2019). 484 Table 2 shows that the CT concentrations in sainfoin pellets differed considerably 485 between the two assays (6.5 and 1.7 g CT/100 g DW) in contrast to the carob meal 486 data (5.8 and 7.2 g CT/100g DW). However, we currently do not know whether the 487 pelleting process enhances the anthelmintic activity of CTs or not. Secondly, up to 488 now most attempts to unravel links between CT structural features and anthelmintic 489 effects have employed in vitro assays. Therefore, in vivo feeding trials such as the 490 present ones are vital to test the laboratory data. It may turn out that the esterified 491 galloyl groups are not stable in the digestive tract and that the prodelphinidins in 492 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.

497 The nutritional and/or anthelmintic properties of sainfoin fed as direct grazing, silage, 498 hay or pellets have been evaluated in both sheep and goats, with promising 499 anthelmintic results when used either alone (Paolini et al., 2005; Heckendorn et al., 500 2006; Ríos-de Alvarez et al., 2008; Gaudin et al., 2016) or in combination with other 501 CT sources (Girard et al., 2013). Previous results have demonstrated that sainfoin 502 consumption under different forms of preservation can reduce FEC and also reduce 503 female worm fecundity of H. contortus (Manolaraki et al., 2010; Arroyo-Lopez et al., 504 2014) or T. colubriformis (Manolaraki et al., 2010); however, in other studies a lack 505 of effect has been observed (Heckendorn et al., 2006). The issue of the variable 506 results has also been addressed in several reviews (Hoste et al., 2015; Hoste and 507 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

515 the other hand, although the differences were not significant, the percentage of 516 reduction compared to the controls (Group C) for *H. contortus* worm numbers were 517 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).

523 Our results confirmed that i) the consumption of CT containing resources can 524 modulate the biology of GINs; ii) that CT were involved in the anthelmintic effects of 525 carob and iii) the concentration in the diet influenced the anthelmintic effects as 526 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 527 528 the worm population and could explain the reduction of FECs: either a reduced 529 fecundity of female adult worms (see Trial 1 and 3) and /or a reduction of the number 530 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.

536

537 Our results suggest that, when worm populations are exposed to CTs in the 538 gastrointestinal tract, upon their ingestion by the host the most evident effect recorded 539 is the reduction of female fecundity. Particularly for *H. contortus*, it appears that 540 fecundity is only affected when the worms are exposed to CTs during maturation 541 (Trial 1 and 3) and not when they are already mature adults (Trial 2). On the other 542 hand, when CaBP was consumed for two weeks by animals in which adult worm 543 populations were already established and patent, the main finding was a significant 544 decrease in H. contortus worm counts. Moreover, the current study adds further 545 support to the observation that most of the CT effect is related to abomasal parasite -546 and not so much to the small intestinal parasite. This is possibly due to the higher CT 547 concentration in the abomasum compared with the rumen and intestines, along with 548 higher prodelphinidin percentage as already shown in studies on the cattle abomasal 549 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/<br>group | Mean BW<br>at start<br>±s.d. (kg) | Day<br>feeding<br>started | Infection<br>Day | Inoculation<br>dose (L3)      | Day<br>trial<br>ended |
|-------|------------|-----------------|-----------------------------------|---------------------------|------------------|-------------------------------|-----------------------|
|       | CaBP 0%    |                 | 25.8±1.1                          |                           | 0                | 12,000 <i>H</i> .             |                       |
|       | CaBP 3%    |                 | 26.2±3.7                          |                           |                  | contortus                     |                       |
| 1     | CaBP 6%    | 7               | 25.2±2.6                          | -14                       |                  | &                             | 49                    |
|       | CaBP 9%    |                 | 27.1±3.5                          |                           |                  | 12,000 T.<br>colubriformis    |                       |
|       | CaBP 12%   |                 | 26.4±2.8                          |                           |                  |                               |                       |
|       | С          |                 | 26.4±2.7                          | 21                        | 0                | 8,000 L3 <i>H</i> .           |                       |
| •     | C+PEG      | -               | 26.5±2.6                          |                           |                  | contortus                     | 27                    |
| 2     | CaBP (12%) | 6               | 26.3±2.4                          |                           |                  | &                             | 37                    |
|       | CaBP+PEG   |                 | 27.0±1.7                          |                           |                  | 16,000 L3 T.<br>colubriformis |                       |
|       | С          |                 | 27.0±3.1                          |                           |                  | 12,000 L3 <i>H</i> .          |                       |
| •     | CaBP (12%) | -               | 27.1±2.5                          |                           | 0                | contortus                     | ~-                    |
| 3     | CaBP+S     | 6               | 27.1±2.9                          | -14                       |                  | &                             | 37                    |
|       | S          |                 | 26.8±3.2                          |                           |                  | 12,000 L3 T.<br>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-<br>HCl/butanol) | Tannins<br>(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 <sup>a</sup> | 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)       |  |

<sup>a</sup>The 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         |  |
|           | 2,777    | 2,331          | 5,109    | 2,063            | 1,097  | 3,160         |  |
| 0% CaBP   | (±1,579) | (±1,382)       | (±2,802) | (±534)           | (±471) | (±944)        |  |
| 20/ C-DD  | 2,789    | 2,006          | 4,794    | 2,493            | 1,163  | 3,656         |  |
| 3% CaBP   | (±1,606) | $(\pm 1, 117)$ | (±2,709) | (±986)           | (±568) | (±1,461)      |  |
|           | 3,584    | 2,570          | 6,154    | 2,514            | 903    | 3,417         |  |
| 6% CaBP   | (±1,570) | $(\pm 1, 178)$ | (±2,595) | (±497)           | (±676) | (±1,073)      |  |
| 00/ C-DD  | 3,029    | 2,799          | 5,827    | 1,910            | 633    | 2,543         |  |
| 9% CaBP   | (±1,385) | (±1,415)       | (±2,692) | (±1,043)         | (±427) | $(\pm 1,418)$ |  |
| 120/ C-DD | 2,160    | 2,039          | 4,199    | 1,550            | 944    | 2,494         |  |
| 12% CaBP  | (±1,362) | (±1,212)       | (±2,422) | (±801)           | (±630) | (±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    |  |
| 0         | 897    | 584          | 1,480 <sup>a</sup> | 5,783            | 4,382    | 10,166   |  |
| C         | (±736) | (±461)       | (±1,194)           | $(\pm 2, 104)$   | (±1,529) | (±3,599) |  |
|           | 1,002  | 710          | 1,712 <sup>a</sup> | 6,028            | 4,713    | 10,742   |  |
| C+PEG     | (±323) | (±279)       | (±593)             | $(\pm 2,740)$    | (±2,067) | (±4,776) |  |
| CaDD      | 288    | 243          | 532 <sup>b</sup>   | 5,397            | 4,882    | 10,279   |  |
| CaBP      | (±220) | (±199)       | (±399)             | $(\pm 2,280)$    | (±2,171) | (±4,439) |  |
|           | 701    | 617          | 1,318 <sup>a</sup> | 5,751            | 5,303    | 11,054   |  |
| CaBP+PEG  | (±250) | (±178)       | (±424)             | (±2,387)         | (±1,743) | (±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    |  |
| <u> </u>  | 3,482    | 3,302        | 6,783    | 1,370            | 1,082    | 2,452    |  |
| C         | (±3,170) | (±2,797)     | (±5,924) | (±443)           | (±438)   | (±842)   |  |
| C-DD      | 2,087    | 2,288        | 4,375    | 1,140            | 1,052    | 2,192    |  |
| CaBP      | (±2,339) | (±2,562)     | (±4,788) | (±373)           | (±259)   | (±617)   |  |
| C         | 1,315    | 1,255        | 2,570    | 1,735            | 1,453    | 3,188    |  |
| 5         | (±1,594) | (±1,707)     | (±3,286) | (±1,414)         | (±1,029) | (±2,406) |  |
| CaDD      | 1,470    | 1,685        | 3,155    | 1,088            | 863      | 1,952    |  |
| CaBP+S    | (±1,257) | (±1,393)     | (±2,607) | (±1,198)         | (±961)   | (±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*.





DAY (D)



**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.





i. H. contortus

A.

B.

ii. T. colubriformis

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### **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