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1 **Compilation of 29-year *postmortem* examinations identifies**
2 **major shifts in equine parasite prevalence from 2000**
3 **onwards**

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18

19 **Abstract**

20 Horses are infected by a wide range of parasite species that form complex communities.
21 Parasite control imposes significant constraints on parasite communities whose monitoring
22 remains however difficult to track through time. *Postmortem* examination is a reliable method
23 to quantify parasite communities. Here, we compiled 1,673 necropsy reports accumulated over
24 29 years, in the reference necropsy centre from Normandy (France). The burden of non-
25 strongylid species was quantified and the presence of strongylid species was noted. Details of
26 horse deworming history and the cause of death were registered. Building on these data, we
27 investigated the temporal trend in non-strongylids epidemiology and we determined the
28 contribution of parasites to the death of horses throughout the study period. Data analyses
29 revealed the seasonal variations of non-strongylid parasite abundance and reduced worm
30 burden in race horses. Beyond these observations, we found a shift in the species responsible
31 for fatal parasitic infection from the year 2000 onward, whereby fatal cyathostominosis and
32 *Parascaris* spp. infection have replaced death cases caused by *S. vulgaris* and tapeworms.
33 Concomitant break in the temporal trend of parasite species prevalence was also found within
34 a 10-year window (1998-2007) that has seen the rise of *Parascaris* spp. and the decline of
35 both *Gasterophilus* spp. and tapeworms. A few cases of parasite persistence following
36 deworming were identified that all occurred after 2000. Altogether, these findings provide
37 insights into major shifts in non-strongylid parasite prevalence and abundance over the last 29
38 years. They also underscore the critical importance of *Parascaris* spp. in young equids.

39

40 **Keywords:** horse; parasite; necropsy; *Parascaris*; *Strongylus*; cyathostomin; *Gasterophilus*;
41 *Anoplocephala*;

42

43 **1. Introduction**

44 Horses harbour complex macroparasite communities along their digestive tract, encompassing
45 among other *Gasterophilus* larval stages (bots), nematodes (mainly strongylids and ascarids)
46 and tapeworms (Anoplocephalidae). The control of this vast parasite community has largely
47 relied on the regular use of anthelmintic drugs. Following a few decades of treatments,
48 parasitologists have reported alteration of strongylid communities: *Strongylus* spp. prevalence
49 drastically decreased over time whereas cyathostomin infection has become a major issue
50 (Herd, 1990; Love and Duncan, 1991). This shift in species importance was derived from
51 independent scattered pieces of evidence in the field (Herd, 1990; Love and Duncan, 1991). It
52 was also associated with the development of drug resistant cyathostomin populations across
53 the world (Fischer et al., 2015; Sallé et al., 2017; Tzelos et al., 2017; Nielsen et al., 2018).
54 Independent evidence of ivermectin resistant *Parascaris* spp. populations have also been
55 accumulating in recent years (Lyons et al., 2008; Laugier et al., 2012; Martin et al., 2018).
56 However, the quantification of parasitic community evolution through time remains difficult and
57 limited reports have been made so far.

58 Necropsy is a reliable method for the diagnosis of infection with equine intestinal parasites,
59 especially those which are not detected (immature or larval stages) or may be underestimated
60 (tapeworms) by routine coproscopy (Lyons et al., 1981; Lyons et al., 1984; Proudman and
61 Edwards, 1992; Rehbein et al., 2013). Moreover, large parasites are easily recovered by
62 careful examination of bowel contents and intestinal mucosal surfaces. Data derived by
63 specific *post-mortem* examinations are hence more definitive than those obtained from other
64 methods of investigation. This technic provides an invaluable tool for measuring parasite
65 abundance and prevalence (Lyons et al., 1981). Using this technique, Tolliver et al. reported
66 observations on the composition of equine parasite communities and their respective
67 prevalence over a 28-year period in Kentucky (Tolliver et al., 1987). This work is, to our
68 knowledge, the most extensive time series to date but it was based on a subset of horses
69 selected to have patent strongyle infection (Tolliver et al., 1987). A decade later, further reports

70 suggested a decrease in the prevalence of *Gasterophilus* spp. but a steady rate of infection by
71 *Parascaris* spp. in young horses (Lyons et al., 2000). Most recent report from the same region
72 suggested an increased infection rate by bots and tapeworms but was in line with the decline
73 of *Strongylus* spp. (Lyons et al., 2018).

74 To our knowledge, there is no other example of comprehensive longitudinal analysis of equine
75 parasite communities in any other region than Kentucky. In France, only scarce data are
76 currently available on prevalence and intensity of equine intestinal parasites and they mainly
77 rely on coproscopy results (Laugier et al., 2012; Traversa et al., 2012). The equine necropsy
78 unit from the French agency for food, environmental and occupational health safety (ANSES,
79 France) has been performing around 300 necropsies a year since 1987, following the same
80 procedure.

81 Here, we present the parasitological data recorded in 1,673 young horses examined during a
82 29-year period to establish overall infection pattern in relation to host and environmental
83 factors. Using known deworming history, we identified likely cases of drug resistance, and we
84 relied on histo-pathological conclusions to determine parasite species contribution to the death
85 of examined horses. We also estimated changes in species prevalence and abundance over
86 time.

87

88 **2. Material and methods**

89 **2.1. Animals**

90 Material consisted of 1,740 young equids (2 to 24 months old) from 758 stud farms. Animals
91 were derived from an equine population of 8,564 equids that were submitted for routine
92 necropsy at the ANSES laboratory for animal health in Normandy from January 1987 to
93 December 2015. This source of dead horses is hence relatively unbiased toward drug
94 resistance. It also offers both a snapshot of non-strongylid communities and an assessment of
95 cyathostominosis and verminous arteritis in the same young equid population from Normandy.

96 We restricted our analysis to this subset of the total population as they more often suffer
97 parasitic infection, and to avoid heterogeneity in the data.

98 Foals younger than 2 months were excluded from the study because none of them harbored
99 the parasites searched. Necropsies were usually performed within a few hours after death but
100 some were delayed for periods up to 24 hours. About 90% of the equids examined were from
101 Normandy, which is the leading horse breeding region in France with roughly 10,500 foal births
102 per year.

103 For every animal, relevant metadata including age, sex, breed, date of death, original stud farm
104 were recorded. Indications about the last anthelmintic treatment administered were also
105 collected in most cases.

106 A few observations were discarded before analysis, including an individual whose age was
107 unknown and 32 records collected from geldings (as it was not possible to estimate any sex
108 effect with so few observations). In addition, 33 cases from a bankrupted stud farm and one
109 horse from a farm where no anthelmintic drug were given were also removed from the dataset
110 to avoid spurious signal linked to a lack of parasite management. In the end, 1,673 cases from
111 735 studs were retained for analysis.

112 Foals were generally born in spring thereby resulting in a collinear relationship between the
113 age at necropsy and the month when the necropsy took place. To account for this structure in
114 the data, age was rounded to the closest month and clustered into two categories, being either
115 less than or 1-year-old (foal) or older (yearling). Month of necropsy was binned into seasonal
116 categories: winter (January to March), spring (April to June), summer (July to September) and
117 autumn (October to December).

118

119 **2.2. Necropsy technique and parasitological procedures**

120 This study builds on data collected during routine necropsy. As a result, fine-grained
121 parasitological examinations could not be systematically carried out due to time and material
122 limitations. Parasite count was therefore performed for non-strongylid large parasites as these

123 species are relatively easy to detect upon visual inspection. In addition, routine examination
124 also included a research of encysted cyathostomin larvae in the mucosa and submucosa of
125 the large intestine and of migrating *Strongylus vulgaris* larvae in the major arteries of the
126 gastro-intestinal tract.

127 Throughout the study period, all necropsies were performed using the same complete protocol
128 (Rooney, 1970; Collobert, 1995) and were implemented by the same team. Particularly,
129 evisceration of the different parts of the digestive tract was carried out according to the
130 procedure described by Rooney (1970) and specific examinations were performed for parasite
131 recovery.

132 During evisceration, the stomach, small intestine, cecum and ascending colon were isolated
133 with ligatures. Then, every organ was opened with scissors and their content was collected
134 separately, spread on large trays and examined grossly for parasites. The mucosal surfaces
135 were gently flushed with tap water and visually inspected for attached parasites. The other
136 parts of the digestive tract such as the pharynx and the oesophagus were also examined.
137 Species were searched in the vicinity of their preferential niche. Therefore, special attention
138 was paid to bots in the oral cavity, pharynx, oesophagus and stomach; tapeworms were looked
139 for in the small intestine, ileocaecal junction, cecum and right ventral colon, whereas ascarids
140 and pinworms were looked for in the small intestine and in the ascending and small colon
141 respectively.

142 Parasite specimens recovered were identified as to family (Anoplocephalidae), subfamily
143 (Cyathostominae), genus (*Gasterophilus*, *Parascaris*, *Strongylus*) or species (*Oxyuris equi*)
144 according to their anatomical location and published keys and illustrations (Lichtenfels, 1975;
145 Jacobs, 1986; Price and Stromberg, 1987). *Gasterophilus* spp larval stage and species were
146 determined during two consecutive years only (from March 1990 to February 1992) using a
147 dissecting microscope (x30) and following appropriate guidelines (Wells and Knipling, 1938).
148 Regarding tapeworms, specimens recovered from the small intestine, caecum and right ventral
149 colon were preserved separately in 10% formalin in order to be microscopically examined later
150 for the purpose of specific identification (Euzéby, 1966; Lichtenfels, 1975). Tapeworms

151 recovered from the small intestine were all examined and identified. For tapeworms present in
152 other intestinal segments, the following protocol was implemented: when less than 100
153 specimens were counted, all the worms were identified. In cases of heavier infection (more
154 than 100 tapeworms), a 10% aliquot was examined.

155 The cranial mesenteric artery and its major branches were opened and evaluated for lesions
156 secondary to the migration of *Strongylus vulgaris* larvae. Adherent thrombi and granulation
157 tissue were removed by scraping the intimal surface; then the parasites were recovered by
158 dissecting carefully all these fragments and counted.

159 A special procedure was applied to detect cyathostomin larvae. At each site of the large
160 intestines where the presence of parietal larvae was suspected by careful visual inspection, a
161 10 cm² fragment of the digestive wall was removed, examined by mural transillumination
162 technique (Reinemeyer and Herd, 1986) and then dissected under a dissection microscope at
163 10 to 30X magnification to confirm the presence of cyathostomin larvae. The number of larvae
164 per cm² was recorded. This technic is less sensitive than artificial digestion and can only detect
165 large developing larvae (late third stage larvae, fourth stage larvae) and will miss early third
166 stage larvae (Chapman et al., 1999). Nevertheless, it was chosen as an optimum between
167 material capacities and detection sensitivity of developing cyathostomins. Fourth stage larvae
168 were searched for by visual inspection of the caeco-colic content, recovered and inspected
169 under a microscope to confirm their developmental stage from morphological criteria, i.e. cup-
170 shaped buccal capsule and no visible cuticle (Brianti et al., 2009).

171

172 **2.3. Determination of the cause of death**

173 The cause of death was determined according to horse clinical history (duration of the disease
174 and evolution, clinical signs and results of laboratory tests), observed lesions and the
175 epidemiological context. The same person was in charge of categorizing the cause of deaths
176 throughout the study period, thereby making observations comparable across the years.

177 Parasites were declared as the most likely cause of death when parasite recovery was
178 associated with the following lesions:

- 179 - *Parascaris sp.*: intestinal obstruction, intussusception or rupture, toxemia and allergic
180 shock following treatment in heavily infected foals (more than 30 worms);
- 181 - Tapeworms: presence of at least one tapeworm associated with ileal, ileo-caecal,
182 caeco-caecal and caeco-colic intussusception, thickening of the ileal wall with
183 obstruction, paralytic ileus at the ileocaecal valve;
- 184 - Larval cyathostominosis was suspected in case of extensive typhlocolitis including
185 mucosal congestion, oedema, ulceration and necrosis, along with the presence of
186 numerous encysted larvae (more than 10 larvae per cm²) and or hundreds of emerged
187 L4 larvae in the bowel content.
- 188 - Infection with *S. vulgaris* larvae was considered as the cause of death when arterial
189 infarction and necrosis of a bowel segment was diagnosed and was associated with
190 verminous arteritis and thromboembolism.

191

192 **2.4. Statistical analyses**

193 Statistical analyses were carried out with the R software v3.5 (R Core Team, 2016).
194 Parasitological data were analyzed following a binary outcome, *i.e.* infected or not, or as a
195 continuous trait that quantifies the severity of the infection. The binary trait was modeled using
196 logistic regression and a binomial link function, while raw worm counts were assumed to follow
197 a negative binomial distribution, which is common for over-dispersed count data. This latter
198 assumption was confirmed using the *fitdistrplus* v.1.0-14 package (Delignette-Muller and
199 Dutang, 2015) by visual inspections of scatterplots of observed quantiles against the
200 theoretical quantiles from three distributions, *i.e.* normal, Poisson and negative binomial
201 (supplementary Figure 1).

202 For both type of trait, models were built as the sum of known fixed effects, *i.e.* horse sex, breed
203 (French trotter, Thoroughbred, miscellaneous), age class (older than one year of age or not),

204 and the season at which the horse died. We also added a binary variable encoding the time
205 period, *i.e.* before or after the observed break in species prevalence through time. The break
206 in parasite prevalence occurring around the year 2000 was inferred after regression of their
207 estimated prevalence upon the year, using the segmented package (Muggeo, 2017). This
208 strategy was chosen to account for the temporal trends in species prevalence and abundance;
209 more complex mixed models including year as a random effect did not provide precise
210 estimates and were faced with convergence issues when combined with a negative binomial
211 link function. Fixed effects were subsequently kept or discarded by an AIC-based variable
212 selection using the *stepAIC()* function from the MASS package (Venables and Ripley, 2002).
213 This procedure aims at minimizing the residual variance while avoiding model overfitting.
214 Horse sex was never retained during the variable selection procedure.
215 The cause of death was registered and classified as a binary outcome, *i.e.* of parasitic origin
216 or not, and regressed upon horse breed and the season at which the horse was examined
217 using logistic regression. The prevalence of fatal parasite infection was regressed upon the
218 year of examination to establish whether it varied significantly between 1987 and 2015.
219 Mean estimates of the logistic regressions were exponentiated to obtain the relative risk
220 associated with each variable level.
221 Due to the very low prevalence of *Parascaris* spp. in yearlings, modelling of worm burden and
222 prevalence for these species was performed on the foal data only (n = 1,174 out of the 1,673
223 available observations).
224 Any test with *P*-value below 5% was deemed significant.

225

226 **3. Results**

227 **3.1. Overall infection pattern by non-strongylid species**

228 Average non-strongylid parasite burden and prevalence were in a lower range of values
229 (Figure 1). Only 14 horses harboured *O. equi* and this species was not considered further. Bots
230 were recovered from 409 out of the 1,673 equids examined *post-mortem* (24.4% prevalence,

231 95% c.i. : 22% - 26%). The number of *Gasterophilus* spp. per infected equid ranged between
232 1 and 889 (mean = 65.03 ± 90.46 and median = 35) and these were found in the stomach in
233 most cases (380 out the 385 cases with observations; 10 horses presented instar attached to
234 the oesophagus and five horses had larvae attached to their pharynx). Species determination
235 was performed between March 1990 and February 1992 on 4,650 larvae sampled from a
236 subset of 153 horses. At that time, *G. intestinalis* was the most prevalent species (37.9%)
237 followed by *G. nasalis* (12.4%) and *G. haemorrhoidalis* (0.7%). Tapeworms were found in 289
238 equids (17.2% prevalence, 95% c.i. : 15% - 19%), and were almost exclusively located in the
239 caecum (n = 224 cases). *A. magna* and *P. mamillana* were recovered in 2 horses each,
240 including a co-infection with *A. perfoliata* in both cases. *Parascaris* sp. was recovered from the
241 small intestine of 207 foals (17.6% prevalence, 95% c.i. : 15.5% - 19.9%) with an average
242 abundance of 95 worms recovered (ranging from 1 to 1605 individuals).

243 Co-infection by three non-strongylid species rarely occurred (n = 20), but 12.4% of the
244 examined cases presented two non-strongylid species. In that latter case, parasites were twice
245 as likely to be responsible for the death of the horse (14.4% of cases against 6.8% in the total
246 population of cases).

247 The youngest foals with gastro-intestinal macroparasites were two months of age. Bots and
248 *Parascaris* spp. were found in respectively 6 and 10 foals of that age with counts ranging from
249 1 to 34 and 1 to 75 individuals for bots and *Parascaris* nematodes respectively. The youngest
250 foals infected with tapeworms were four months of age (n = 3) and harboured between 2 and
251 34 cestodes.

252 Parasite burden and prevalence followed seasonal fluctuations (Figure 1). *Parascaris* spp.
253 were significantly more abundant in summer and autumn (averaged corrected burden of $34 \pm$
254 1.32 and 19.5 ± 1.42 nematodes/horse, $P < 10^{-4}$ and 2×10^{-3}), with a peak of infection in
255 August. The same pattern was found for prevalence, whereby the highest frequency of
256 infection was observed in autumn (odd ratio = 3.43, 95% c.i. = 2.12 - 5.57; $P < 10^{-4}$). Bots and

257 tapeworms hit their highest abundance later in the second half of the year, *i.e.* in autumn ($P <$
258 10^{-4}) and winter ($P = 0.018$). During their respective most favourable season, bot and tapeworm
259 prevalences was 14.67- (95% c.i. : 9.34 - 23.03) and 4.23-fold (95% c.i.: 2.77 - 6.46) as high
260 as that observed in spring, respectively. Of note, tapeworms were more frequently found in
261 yearlings than in foals (odd ratio = 2.54, 95% c.i. = 1.94 - 3.32) whereas horse age category
262 neither contributed to bot burden variance nor to their prevalence variance.

263 Horse breeds were variously infected by non-strongylid parasites. Equids fell into three breed
264 types: Thoroughbred (TB), French Trotter (FT) and miscellaneous (MISC) that encompassed
265 French Saddlebreds (66.9%), other sport horses (6.6%), ponies (13.3%), Arabians (5.9) and
266 draft horses (4.8%). Substantial variation was found in bot abundance and prevalence across
267 the considered breed categories: Thoroughbred horses were significantly twice less likely to
268 be infected by bots as miscellaneous horses (difference in relative risk = 0.45, 95% c.i. = 0.32
269 - 0.64; $P < 10^{-4}$). In that case, parasite abundance was lower (8.58 ± 1.17 bots on average, P
270 = 9.8×10^{-3}) in Thoroughbred horses than for the two other breeds (13.46 ± 1.16 and $18.54 \pm$
271 1.29 bots on average for French trotter and miscellaneous horses respectively). Thoroughbred
272 horses also displayed lower tapeworm burden on average (average burden of 11 ± 1.24
273 cestodes vs. 20 ± 1.24 and 39 ± 1.39 in French trotters and miscellaneous horses). However,
274 their infection rate was not significantly different from that observed in other breeds ($\chi^2 = 2.63$,
275 $d.f. = 2$; $P = 0.27$). No difference in *Parascaris* spp. ($\chi^2 = 0.92$, $d.f. = 2$; $P = 0.63$) infection rate
276 was found between the three breed types considered.

277

278 **3.2. A shift in parasite species causing the death of young horses** 279 **through time**

280 Out the 1,673 horses, most of them died spontaneously ($n = 1347$) whereas the remainder
281 were euthanized by a veterinarian ($n = 326$). Overall, the cause of death was ascertained in
282 93.4% of horses ($n = 1563$), suspected for 92 cases or remained unknown in 18 cases.

283 Parasite were identified as being responsible for the death of 111 horses and highly suspected
284 for 3 additional horses (Figure 2). Out of these, cyathostomiasis was the most frequent cause
285 of death (n = 38), followed by caeco-colic invagination caused by *Anoplocephala* sp. infection
286 (n = 25). Thrombo-embolic disease caused by *S. vulgaris* (n = 22) and fatal *Parascaris* spp.
287 infection (n = 19) were the main remaining causes of parasitic death.

288 The annual proportion of death caused by parasites remained relatively constant ($6.5\% \pm 3.8\%$
289 of total deaths) throughout the considered 29 years ($F_{1,27} = 0.34$; $P = 0.56$). It reached its
290 highest in 2010 (18% of young horses necropsied) but was null in 2013. The relative risk of
291 fatal parasitic infection was higher late in the year (4.1- and 6.4-fold increase in relative risk in
292 autumn and winter respectively, $P < 10^{-4}$). It was also significantly reduced in race horses (odd
293 ratios of 0.37 and 0.39, $P < 10^{-4}$ for both Thoroughbreds and French trotters respectively). In
294 miscellaneous horses, cyathostomiasis represented more than half of total deaths of parasitic
295 origin (29 out of 38 cases) but this affection was less often seen in race horses (5 and 4 cases
296 out of 31 Thoroughbreds and 37 French trotters respectively; Figure 2). French trotters were
297 however more subject to fatal infection by tapeworms and *S. vulgaris* infection (Figure 2).

298 Of note, the yearly number of deaths caused by parasitic infection significantly increased after
299 2000 (2.53 ± 0.64 cases more, $P = 10^{-3}$; supplementary Figure 2). A shift in the species
300 responsible for the death of horses was also found from 2000 onward, whereby *S. vulgaris* and
301 tapeworms have been progressively replaced by cyathostomins and *Parascaris* spp. in more
302 recent times. *S. vulgaris* and tapeworms were responsible for 4.53 ± 0.92 ($P < 10^{-4}$) more cases
303 per year before 2000.

304

305 **3.3. Persistence of gastro-intestinal helminths in recently dewormed** 306 **horses**

307 Complete deworming history including the date and class of the last anthelmintics used for
308 deworming was available in 647 cases, 552 of which had been dewormed within the last 90

309 days. We found five cases (one French trotter, four Thoroughbred horses) of patent *Parascaris*
310 spp. infection in foals that had been treated with ivermectin within the last 30 days before
311 necropsy (4 to 22 days before death). These cases were noticed between 2004 and 2010. Two
312 foals died because of *Parascaris* spp. mediated intestinal perforation, but the three others had
313 non-parasitic causes of death.

314 *Parascaris* spp worms were found in foals treated with pyrantel two or six days before necropsy
315 in 1999 and 2015 respectively. The former French trotter died from enterotoxaemia
316 consecutive to deworming, while the latter Thoroughbred was euthanized because of a canon
317 fracture.

318 A last case of patent *Parascaris* sp. infection was noticed on a 7.5 month-old Thoroughbred
319 foal that had been drenched with fenbendazole four days before its death but harboured 54
320 worms.

321 An 18-month old Thoroughbred horse euthanized for a jaw lymphosarcoma in 2012, exhibited
322 836 *A. perfoliata* whereas he had been treated with a mixture of ivermectin and praziquantel
323 45 days before.

324

325 **3.4. An increased prevalence of *Parascaris* spp. from 2008 onward**

326 In relationship with the observed shift in species responsible for the death of young equids, we
327 quantified the temporal variation of parasite prevalence and abundance across the 29-year
328 period (Figure 3, supplementary Table 1). Breakpoints in non-strongylid prevalence were found
329 to occur within a ten-year period around 2000, *i.e.* 1998, 2005 and 2007 for bots, tapeworms
330 and *Parascaris* spp. respectively (Figure 3).

331 This analysis revealed a 1.97-fold increase in the prevalence of *Parascaris* spp. infection after
332 2007 (95% c.i. = 1.41 - 2.75; $P < 10^{-4}$). On average, 2.2 as many worms were observed in foals

333 after 2007 relative to pre-2007 observations ($P = 0.03$). This suggests that following 2007, foals
334 were significantly more often infected by *Parascaris* spp. and had increased worm loads.

335 An opposite pattern was found for bots and tapeworms (Figure 3). A break occurred in bots
336 prevalence from 1998 onward, that resulted in a 1.35-fold (95% c.i.: 1.33 - 2.18; $P < 10^{-4}$)
337 reduction of its infection rate. This trend was also conserved for the abundance of bots found
338 upon necropsy, with average count shifting from 17.8 ± 1.18 to 9.58 ± 1.15 after 1998 ($P = 2.3$
339 $\times 10^{-3}$). A similar significant reduction in the frequency of infection was found for tapeworm after
340 2005 (odds ratio = 0.62; 95% c.i.: 0.46 - 0.82; Figure 3), but their abundance was not
341 significantly altered through time ($\chi^2 = 0.12$, $d.f. = 1$; $P = 0.73$).

342

343

344 **4. Discussion**

345 Our survey provides one of the most comprehensive long-term surveys of equine gastro-
346 intestinal parasite dynamics. It is similar to a previous extensive report of 513 *postmortem*
347 examinations performed between the mid-1950's and 1983 in the USA (Tolliver et al., 1987).
348 These horses had been however chosen because of their patent strongylid infection and the
349 authors had limited information regarding their deworming history (Tolliver et al., 1987). This
350 latter piece of information is difficult to obtain in field conditions and is often missing in
351 *postmortem* examination (Lyons et al., 2000; Lyons et al., 2018) or abattoir surveys (Rehbein
352 et al., 2013). In some studies, specific parasite species are searched for in a subset of
353 individuals (Lyons et al., 2000). Here, we analyzed the long-term dynamics of parasite
354 population in young horses, using the same examination protocol and the relevant background
355 for each horse. The working subset of young animals reflected the diversity of equine
356 production in Normandy. Indeed, horses were coming from 25% of the 2,981 stud-farms
357 present in Basse-Normandy in 2014 (Anonymous, 2015). In addition, the diverse aetiologies
358 underpinning the death of young equids suggest that these horses were not biased towards

359 stud farms facing major issues in parasite control. A sampling bias remains however possible,
360 as it is likely that all dead horses in the region were not sent for necropsy.

361 The data collected on this subset of young horses highlighted a seasonal pattern in non-
362 strongylid parasite abundance and prevalence. In agreement with previous reports from
363 temperate areas, bots (Lyons et al., 1985; Price and Stromberg, 1987; Mfitlodze and
364 Hutchinson, 1989; Lyons et al., 1994; Bucknell et al., 1995; Höglund et al., 1997; Lyons et al.,
365 2000; Rehbein et al., 2013) and tapeworms (Benton and Lyons, 1994; Bucknell et al., 1995;
366 Nilsson et al., 1995; Meana et al., 2005; Rehbein et al., 2013; Tomczuk et al., 2015) were more
367 abundant and prevalent in autumn and winter seasons. This suggests that the subset of young
368 equids, that were examined throughout the year, was a good proxy to investigate the regional
369 parasite community dynamics. However, a peak of *Parascaris* spp. abundance was found in
370 August and highest prevalence occurred in autumn. This slight seasonal disconnection
371 between the occurrence of tapeworms and *Parascaris* spp may contribute to explain the limited
372 extent of co-infection between the three non-strongylid species types. A similar seasonality
373 was found in Northern Queensland (Australia), whereby *Parascaris* spp. infection was more
374 prevalent in wetter months (Mfitlodze and Hutchinson, 1989). This finding is in contrast with
375 multiple reports that did not find any evidence of a seasonal pattern (Lyons et al., 1994;
376 Bucknell et al., 1995; Rehbein et al., 2013; Fabiani et al., 2016) and could result from the
377 collinearity between the season when necropsies were performed and foal age. To this regard,
378 the median foal age in August, when *Parascaris* spp. were the most abundant, was 4 months
379 of age, which corroborates recent report (Fabiani et al., 2016).

380 Our prevalence estimates for bots and tapeworms were in the lower range of previously
381 reported values, that varied between 15% (Lyons et al., 2000) to 94% (Tolliver et al., 1987) for
382 bots and 30% (Mfitlodze and Hutchinson, 1989) to 80% for tapeworms (Benton and Lyons,
383 1994). This certainly reflects the important contribution of race horses to our dataset (84%), as
384 these are usually subjected to intensive deworming programs. For instance, a 2013-survey
385 across eight French Trotter studs revealed that foals were given eight anthelmintics a year
386 (*Sallé et al.*, unpublished observations).

387 Of note, a significant break in non-strongylid prevalences has occurred within a ten-year
388 window ranging from 1998 to 2008, whereby *Parascaris* spp. arose in contrast to bots and
389 tapeworms that suffered strong reduction in their respective prevalences. The sharp decline of
390 tapeworm prevalence followed closely the release of praziquantel between 2001 and 2005,
391 commercialized either alone (marketing authorization number FR/V/8052367 3/2001) or
392 combined with ivermectin (marketing authorization number FR/V/1889939 3/2004) or with
393 moxidectin (marketing authorization number FR/V/3281212 3/2005). The increased
394 awareness of the association between tapeworm infection and clinical intestinal disease in
395 horses in the 1990's (Pearson et al., 1993; Proudman and Edwards, 1993; Proudman et al.,
396 1998) has also certainly contributed to the implementation of a tapeworm-killing treatment in
397 late fall or winter. At that time, tapeworm control relied on the off-label use of niclosamide (100
398 mg/kg) or a double dose of pyrantel embonate. On the contrary, the decrease in bot prevalence
399 did not match the first release of ivermectin in 1983 (marketing authorization number
400 FR/V/6151318 9/1983). Their decline was however lower than the 85% drop-off in *G.*
401 *intestinalis* prevalence found between 1980 and 2000 in Kentucky (Lyons et al., 2000).
402 Because species determination was performed during two years only, it was not possible to
403 ascertain whether shifts in bots communities occurred. The observed decline remains hence
404 difficult to explain with available data.

405 We also identified a significant shift in the species responsible for the death of young horses.
406 Fatal tapeworm and *S. vulgaris* infections strongly declined after 2000, before a rise in
407 *Parascaris* spp. and cyathostomin mediated deaths occurred. In the lack of farm management
408 data or any climatic trend, a definitive explanation remains elusive. The decrease in fatal
409 tapeworm infection is likely connected to its reduced prevalence starting in early 2000's. The
410 *S. vulgaris* decline has been reported since the 1990's from various strands of evidence (Herd,
411 1990), although this is, to our knowledge, the first longitudinal quantification of this
412 phenomenon. This observation is in strong contrast with recent observations from
413 Scandinavian countries where increased prevalence of *S. vulgaris* was associated with
414 evidence-based drenching regimens (Nielsen et al., 2012; Tydén et al., 2019). France,

415 Denmark and Sweden fall under the same European regulation that imposes that anthelmintic
416 drugs are delivered upon prescription by a veterinarian (Anonymous, 2001). However, the drug
417 can be delivered without any coprological analysis in Sweden and France as opposed to the
418 Danish setting (Anonymous, 1998). The limited uptake of evidence-based drug treatment in
419 combination with the significant proportion of breeders buying anthelmintic drugs on their own
420 in France (Sallé et al., 2017) is likely to explain the reduction in *S. vulgaris* prevalence.

421 Of note, cyathostomiasis has remained the most frequent aetiology in death cases of parasitic
422 origin. The miscellaneous horse category was particularly at risk in comparison to race horses.
423 This higher incidence can arise from a poor control of cyathostomin populations or from a poor
424 diagnostic and management of the horse condition. Recent survey on a limited number of
425 premises in this region indicated that stud farms heavily relied on their veterinarians for the
426 design of parasite management scheme but that it was not the case in riding schools (Sallé et
427 al., 2017). This may contribute to increase cyathostomin prevalence outside stud farms. Non-
428 professional horse owners or their veterinarians or both may also have a reduced awareness
429 of cyathostomiasis management which contribute to increase the incidence of fatal cases.

430 In the case of *Parascaris* spp., our observations suggest that a few cases of suboptimal drug
431 efficacy occurred over the same time period. This is in line with other observations gathered
432 from the same region (Laugier et al., 2012), from other European countries (Boersema et al.,
433 2002; Schougaard and Nielsen, 2007; von Samson-Himmelstjerna et al., 2007; Näreaho et al.,
434 2011; Martin et al., 2018) or from more distant areas, like in Australia (Beasley et al., 2015) or
435 in the USA (Craig et al., 2007). This epidemiological context would hence suggest that the rise
436 of *Parascaris*-mediated death might be linked to a decrease in anthelmintic efficacy.

437 As a conclusion, this compilation of *postmortem* examination over a 29-year period in a unique
438 spatial entity, quantified major shifts in equine parasite communities that occurred within a 10-
439 year window from early 2000 onwards. Observed patterns suggested that the release of
440 macrocyclic lactones and praziquantel were major drivers of these shifts. The prevalence of
441 fatal parasite infection remained constant through time, but fatal cyathostomiasis cases have
442 been increasing since the year 2000. This likely mirrors both a confusion with other causes of

443 chronic diarrhoea by veterinarians in the field and a lack of awareness about drug resistance
444 in cyathostomin populations resulting in a poor control of these populations. Worryingly, the
445 rise of *Parascaris* spp. infection cases was concomitant with suboptimal anthelmintic efficacy
446 cases that have appeared within the last decade. While additional education efforts among
447 veterinarians and horse owners should contribute to dampen cyathostominosis cases, other
448 strategies should be leveraged for the control of *Parascaris* spp. in foals.

449

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590

591 **Legend to figures**

592 **Figure 1. Non-strongylid species distribution across season and horse breed**

593 The figure depicts the distribution of parasite burden measured in each breed type (MISC:
594 Miscellaneous; TB: Thoroughbred; FT: French Trotter) and across seasons.

595

596 **Figure 2. Parasites responsible for the death of young horses across breeds**

597 The relative contribution of equine parasites to the death of young horses (114 cases) is plotted
598 for each breed type considered (FT: French trotter; MISC: miscellaneous; TB: Thoroughbred).

599 The figure highlights the higher contribution of cyathostomiasis cases in the miscellaneous
600 breed type.

601

602 **Figure 3. Temporal variation of non-strongylid parasite prevalence**

603 The figure illustrates the breakpoints in parasite prevalence around the year 2000 for each of
604 the three non-strongylid parasites considered, *i.e.* 1998, 2007 and 2005 for *Gasterophilus* spp.,
605 *Parascaris* spp. and tapeworms respectively. Points are coloured according to the considered
606 time period, and the respective regression line is given with associated 95% confidence interval
607 (shaded area).

608

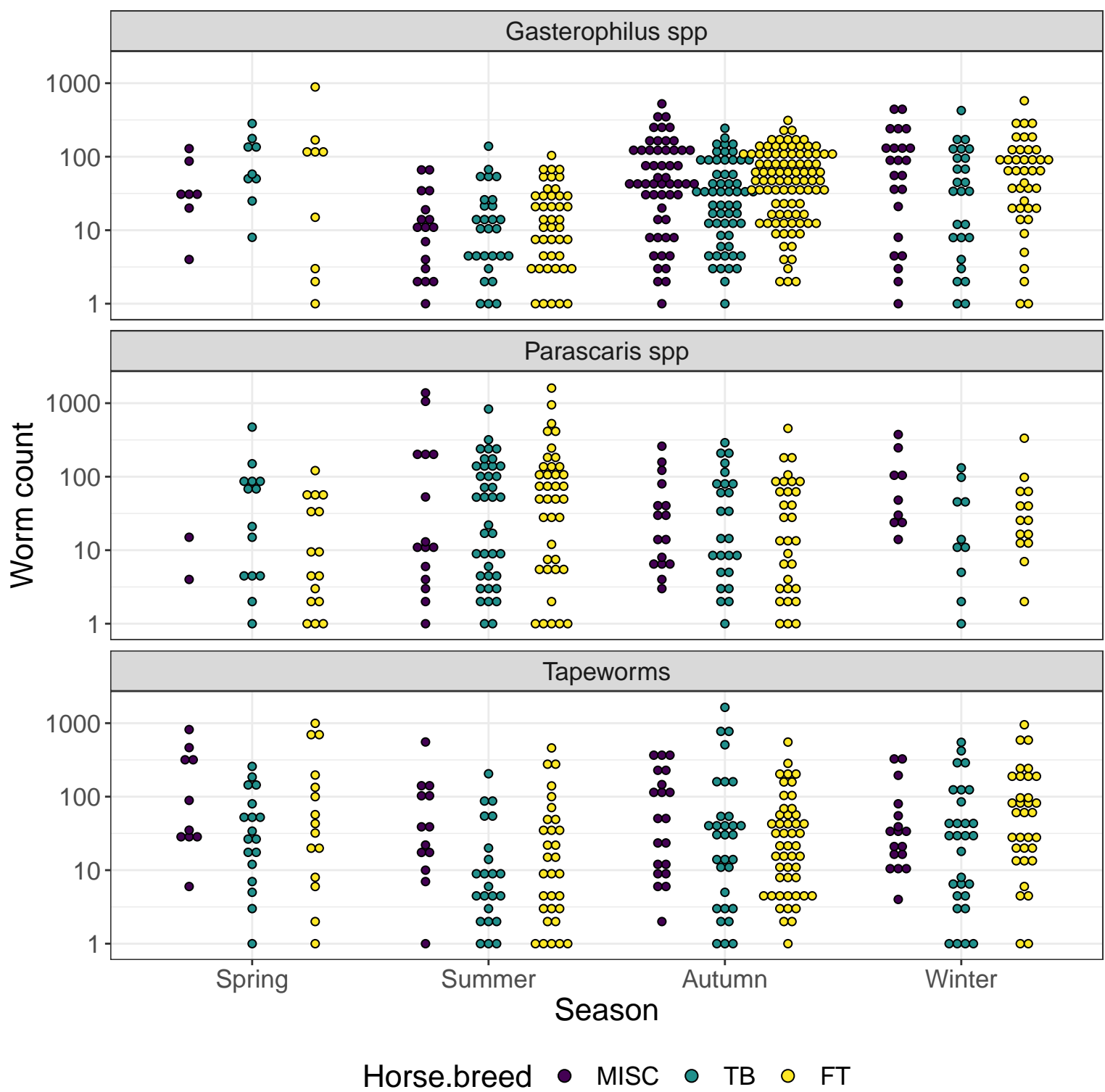
609 **Supplementary Figure 1. Quantile-Quantile plot of non-strongylid parasite burden data**

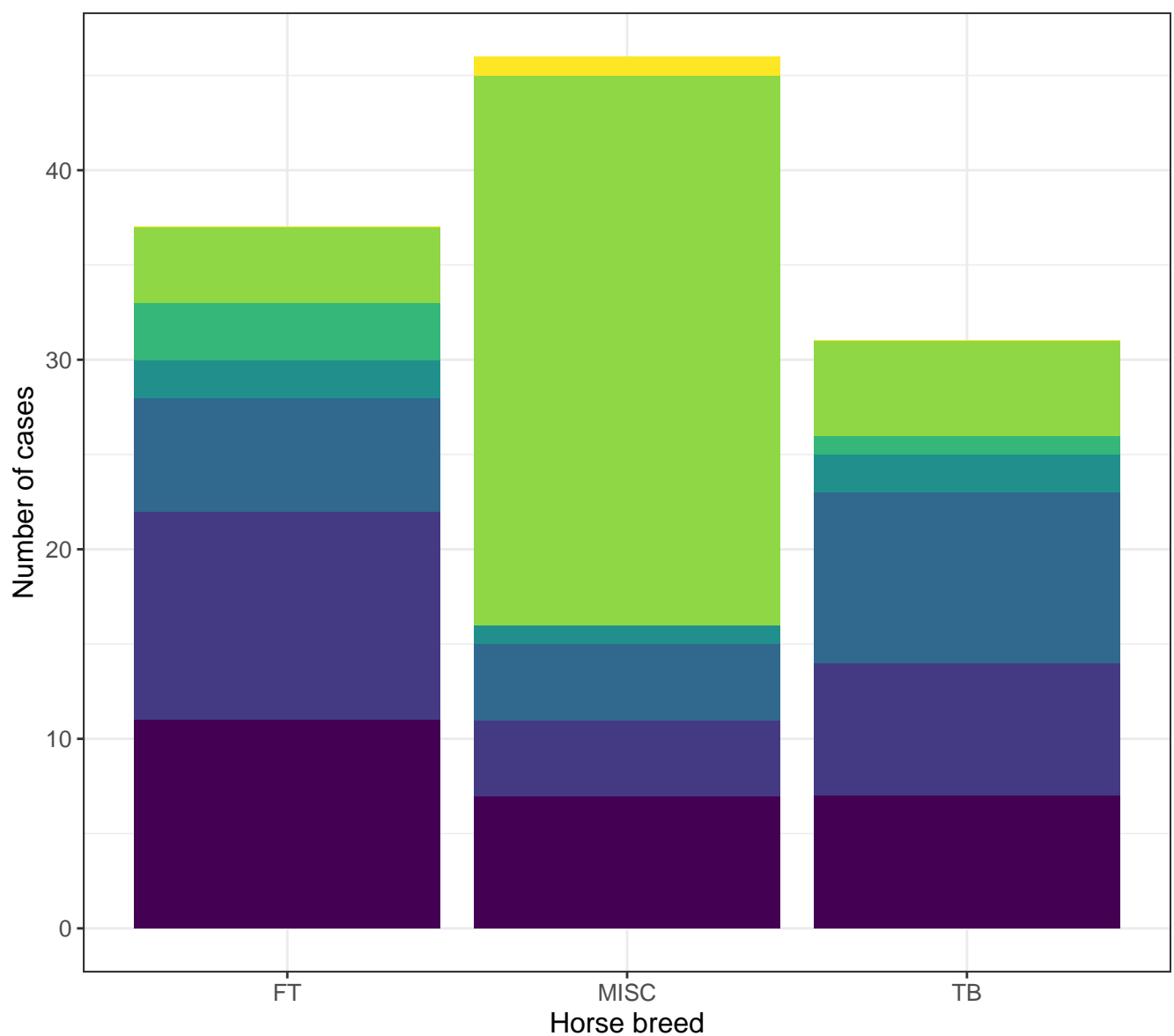
610 For every considered non-strongylid parasite (by row), empirical quantiles are plotted against
611 theoretical quantiles from the normal (left column), Poisson (middle column) or negative
612 binomial (right column) distributions. Any deviation from the black reference line is in favour of
613 a mismatch between theoretical expectations associated with the chosen distribution and the
614 observed data. For every parasite, data had their best fit against the negative binomial
615 distribution.

616

617 **Supplementary Figure 2. Distribution of cyathostomiasis cases throughout the study** 618 **period**

619 The number of fatal cyathostomiasis cases seen over the 29-year period is plotted for each
620 considered year.

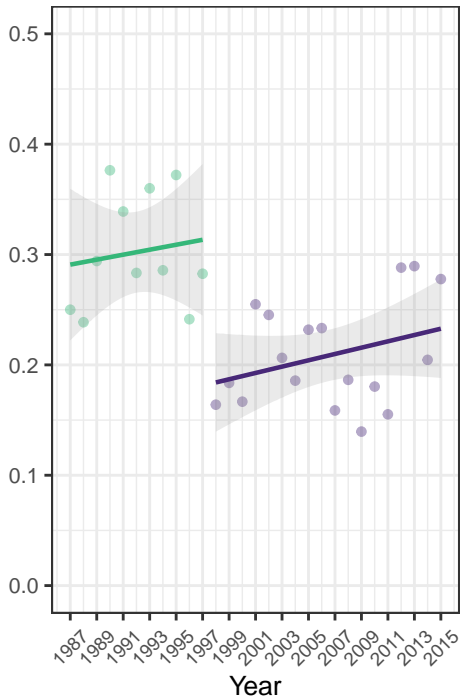




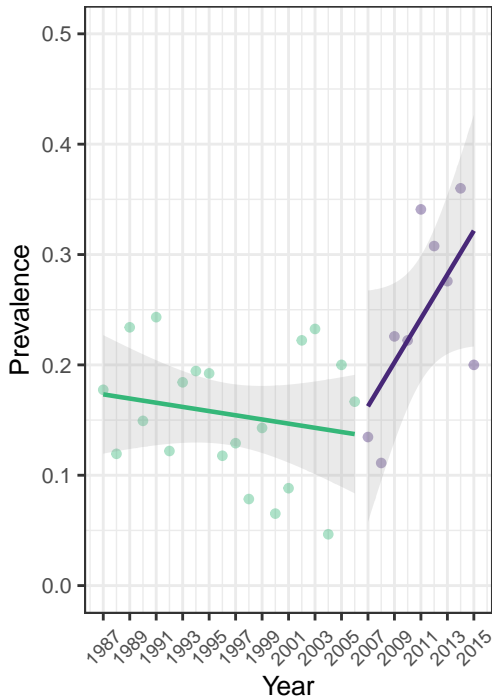
Cause

- Bots
- Cyathostomin
- Death upon deworming
- Other
- Parascaris spp*
- S.vulgaris*
- Tapeworms

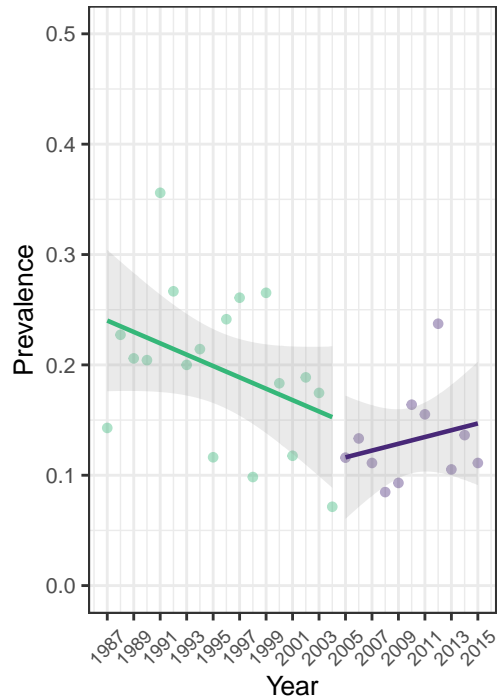
Gasterophilus spp.



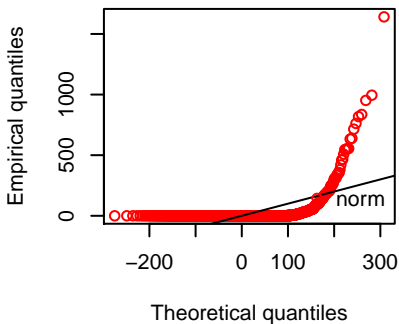
Parascaris spp.



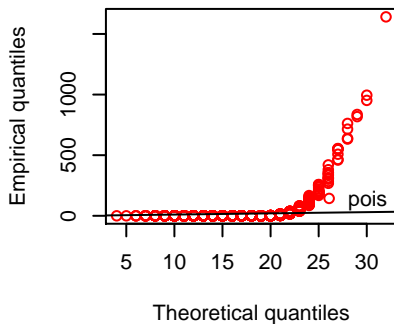
Tapeworms



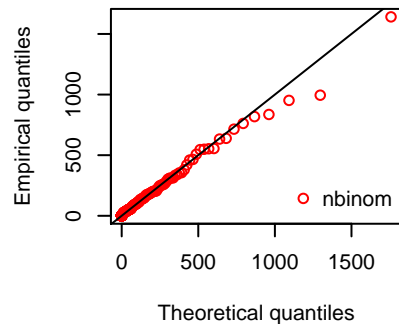
Tapeworms spp.
QQ-Plot – Normal



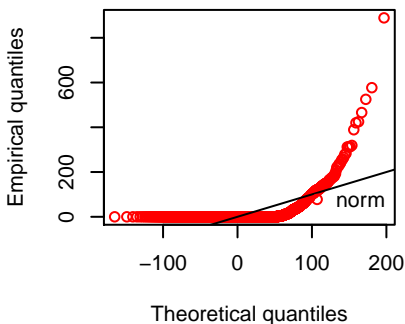
Tapeworms
QQ-Plot – Poisson



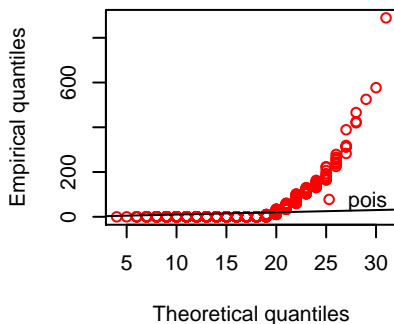
Tapeworms
QQ-Plot – Negative binomial



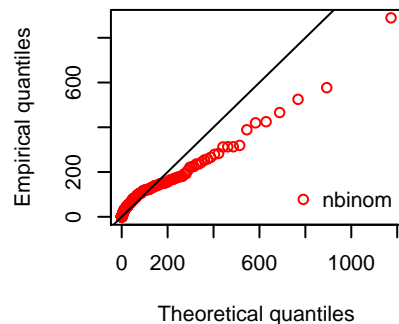
Gasterophilus spp.
QQ-Plot – Normal



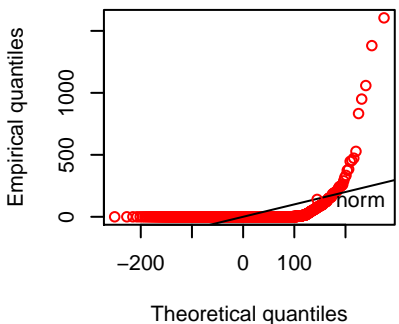
Gasterophilus spp.
QQ-Plot – Poisson



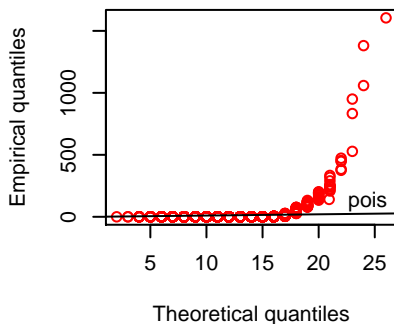
Gasterophilus spp.
QQ-Plot – Negative binomial



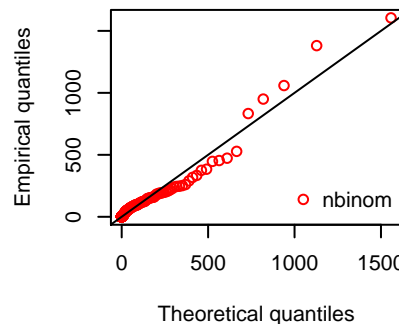
Parascaris spp.
QQ-Plot – Normal

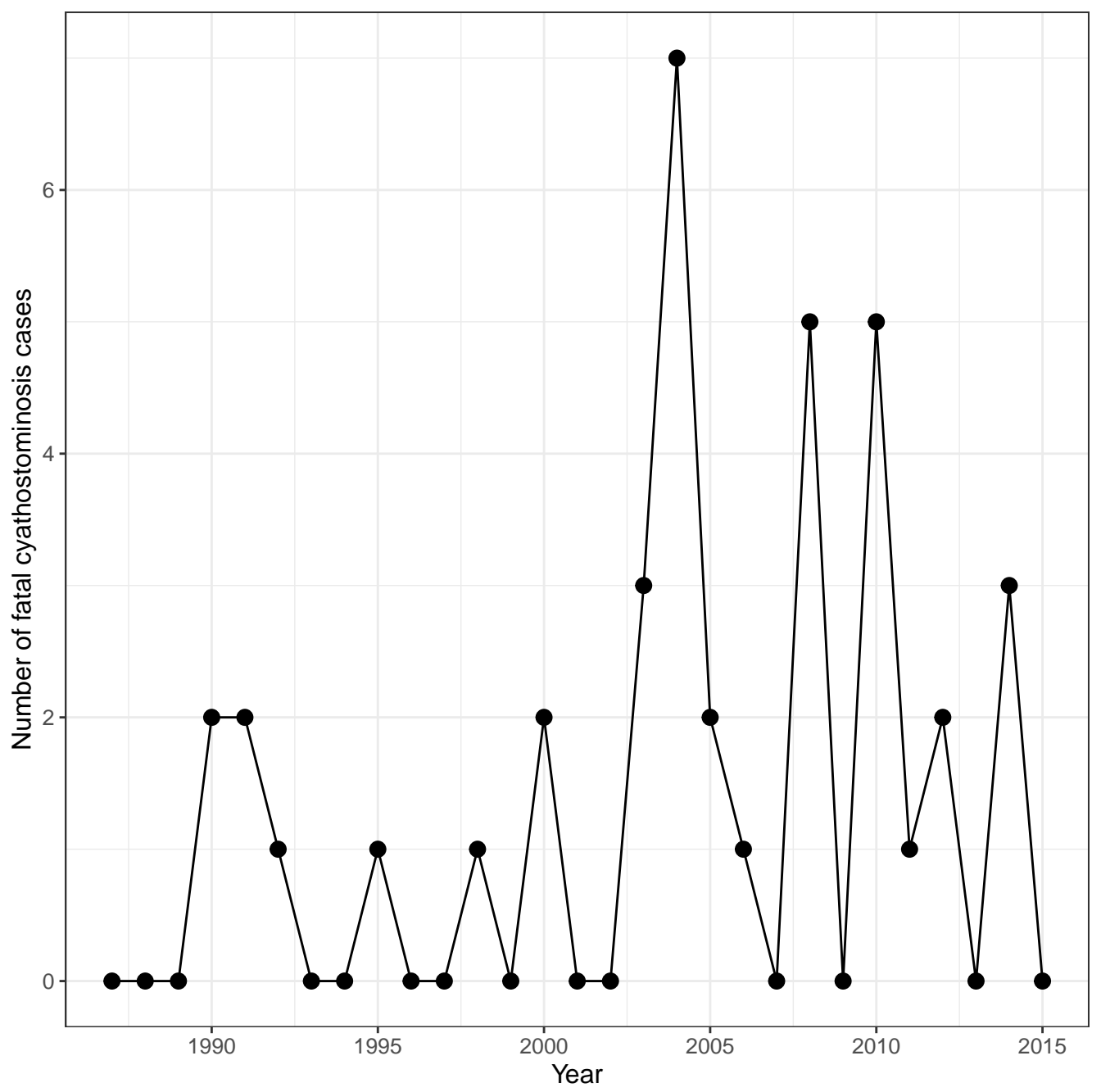


Parascaris spp.
QQ-Plot – Poisson



Parascaris spp.
QQ-Plot – Negative binomial





Species	Trait	Variable	Estimate	s.e.	Z-score	P-value
<i>Parascaris sp.</i>	Prevalence	Intercept	-2,53	0,21	-11,79	2,00E-16
		Before 2000 vs. After 2000	0,68	0,17	3,98	6,87E-05
		Spring vs. Summer	0,90	0,24	3,74	1,83E-04
		Spring vs. Autumn	1,27	0,25	5,08	3,86E-07
		Spring vs. Winter	0,75	0,30	2,52	1,17E-02
	Abundance	Intercept	1,10	0,33	3,28	1,02E-03
		Before 2000 vs. After 2000	0,80	0,38	2,11	3,48E-02
		Spring vs. Summer	2,01	0,41	4,86	1,15E-06
		Spring vs. Autumn	1,42	0,46	3,09	1,98E-03
		Spring vs. Winter	1,11	0,55	2,03	4,25E-02
Bots	Prevalence	Intercept	-1,99	0,26	-7,76	8,62E-15
		Miscellaneous vs. Thoroughbred	-0,78	0,18	-4,44	8,86E-06
		Miscellaneous vs. French Trotter	-0,30	0,17	-1,80	7,12E-02
		Before 2000 vs. After 2000	-0,53	0,13	-4,18	2,89E-05
		Spring vs. Summer	0,97	0,24	4,10	4,09E-05
		Spring vs. Autumn	2,66	0,23	11,52	2,00E-16
		Spring vs. Winter	2,17	0,25	8,73	2,00E-16
	Abundance	Intercept	2,68	0,35	7,73	1,04E-14
		Miscellaneous vs. Thoroughbred	-0,74	0,30	-2,48	1,30E-02
		Miscellaneous vs. French Trotter	-0,31	0,29	-1,07	2,86E-01
		Before 2000 vs. After 2000	-0,63	0,21	-3,05	2,32E-03
		Spring vs. Summer	-0,86	0,26	-3,28	1,06E-03
		Spring vs. Autumn	1,49	0,29	5,19	2,11E-07
		Spring vs. Winter	1,60	0,33	4,81	1,55E-06
Tapeworms	Prevalence	Intercept	-2,33	0,18	-13,04	2,00E-16
		> 1-year of age	0,91	0,14	6,68	2,40E-11
		Before 2000 vs. After 2000	-0,49	0,15	-3,25	1,16E-03
		Spring vs. Summer	0,10	0,21	0,49	6,22E-01
		Spring vs. Autumn	0,98	0,20	4,87	1,13E-06
		Spring vs. Winter	1,41	0,22	6,54	6,14E-11
	Abundance	Intercept	2,75	0,43	6,46	1,08E-10
		Miscellaneous vs. Thoroughbred	-1,23	0,39	-3,16	1,58E-03
		Miscellaneous vs. French Trotter	-0,67	0,38	-1,75	8,04E-02
		> 1-year of age	1,92	0,29	6,68	2,41E-11
		Spring vs. Summer	-1,78	0,34	-5,19	2,07E-07
		Spring vs. Autumn	0,41	0,37	1,09	2,74E-01
		Spring vs. Winter	1,23	0,43	2,82	4,78E-03