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Rabbit dietary supplementation with pale purple coneflower (Echinacea pallida). 1. Effects on the reproductive performance and immune parameters of does

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(Article begins on next page)

| 1 | Effects of dietary supplementation with pale purple coneflower (Echinacea |
|---|---|
| 2 | pallida) on reproductive performance and immunity of rabbit does and on |
| 3 | productive results of their kits |

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- 21 Running head: Pale purple coneflower in rabbit nutrition
- 22

23 Abstract

Echinacea pallida (EPAL), also known as pale purple coneflower, is an herbaceous flowering plant with immune-enhancement and antioxidative properties. EPAL effect was studied on rabbit does' reproductive performance, serum biochemistry and

haematological parameters as well as on their kits growth performance. One hundred 27 21-weeks-old Grimaud rabbit does were randomly assigned to two groups. One 28 group was fed a basal diet supplemented with 3 g EPAL /kg diet (Echinacea group, 29 E) while the other was fed the basal diet without the supplementation (Control group, 30 C). Reproductive performance of does was not affected by the treatment (P>0.05). 31 Haematological parameters of pregnant rabbits showed that any interaction between 32 gestational day and treatment was observed except for neutrophils cells (P=0.033). 33 The control group was significant higher than the treatment group for basophils cells 34 (0.55 and 0.29 %, respectively; P=0.049). Gestational day significantly affected most 35 haematological parameters (P<0.05). No significant effect of gestational day or 36 treatment was observed on blood serum chemistry. Regarding the immune 37 parameters, no significant differences were observed between groups; while a 38 significant effect of gestational day was observed for lysozymes (6.02 vs 7.99 vs 39 1.91; for 0, 14 and 28 days respectively; *P*=0.014). Eighty weaned kits (40 born from 40 C does and 40 born from E does) were randomly assigned to four groups of 20 41 animals each fed a growing commercial diet supplemented with or without 3 g EPAL 42 /kg diet. The following experimental groups were formed: CC (rabbits fed the C diet 43 and born from the C does), CE (rabbits fed the E diet and born from the C does), EC 44 (rabbits fed the C diet and born from the E does) and EE (rabbits fed the E diet and 45 born from the E does). Dietary EPAL treatment did not significantly (P>0.05) affect 46 the growth performance of weaned rabbits. In conclusion, a lack of effect of EPAL 47 was reported. Indeed, its dietary supplementation did negatively influence the 48 reproductive and haematological parameters of does nor the growing performance of 49 fattening rabbits. 50

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Keywords: pale purple coneflower, Echinacea pallida, rabbit does, haematology,
fattening rabbits.

54

55 Implications

In recent years, after the ban on the use of antibiotics as growth promoters, phyto-56 additives have been proposed to improve rabbit health and reduce post-weaning 57 mortality. The present study describes the effects of dietary supplementation with 58 Echinacea pallida (known to possess immune-enhancement and antioxidative 59 properties) on rabbit does reproductive performance and immunity and on their kits 60 productive results. The EPAL dietary supplementation did not influence the 61 reproductive and haematological parameters of rabbit does nor did promote the 62 growth performance of their kits. 63

64

65 Introduction

Animal health is a critical issue in animal production strongly affecting the income 66 generated from husbandry activity. Moreover, since the European Union has banned 67 the use of antibiotics as feed additives, many researches in the animal nutrition area 68 have been focused on gauging alternative feeding strategies preventing digestive 69 diseases while enabling the achievement of a satisfactory growth performance. Given 70 the advance in modern biotechnology, the application of naturally-occurring 71 antimicrobial and antioxidant compounds has been preferably employed in animal 72 nutrition due to its potential health benefits on the host physiology (Chrastinová et al., 73 2010). The immunomodulatory and anti-oxidative properties of officinal plants are 74 well known, as well as their ability to promote positive outcomes on animal health and 75 performance (Böhmer et al., 2009; Arafa et al., 2010). 76

Echinacea is a genus of herbaceous flowering plants belonging to the Asteraceae 77 78 botanical family. It presents high levels of production and economic importance in the United States of America, Canada and European countries. The use of a mixture of 79 Echinacea purpurea, Echinacea angustifolia and Echinacea pallida (EPAL) has been 80 reported to have immune-enhancement properties and benefits, such as the 81 prevention and treatment of upper respiratory tract infections (Barnes et al., 2005). 82 Active components from *Echinacea* extracts (mainly alkylamides, polysaccharides 83 and proteoglycans) have been shown to exert immunomodulatory, anti-inflammatory 84 and anti-viral activities (Barnes et al., 2005). Extracts of EPAL have been proposed 85 86 as phyto-immunostimulating agents and their activities are mainly directed towards the innate immune system. Most studies performed on the immunotropic properties 87 of EPAL were related to its effect on nonspecific immunity (activation of macrophage 88 89 functions, phagocytosis of granulocytes, NK cells cytotoxicity), while other studies have investigated the adaptive immune modulation of EPAL (Egger et al., 2008). 90

Improvement of immunity parameters and productive performance has been reported 91 in various livestock species (poultry, quails and rabbits) fed diets supplemented with 92 Echinacea spp. (Maass et al., 2005; Ahmed et al., 2008; Böhmer et al., 2009; Arafa 93 et al., 2010; Nasir and Grashorn, 2010; Sahin et al., 2012). Nevertheless, scarce and 94 conflicting evidence is available concerning the use of Echinacea spp. products in 95 rabbit does during pregnancy. Based on this evidence, the aim of this study was to 96 evaluate the effects of EPAL dietary supplementation on reproductive performance, 97 blood parameters and immune indices in rabbit does as well as on the productive 98 performance of their kits. 99

100

101 Material and methods

103 Animals, housing, diets and management of rabbit does

One hundred nulliparous does (14 week old) of a strain of Grimaud rabbits, obtained from Grimaud Italy, were housed individually in a closed rabbitry, with flat-deck wire net cages (40×50 cm², including nest boxes: 41×26 cm²), and under a constant photo-period of 16 h of light per day. The rabbitry temperature was kept within 18°-22°C. A relative humidity of 60-75% was maintained by a forced ventilation system.

The does were randomly assigned to two groups (50 does per group). The first group was fed *ad libitum* a commercial pelleted diet (control diet, C) while the second one was fed the same diet supplemented with 3 g of EPAL powder /kg diet (*Echinacea* diet, E).

The doe rabbit diets were provided by the Ferrero S.p.A. feed manufacturer 113 (Farigliano CN, Italy). Dry ground EPAL roots, obtained from Biotrade Snc[®] (Via 114 Pacinotti, 21, Mirandola, Italy), was included in the treated diets during the raw 115 material mixing process. The feeding program consisted of a diet provided from 116 insemination to 21 days after parturition and another diet provided from 21 days after 117 parturition to kits weaning. The diets contained the following ingredients in 118 decreasing order: alfalfa meal, sunflower meal, barley, wheat bran, dried beet pulp, 119 maize germ, roasted soybean meal, cane molasses, soybean oil, calcium carbonate, 120 sodium chloride. The diets were analyzed for dry matter (DM, AOAC 925.40), crude 121 protein by total nitrogen contents (AOAC 984.13), ether extract (AOAC 945.16), 122 crude fiber (AOAC 962.09) and ash by ignition to 550°C (AOAC 923.03) according to 123 the Association of Official Analytical Chemists (AOAC, 2000). NDF, ADF and ADL 124 were determined according to Van Soest et al. (1991). Starch was determined by 125 means of Ewer's polarimetric method (European Economic Community, 1972). The 126

chemical composition of the different diets was reported in Table 1. Water was 127 available ad libitum from nipple drinkers. The diets were completely exempt from 128 medication (antibiotics or coccidiostat). All animals were reared under the same 129 environmental and management conditions during the whole experimental period. 130 Rabbit does were first artificially inseminated at 21 weeks of age (mean body weight: 131 3712 ± 176g). Then, artificial insemination was applied at 18 days post-partum (49 132 day reproductive rhythm and single batch system). Cross-fostering was applied within 133 the experimental groups with a maximum of 8, 9 and 10 kits per litter at first, second 134 and following kindling, respectively. The kits were freely nursed by their doe and 135 136 weaned at 35 days of age.

137

138 Does performance

Data of the first five consecutive reproductive cycles were evaluated. Body weight of 139 does at first and final kindling, does mortality and reproductive performance variables 140 were studied. The following variables were calculated on the basis of IRRG's 141 recommendations (International Rabbit Reproduction Group, 2005): total born; born 142 alive; stillborn; litter size at 21 and 35 days of age; litter weight at 21 and 35 days of 143 age; individual body weight of kits at 21 and 35 days of age; Kindling rate (%) = 144 number of kindled does per number of inseminated does x100; Prolificacy = number 145 of born kits per number of does kindled; Numerical productivity at birth = number of 146 born alive per inseminated doe; Overall productivity at birth = weight of born alive per 147 inseminated doe; Perinatal mortality (%) = number of stillborn kits per number of total 148 born x 100; mortality between 0-21 and 0-35 days of age. 149

150

151 Haematological, serum biochemistry and serum electrophoresis of rabbit does

Blood samples were collected from 8 rabbits per group at different time points during 152 the second gestation. Considering the day of artificial insemination as starting day 153 (T0), blood samples were collected at: day 0, day 14 and day 28, respectively. The 154 samples were collected from the lateral saphenous vein with a heparinized syringe to 155 prevent the blood clot. At each sampling time point, one ml of blood was collected 156 into sterile tubes containing ethylenediaminetetraacetic acid -2K (SB-41: Sysmex 157 Corporation) for the evaluation of haematological parameters. Meanwhile, serum 158 obtained by collecting four ml blood samples in a sterile serum plain tube, after 159 incubation at room temperature (22°C) for two hours and centrifugation at 2500 g for 160 10 minutes, was used for serum biochemistry and serum electrophoresis. Serum was 161 stored at -80° C until analysis. Full blood count was performed using an automated 162 laser cell counter calibrated for rabbits (MS4-S Hematology Analyzer, Melet 163 164 Schloesing, Osny - France) to assess the following parameters: red blood cells (RBC, M/mm³), haemoglobin (Hb, g/dl), haematocrit (HCT, %), mean corpuscular volume 165 (MCV, fl), mean corpuscular haemoglobin (MCH, pg), mean corpuscular 166 haemoglobin concentration (MCHC, g/dl), red cell distribution width (RDW, %), 167 platelets (PLT, m/mm³), relative volume of thrombocytes (PCT, %), mean platelet 168 volume (MPV, fl), platelet distribution width (PDW, %), white blood cell count (WBC, 169 m/mm³), lymphocytes (LYM, %), monocytes (MON, %), neutrophils (NEUT,%), 170 eosinophils (Eos, %), basophils (Bas, %). For the serum blood chemistry, the 171 concentrations of total protein (TP, g/dl), glutamate oxaloacetate transaminase (GOT, 172 UI/L), blood urea nitrogen (BUN, mg/dl), albumin (g/dl), urea (mg/dl) and cholesterol 173 (mg/dl) were measured using an automated system photometer (Screen Master 174 Touch, Hospitex Diagnostics, Sesto Fiorentino, FI, Italy). 175

For immune indices, the serum electrophoretic patterns were obtained using a semi-176 automated agarose gel electrophoresis system (Sebia Hydrasys, EVRY, France) to 177 determine serum protein. Serum lysozyme was measured with a lysoplate assay, 178 carried out in a moist incubator at 37°C for 18 min. The method is based on the lyses 179 of Micrococcus lysodeikticus in 1% agarose. The diameter of the lysed zones was 180 measured with a ruler and compared with the lysed zones of a standard lysozyme 181 preparation (Sigma Aldrich, Milan, Italy). The value was expressed as µg/ml 182 (Osserman and Lawlor, 1996). The haemolytic complement assay was carried out in 183 microtitre plates. The complement titre is the reciprocal of the serum dilution causing 184 50% lysis of red blood cells of rams. Its concentration was expressed as CH_{50%} 185 (Moscati *et al.*, 2008). 186

187

188 Performance of fattening rabbits

At the second parturition, forty weaned kits were randomly chosen from both C and E 189 does. Rabbits were allocated into individual wire cages (0.41 m long × 0.30 m wide × 190 0.28 m high) and randomly assigned to four equal-size experimental groups (n=20). 191 Two groups of rabbits were fed a growing commercial basal diet (C) while the 192 remaining two groups were fed the same diet supplemented with 3 g of EPAL powder 193 / kg diet (E). According to the maternal diet, the following experimental groups were 194 formed: CC group (rabbits fed the C diet and born from the C does), CE group 195 (rabbits fed the E diet and born from the C does), EC (rabbits fed the C diet and born 196 from the E does) and EE group (rabbits fed the E diet and born from the E does). The 197 chemical composition of the different diets is reported in Table 2. The diets were 198 completely exempt from medication (antibiotics or coccidiostat). Feed and water were 199 provided ad libitum. During the whole trial, temperature was maintained at 22±2°C 200

201 and a 16L: 8D photoperiod was applied. Health status was monitored daily from 202 weaning to 77 days of age.

Rabbits were weighed at 35, 49 and 77 day of age and the following performance
parameters were calculated: daily feed intake, daily weight gain and feed conversion
ratio at different periods of age.

- 206
- 207 Chromatographic identification of Echinacea ingredients

208 Chemicals

Echinacoside (purity 98%), chlorogenic acid (purity \geq 95%), HPLC-MS and analytical

- grade solvents were purchased from Sigma-Aldrich (Milan, Italy).
- 211 Extraction procedure

500 mg of dry ground EPAL roots, were sonicated for 10 min with 10 ml of a mixture
of MeOH/H2O (70/30) three times. The resulting total extract (30 ml) was filtered and
analyzed by UHPLC-PDA-MS/MS system.

215 HPLC Analysis

EPAL extract analyses were carried out on a Shimadzu Nexera X2 system equipped 216 with a photodiode detector SPD-M20A in series to a triple quadrupole Shimadzu 217 LCMS-8040 system provided with electrospray ionization (ESI) source (Shimadzu, 218 Dusseldorf Germany). An Ascentis® Express C18 column (150 mm x 2.1 mm i.d., 2.7 219 µm particle size), (Supelco, Bellefonte, PA) was used (operated at 30°C). The mobile 220 phase consisted of 0.1% formic acid in water (A) and 0.1% formic acid in acetonitrile 221 (B), at a flow rate of 0.4 ml min⁻¹. Polyphenols elution was achieved using the 222 following linear gradient: starting condition, 95% A, 5% B; 3 min, from 5 to 15% B; 17 223 min, from 15 to 100% B; 5 min and 100% B for 2 min. The injection volume was 5 µl. 224 UV spectra were acquired in the 210-450 nm wavelength range. The identification of 225

the components was based on the co-injection of pure standards and on their UV spectra and mass spectral information in both positive and negative ionization mode (respectively, ESI+ and ESI-).

229 Quantification of Echinacoside: A standard stock solution (1mg/ml) of Echinacoside 230 was prepared in methanol and stored at -18°C. Suitable dilutions of the standard 231 stock solution in methanol/water (1/10) were prepared to obtain final concentrations 232 from 10 to 100 mg/ml. Calibration curve was built by analysing the resulting standard 233 dilutions three times by HPLC-PDA.

234

235 Statistical analysis

Statistical analyses were performed using SPSS software package (IBM SPSS, 236 2012). Data concerning the reproductive parameters from the first to the fifth 237 reproductive cycles were combined and analyzed in a single dataset. Statistical 238 analyses for significant differences in reproductive performance between the control 239 and Echinacea groups were performed using a Student's t-test. Mortality, kindling 240 rate and prolificacy were analyzed using Chi-square test. The effect of dietary 241 treatments on blood indices and immune parameters across three gestational periods 242 (day 0, day 14, day 28) was statistically analyzed with a mixed between-within 243 subjects model (GLM Repeated Measures). Performance of the fattening rabbits was 244 analyzed using a one-way ANOVA with group as fixed factor. Duncan's New Multiple 245 Range test was used for post-hoc comparisons. The significance was declared at 246 *P*<0.05. 247

248

249 **Results**

250 HPLC profile of EPAL

The HPLC profiles of EPAL root extract are shown in Figure 1. The analysis identified the presence of caftaric acid, cichoric acid, chlorogenic acid and Echinacoside which specifically characterized EPAL species (Hu and Kitts, 2000; Speroni *et al.*, 2002; Barrett, 2003). Chromatographic analysis was reported to find 0.37 % Echinacoside.

Echinacoside was found to be the main caffeic acid derivative in EPAL extract, 255 responsible for the immunostimulatory action of Echinacea extracts (Hu and Kitts, 256 2000; Pellati et al., 2005; Dalby-Brown et al., 2005). Echinacoside has been studied 257 for its antioxidant, anti-inflammatory and cicatrizing activities (Speroni et al., 2002). 258 However, a purified phytochemical does not imitate the immunological effects of 259 whole plant extracts. It appears that the immunopharmacological activities of 260 Echinacea depend on a combination of several active compounds (Randolph et al., 261 2003). 262

263

264 Reproductive performance

Reproductive performances of the first five reproductive cycles are reported in Table 3. There were no significant differences between groups for any of the studied parameters. Numerical and overall productivities calculated during the five cycles were: born alive, 1438 and 1471 kits; number of kits at day 35, 1229 and 1260 for control and E groups, respectively.

270

271 Haematological findings

The haematological parameters of pregnant rabbits are reported in Table 4. The results indicated a significant (P<0.05) effect of treatment and gestational day on some haematological parameters. The control group was significant higher than the treatment group for basophils cells (0.55 and 0.29 %, respectively; P=0.049). Gestational day significantly affected RBC, Hb, HCT, MCV, MCH, MCHC, RDW, MPV, PDW, WBC, LYM, NEUT and Eos (P<0.05). For any studied variables, no significant interaction between treatment and gestational period was reported except for NEUT (P=0.033).

No significant effect of gestational day or treatment was observed on blood serum chemistry. Regarding the immune parameters, no significant differences were observed between groups; while a significant effect of gestational day was observed for lysozymes (P=0.014). The higher concentration of lysozymes was observed in day 14 of gestation in comparison with days 0 (+32.7%) and 28 (+318.3%) (6.02 *vs* 7.99 *vs* 1.91; for 0, 14 and 28 days respectively).

286

287 Fattening rabbit performance

The results of fattening rabbits performance are illustrated in Table 5. For all studied variables, no statistically significant differences were reported amongst the experimental groups (P>0.05). In addition, regarding the health status, no illness and death were observed during the fattening period.

292

293 Discussion

294 Reproductive performance

Body weight of does at kindling, kindling rate, litter size at birth, at days 21st and 35th of age, and the mortality of kits did not differ between the two groups. This indicates that *Echinacea* supplements in does' diets did not exert a promoting effect on reproductive function when administered at 3 g EPAL/kg of diet. Our results differ from those obtained in mice by Barcz *et al.* (2007) who found that two *Echinacea* drugs (*Esberitox* and *Echinapur*) lowered the number of embryos in one litter, even if the results were on the edge of statistical significance. During murine pregnancy, *Echinacea purpurea* reduced the number of viable foetus (Chow *et al.*, 2006). A prospective study suggested that the use of *Echinacea* in pregnancy during organogenesis is not associated with an increased risk of major malformations (Gallo *et al.*, 2000). Further theoretical evidence via an expert panel on botanical medicine reported that oral consumption of *Echinacea* in recommended doses appeared safe and effective to use during pregnancy (Perri *et al.*, 2006).

308

309 Haematological findings

Blood parameters in rabbits are used as an aid for the clinical diagnosis of metabolic, 310 infectious and parasitic diseases and to assess animal condition. A variety of factors 311 312 can affect animal haematological and biochemical parameters, including breed, gender, diet, age, reproductive status and seasonal variations (Ozegbe, 2001; Wells 313 et al., 1999). The haematological and biochemical parameters of this study were 314 within normal ranges for rabbit species (Archetti et al., 2008; Özkan et al., 2012). The 315 application of Echinacea extract should booster immunological reactivity and should 316 317 contribute to improve health status (Böhmer et al., 2009). In the present trial, EPAL had no influence the heamatological and health status of rabbit does. The change in 318 blood coagulation-related parameters during the later stage of gestation is a common 319 320 physiological response for the protection against excessive haemorrhage or for the preservation of the homeostasis at parturition (Mizoguchi et al., 2010). In our study, 321 the modulation of RBC and HCT may be related to physiological anemia resulting 322 323 from haemodilution (Ozegbe, 2001). Watery supplementation with Echinacea purpurea extract induced higher results of Hb, PCV and RBC in growing rabbits 324 (Ahmed et al., 2008). Likewise, a study by Chow et al. (2006) found an increase in 325

RBC in pregnant mice when fed Echinacea purpurea. In addition, the increment of 326 erythropoietin level (glycoprotein hormone which controls erythropoiesis) has been 327 reported in Echinacea purpurea-treated men. This should support the RBC increment 328 deriving from the supply of phyto-additives (Whitehead et al., 2007). On the other 329 hand, Maass et al. (2005) did not find any significant difference for these parameters 330 in sows, piglets and grower/finisher pigs that received dried Echinacea purpurea herb 331 as feed additive in their diets. Differences concerning plant species tested (EPAL vs 332 Echinacea purpurea), preparation methods (raw material vs extraction), physiological 333 status (pregnant vs non-pregnant) and species (rabbit vs swine, mice and human 334 beings) could explain these contrasting results. An author showed that WBC 335 parameters increased during the whole period of gestation in pregnant women 336 (Cincotta et al., 1995), in rabbit does (Haneda et al., 2010) and also in rats (DeRijk et 337 al., 2002). Cundell et al. (2003) found a significant increase of lymphocytes after one 338 week in rats fed with dried Echinacea preparations. A higher proliferation rate of 339 spleen lymphocytes in EPAL supplemented mice has been reported in an in vitro 340 study, but the haematology indices were not influenced (Zhai et al., 2007). The 341 increase in WBC generally is a good indicator of immunity efficiency increase 342 (Wieslaw et al., 2006). In our study, the effect of EPAL was observed only for Bas. 343 According to other studies, this effect may be related to its phytochemically active 344 constituents of EPAL (Hu and Kitts, 2000; Pellati et al., 2005; Dalby-Brown et al., 345 2005). 346

With respect to blood serum chemistry, no significant difference was observed in total protein. In contrast, Wells *et al.* (1999) reported a decrease in total protein and albumin in pregnant rabbits and this is thought to reflect the increased blood volume.

Innate immunity has an important role to prevent the infection as first-line defence 350 and also contributes antigen-presenting cells that activate the adaptive immune 351 response, which is specific and powerful (Tizard, 2013). Dietary supplementation with 352 Echinacea can stimulate the innate immunity by increasing cytokine production 353 (Hwang et al., 2004) and phagocyte-stimulation (Böhmer et al., 2009). Lysozymes 354 and the complement system are interesting indicators to study the innate immune 355 function. In our experiment, only lysozyme results showed a time related change. It 356 must be highlighted that our work was performed in a standard environment without 357 infection, stress or other factors influencing immune responses. Therefore, the 358 experimentation in normal conditions may hardly result in a significant effect on 359 immunity despite the supplementation with an immunomodulating agent. 360

361

362 *Fattening rabbit performance*

Growth performance of Echinacea supplemented groups did not showed significant 363 differences. Our results differ from Arafa et al. (2010) who found, in a similar study 364 using Echinacea purpurea at 130 mg/kg body weight, a significant decrease in 365 mortality rate and an increase of live weight in 6-week-old growing rabbits fed E diets 366 (P<0.05). Usually, dietary herb supplementation leads to an improvement of the 367 flavour, which accounts for an increase of feed ingestion and better performance 368 (Franz et al., 2010; Christaki et al., 2012). Ahmed et al. (2008) highlighted a 369 significant improvement of final body weight, daily weight gain and feed conversion 370 ratio in growing rabbits to which were orally given in liquid 7.5 mg of Echinacea 371 purpurea extracts/kg body weight and day. However, the outcomes of above reported 372 references are not fully comparable with our trial due to some dissimilarities in 373 experimental plans concerning: tested Echinacea species, concentration of the 374

supplement, administration route (oral by liquid mixture), supplement preparation
 (extraction) and supplemented periods in doe's diet

Generally, mixtures of *Echinacea purpurea*, *Echinacea angustifolia* and EPAL are used in human medicine and animal production. To this regard, positive outcomes on productive performance were reported in rabbits with *Echinacea purpurea* addition (Ahmed *et al.*, 2008; Arafa *et al.*, 2010), whereas studies conducted with other livestock species did not find any improvement (Hermann *et al.*, 2003; Maass *et al.*, 2005; Böhmer *et al.*, 2009; Sahin *et al.*, 2012).

In conclusion, there is no evidence that diets supplemented with EPAL cause any beneficial effects in normal management condition. Nonetheless, further studies are suggested in order to evaluate the effect of *Echinacea pallida* on animal performance and to study the relation between its active components and physiological functions.

387

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| | Does diet (| from artificial | Does diet (from 21 days | | | |
|------------------------------------|--------------|------------------|-------------------------|----------------|--|--|
| | insemination | to 21 days after | after parturiti | ion to 35 days | | |
| | partu | urition) | after pa | arturition) | | |
| | Control | Treatment | Control | Treatment | | |
| Chemical composition ¹ | | | | | | |
| Dry matter (DM) | 89.3 | 90.2 | 89.9 | 89.9 | | |
| Crude protein (% DM) | 18.7 | 18.8 | 17.5 | 17.2 | | |
| Ether extract (% DM) | 2.6 | 2.9 | 4.5 | 4.6 | | |
| NDF (% DM) | 35.0 | 33.7 | 32.4 | 32.2 | | |
| ADF (% DM) | 22.4 | 22.2 | 17.5 | 17.9 | | |
| ADL (% DM) | 5.5 | 5.7 | 5.4 | 5.4 | | |
| Ash (% MS) | 9.5 | 9.5 | 7.5 | 7.9 | | |
| Starch (% DM) | 26.2 | 27.2 | 17 | 17.4 | | |
| Echinacea pallida (g/kg) | 0 | 3 | 0 | 3 | | |
| Minerals and vitamins ² | | | | | | |
| Calcium (% DM) | 0.9 | 0.9 | 1 | 1 | | |
| Lysine (% DM) | 0.8 | 0.8 | 0.7 | 0.7 | | |
| Phosphorus (% DM) | 0.5 | 0.5 | 0.4 | 0.4 | | |
| Methionine (% DM) | 0.3 | 0.3 | 0.4 | 0.4 | | |
| Sodium (% DM) | 0.3 | 0.3 | 0.3 | 0.3 | | |
| Vitamin A (UI/kg) | 12.5 | 12.5 | 12.5 | 12.5 | | |
| Vitamin D3 | 1.2 | 1.2 | 1.2 | 1.2 | | |
| Vitamin E | 100 | 100 | 100 | 100 | | |
| Ferrous carbonate (mg/kg) | 662 | 662 | 704 | 704 | | |
| Manganese oxide (mg/kg) | 195 | 195 | 209 | 209 | | |
| Zinc oxide (mg/kg) | 186 | 186 | 186 | 186 | | |
| Copper sulfate (mg/kg) | 98 | 98 | 98 | 98 | | |
| Potassium iodide (mg/kg) | 2.4 | 2.4 | 2.5 | 2.5 | | |
| Sodium selenite (mg/kg) | 0.6 | 0.6 | 0.6 | 0.6 | | |

¹The experimental diets were analyzed by the laboratory of the Department of Agricultural,

534

4 Forest and Food Sciences, Turin, Italy. ²These data were provided by the Ferrero Mangimi

535 S.p.A, (Farigliano CN, Italy), which formulated and prepared the experimental diets.

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537

538 **Table 2** Growing rabbits diets composition

| | Diets ² | | |
|------------------------------------|--------------------|-----------|--|
| | Control | Treatment | |
| Chemical composition ¹ | | | |
| Dry matter (DM) | 89.8 | 89.8 | |
| Crude protein (% DM) | 17.1 | 17.3 | |
| Ether extract (% DM) | 3 | 3 | |
| NDF (% DM) | 39.4 | 39.6 | |
| ADF (% DM) | 23.7 | 24 | |
| ADL (% DM) | 6.6 | 6.6 | |
| Ash (% DM) | 9.7 | 10.4 | |
| Starch (% DM) | 12 | 12.3 | |
| Echinacea pallida (g/kg) | 0 | 3 | |
| Minerals and vitamins ² | | | |
| Calcium (% DM) | 1 | 1 | |
| Lysine (% DM) | 0.7 | 0.7 | |
| Methionin (% DM) | 0.4 | 0.4 | |
| Phosphorus (% DM) | 0.4 | 0.4 | |
| Sodium (% DM) | 0.3 | 0.3 | |
| Vitamin A (UI/kg) | 12.5 | 12.5 | |
| Vitamin D3 | 1.2 | 1.2 | |
| Vitamin E | 100 | 100 | |
| Ferrous carbonate (mg/kg) | 662 | 662 | |
| Manganese oxide (mg/kg) | 195 | 195 | |
| Zinc oxide (mg/kg) | 186 | 186 | |
| Copper sulfate (mg/kg) | 98 | 98 | |
| Potassium iodide (mg/kg) | 2.5 | 2.5 | |
| Sodium selenite (mg/kg) | 0.57 | 0.57 | |

¹The experimental diets were analyzed by the laboratory of the Department of Agricultural,
 Forest and Food Sciences, Turin, Italy. ²These data were provided by the Ferrero Mangimi

541 S.p.A, (Farigliano CN, Italy), which formulated and prepared the experimental diets.

| | Control group | Echinacea group | Standard | P-value |
|---|---------------|-----------------|---------------|--------------------|
| | | | error of mean | |
| | | | difference | |
| No. of does at first kindling | 50 | 50 | - | - |
| No. of does at fifth kindling | 37 | 38 | - | - |
| Mortality of does (%) | 26 | 24 | - | 0.817 ¹ |
| Body weight (LW), g | | | | |
| at first kindling | 3868 | 3869 | - | 0.982 |
| at fifth kindling | 4782 | 4770 | - | 0.929 |
| No. of kindled does/artificial insemination | 148 / 221 | 151 / 221 | - | - |
| Kindling rate,% | 67 | 68 | - | 0.760^{1} |
| Prolificacy | 8.78 | 8.88 | - | 0.852^{1} |
| Total born | 10.5 | 10.5 | 0.36 | 0.978 ² |
| Born alive | 9.72 | 9.74 | 0.37 | 0.945 ² |
| Stillborn | 0.78 | 0.76 | 0.18 | 0.907^{2} |
| Litter size | | | | |
| at 21d | 8.36 | 8.42 | 0.25 | 0.816 ² |
| at 35d | 8.30 | 8.34 | 0.26 | 0.877 ² |
| Litter weight (g) | | | | |
| at 21d | 2750 | 2747 | 101.22 | 0.981 ² |
| at 35d | 7023 | 7038 | 229.82 | 0.946 ² |
| Individual body weight (g) | | | | |
| at 21d | 329 | 326 | 3.80 | 0.495 ² |
| at 35d | 846 | 844 | 4.05 | 0.585 ² |
| Perinatal mortality (%) | 7.40 | 7.25 | - | 0.868 ¹ |
| Mortality (%) | - | - | | |
| 0-21d | 14 | 13.6 | - | 0.788 ¹ |
| 21-35d | 0.65 | 0.86 | - | 0.528 ¹ |

Table 3 Effects of pale purple coneflower (Echinacea pallida) dietary supplementation on reproductive performance of rabbit does

⁵⁴³ ¹: parameter analyzed by Chi-square test; ²: parameter analyzed by Student's t-test

545 **Table 4** Effects of pale purple coneflower (Echinacea pallida) dietary supplementation on blood and immune parameters of

546 pregnant rabbit does (n=8 per group)

| | Treatment | | Gestational day | | | Within subjects effects | | | Between subjects effects | |
|--------------------------|------------------|---------------------------|-----------------|--------|--------|----------------------------|-----------------------------------|-----------------|--------------------------|---------------------|
| | Control group | <i>Echinacea</i> group | 0 | 14 | 28 | Gestational | Gestational day | Root Mean | Treatment | Root Mear Square |
| No. of animals | 8 | 8 | 8 | 8 | 8 | day <i>P</i> -value | × Treatment <i>P</i> -value | Square Error | <i>P</i> -value | Error |
| Haematology | | | | | | | | | | |
| RBC (M/mm ³) | 5.80 | 5.50 | 5.38 | 5.96 | 5.62 | 0.025 | 0.963 | 0.422 | 0.145 | 0.265 |
| Hb (g/dl) | 11.99 | 11.40 | 10.96 | 12.27 | 11.86 | 0.013 | 0.992 | 0.885 | 0.274 | 1.912 |
| HCT (%) | 38.02 | 36.18 | 35.65 | 39.46 | 36.19 | 0.014 | 0.907 | 86.827 | 0.271 | 18.164 |
| MCV (fl) | 65.58 | 65.87 | 66.27 | 66.32 | 64.59 | 0.003 | 0.383 | 33.808 | 0.870 | 22.463 |
| MCH (pg) | 20.61 | 20.70 | 20.29 | 20.57 | 21.10 | 0.048 | 0.763 | 0.677 | 0.894 | 2.870 |
| MCHC (g/dl) | 31.53 | 31.46 | 30.69 | 31.06 | 32.74 | <0.001 | 0.553 | 0.932 | 0.770 | 0.439 |
| RDW (%) | 10.81 | 11.65 | 10.14 | 11.53 | 12.03 | <0.001 | 0.339 | 0.702 | 0.343 | 5.216 |
| PLT (m/mm ³) | 137.07 | 168.00 | 146.20 | 154.50 | 156.90 | 0.897 | 0.441 | 53.741 | 0.223 | 64.048 |
| PCT (%) | 0.09 | 0.11 | 0.09 | 0.10 | 0.11 | 0.432 | 0.304 | 0.032 | 0.158 | 0.045 |
| MPV (fl) | 6.74 | 6.83 | 6.51 | 6.55 | 7.29 | <0.001 | 0.380 | 0.253 | 0.620 | 0.460 |
| PDW (%) | 6.77 | 6.78 | 6.66 | 6.29 | 7.38 | 0.009 | 0.729 | 0.692 | 0.974 | 0.538 |
| WBC (m/mm ³) | 9.59 | 9.38 | 11.14 | 11.11 | 6.22 | <0.001 | 0.507 | 2.070 | 0.829 | 2.588 |
| LYM (%) | 14.97 | 14.57 | 15.41 | 12.56 | 16.34 | 0.011 | 0.803 | 2.541 | 0.850 | 5.688 |
| MON (%) | 6.53 | 5.91 | 6.62 | 5.49 | 6.54 | 0.052 | 0.663 | 1.055 | 0.533 | 2.606 |
| NEUT (%) | 76.87 | 78.17 | 76.87 | 80.49 | 75.21 | 0.009 | 0.033 | 3.396 | 0.657 | 7.717 |
| Eos (%) | 1.08 | 1.04 | 0.56 | 0.99 | 1.63 | <0.001 | 0.626 | 0.475 | 0.851 | 0.565 |
| Bas (%) | 0.55 | 0.29 | 0.54 | 0.44 | 0.28 | 0.092 | 0.671 | 0.249 | 0.049 | 0.300 |
| Blood serum chemistry | | | | | | | | | | |
| BUN (mg/dl) | 20.87 | 16.82 | 14.95 | 15.34 | 26.25 | 0.183 | 0.471 | 9.305 | 0.413 | 8.127 |
| GOT (UI/L) | 29.06 | 32.22 | 26.79 | 35.58 | 29.55 | 0.395 | 0.790 | 14.315 | 0.401 | 9.763 |
| Total Protein (g/dl) | 4.60 | 4.34 | 4.48 | 4.22 | 4.72 | 0.325 | 0.641 | 0.711 | 0.296 | 0.643 |
| Albumin (g/dl) | 2.91 | 2.91 | 2.68 | 2.87 | 3.18 | 0.109 | 0.191 | 0.494 | 0.971 | 0.391 |
| Urea (mg/dl) | 29.28 | 36.10 | 32.09 | 32.92 | 33.06 | 0.973 | 0.143 | 6.929 | 0.319 | 10.851 |
| Cholesterol (mg/dl) | 48.66 | 39.41 | 33.85 | 63.21 | 35.04 | 0.352 | 0.219 | 49,781 | 0.658 | 55.169 |

| Immune parameters | | | | | | | | | | |
|-------------------|-------|-------|-------|-------|-------|-------|-------|--------|-------|--------|
| Lysozymes (µg/ml) | 5.64 | 4.98 | 6.02 | 7.99 | 1.91 | 0.014 | 0.590 | 4.122 | 0.862 | 10.067 |
| Complement | 36.72 | 29.44 | 34.31 | 34.06 | 30.87 | 0.826 | 0.267 | 12.959 | 0.174 | 12.438 |
| Alfa1 (g/dl) | 0.14 | 0.16 | 0.20 | 0.09 | 0.18 | 0.234 | 0.117 | 0.155 | 0.746 | 0.182 |
| Alfa 2 (g/dl) | 0.28 | 0.19 | 0.32 | 0.19 | 0.19 | 0.176 | 0.304 | 0.170 | 0.164 | 0,158 |
| Beta 1 (g/dl) | 0.28 | 0.28 | 0.33 | 0.28 | 0.23 | 0.175 | 0.198 | 0.114 | 0.828 | 0.100 |
| Beta 2 (g/dl) | 0.39 | 0.38 | 0.38 | 0.36 | 0.42 | 0.557 | 0.687 | 0.118 | 0.801 | 0.134 |
| Gamma (g/dl) | 0.60 | 0.42 | 0.56 | 0.43 | 0.54 | 0.697 | 0.515 | 0.355 | 0.115 | 0.281 |

548 RBC: Red Blood Cells; Hb: Haemoglobin concentration; HCT: Haematocrit; MCV: Mean Corpuscular Volume; MCH: Mean Corpuscular Haemoglobin; MCHC:

549 Mean Corpuscular Haemoglobin Concentration; RDW: Red cell distribution width; PLT: Platelets; PCT: Relative volume of thrombocytes; MPV: Mean Platelet

550 Volume; PDW: Platelet distribution width; WBC: White Blood Cells; LYM: Lymphocytes; MON: Monocytes; NEUT: Neutrophils; Eos: Eosinophils; Bas:

- 551 Basophils; BUN: blood urea nitrogen; GOT: glutamate oxaloacetate transaminase.

Table 5 Effect of pre and postnatal dietary supplementation with pale purple coneflower (Echinacea pallida) on growth performance

of fattening rabbits (n=20 per group)

| | | DOD | D 1 | | | |
|----------------------------------|------|------|------------|------|------|---------|
| | CC | CE | EC | EE | RSD | P-value |
| Live weight (g) | | | | | | |
| At 35 day | 885 | 889 | 889 | 882 | 53.8 | 0.976 |
| At 49 day | 1713 | 1711 | 1745 | 1717 | 79.7 | 0.513 |
| At 77 day | 3031 | 2998 | 3107 | 3041 | 160 | 0.190 |
| Growth performance in 35-49 days | | | | | | |
| Daily feed intake (g per day) | 134 | 138 | 140 | 139 | 10.6 | 0.323 |
| Daily weight gain (g per day) | 59.2 | 58.7 | 61.2 | 59.6 | 3.58 | 0.160 |
| Feed conversion ratio | 2.28 | 2.36 | 2.29 | 2.35 | 0.15 | 0.200 |
| Growth performance in 49-77 days | | | | | | |
| Daily feed intake (g per day) | 176 | 178 | 181 | 181 | 10.8 | 0.478 |
| Daily weight gain (g per day) | 45.4 | 44.4 | 46.9 | 45.7 | 4.05 | 0.254 |
| Feed conversion ratio | 3.87 | 4.03 | 3.88 | 3.98 | 0.30 | 0.282 |

| | Growth performance in 35-77 days | | | | | | |
|----------|--|---------------------|------------------|-------------------|-----------------|-------------------|------------------|
| | Daily feed intake (g per day) | 162 | 165 | 168 | 168 | 11.5 | 0.368 |
| | Daily weight gain (g per day) | 49.9 | 49.0 | 51.6 | 50.2 | 3.11 | 0.082 |
| | Feed conversion ratio | 3.25 | 3.37 | 3.26 | 3.34 | 0.19 | 0.122 |
| 62 | CC: rabbits fed the C diet and born from | the C does, CE: rat | bits fed the E c | liet and born fro | m the C does, I | EC: rabbits fed t | the C diet and b |
| 63 | from the E does, EE: rabbits fed the E die | t and born from the | E does. | | | | |
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| 71 72 | | | | | | | |

- **Figure 1**
- 577 LC-PDA profile at 325 nm of Echinacea pallida (Nutt.) Nutt root extract at 325 nm.