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1 Decision tree analysis of clinical data to aid diagnostic reasoning for equine laminitis: a cross-sectional

study.

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24 Abstract

25 The objective of this cross-sectional study was to compare the prevalence of selected clinical signs in 26 laminitis cases and non-laminitic but lame controls to evaluate their capability to discriminate laminitis from other causes of lameness. Participating veterinary practitioners completed a checklist of laminitis-associated 27 28 clinical signs identified by literature review. Cases were defined as horses/ponies with veterinary-diagnosed, clinically apparent laminitis; controls were horses/ponies with any lameness other than laminitis. 29 Associations were tested by logistic regression with adjusted odds ratios (OR) and 95% confidence intervals, 30 with veterinary practice as an *a priori* fixed effect. Multivariable analysis using graphical classification tree-31 32 based statistical models linked laminitis prevalence with specific combinations of clinical signs. Data were 33 collected for 588 cases and 201 controls. Five clinical signs had a difference in prevalence of greater than +50%: 'reluctance to walk' (OR 4.4, 'short, stilted gait at walk' (OR 9.4), 'difficulty turning' (OR 16.9), 34 'shifting weight' (OR 17.7) and 'increased digital pulse' (OR 13.2) (all P<0.001). 'Bilateral forelimb 35 36 lameness' was the best discriminator; 92% of animals with this clinical sign had laminitis (OR 40.5, 37 P < 0.001). If, in addition, horses/ponies had an 'increased digital pulse', 99% were identified as laminitis. 'Presence of a flat/convex sole' also significantly enhanced clinical diagnosis discrimination (OR 15.5, 38 P < 0.001). This is the first epidemiological laminitis study to use decision-tree analysis, providing the first 39 evidence-base for evaluating clinical signs to differentially diagnose laminitis from other causes of lameness. 40 41 Improved evaluation of the clinical signs displayed by laminitic animals examined by first-opinion 42 practitioners will lead to equine welfare improvements.

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44

46 Introduction

Equine laminitis is a painful disease of the foot that affects equidae worldwide (Mellor and others 2001;
Wylie and others 2011). The insidious nature of the disease and potential for unrelenting pain often
necessitates euthanasia of the affected animal on welfare grounds (Hunt 1993; Menzies-Gow and others
2010b). Effective diagnosis is necessary to allow prompt instigation of palliative and therapeutic treatments,
to maximise recovery prospects.

52 In equine medicine, 'laminitis' is used to describe animals presenting with pain localised to the lamellar region of the foot, with or without concurrent solar pain under the distal margin of the distal phalanx (Stashak 53 2002). There are no universally accepted gold-standard techniques for the detection and quantification of the 54 four stages of laminitis (Eustace 2010; Herthel and Hood 1999; Hunt and Wharton 2010; Menzies-Gow and 55 56 others 2010c; Swanson 1999). Acute laminitis arises with the development of clinical signs appreciable as changes in the normal stance and gait of the animal (Baxter 1994; Coffman and Garner 1972; Swanson 1999). 57 Acute laminitis either progresses to the subacute form or to the chronic form of the disease. The subacute 58 59 stage can either persist, develop to chronic laminitis, or lead to complete recovery. Development of chronic 60 laminitis usually results in a cycle of recurrent episodes (Hood 1999). The terminology used to describe chronic laminitis is extremely variable (Parks and Mair 2009), but is often taken to describe progression from 61 62 acute laminitis to failure of the SADP resulting in dislocation of the DP following detachment of the hoof wall (Grosenbaugh and others 1999). 63

Laminitis is necessarily commonly diagnosed solely on the presence of a combination of characteristic clinical signs (Baxter 1994; Vinuela-Fernandez *et al.* 2011a). Diagnostic challenges are compounded by the multifactorial aetiology of the disease, which can arise as a consequence of systemic inflammatory disease, endocrine disease or abnormal weight/load bearing which may initiate distinct pathophysiological processes as reviewed by Eades (2010). However, the common feature of all cases of laminitis is the induction of pathological changes within the SADP, resulting in overt foot pain and clinical signs related to lameness (Baxter 1994; Budras and others 2009a; Budras and others 2009b).

71 Despite the perceived importance there is remarkably little evidence-based data regarding the clinical presentation of laminitis (Eustace 2010; Hunt and Wharton 2010; Mellor and others 2001; Wylie and others 72 2013a), adding to inherent difficulties in establishing accurate diagnosis of laminitis due to the non-specific 73 74 nature of clinical signs and the absence of robust case definitions. Furthermore, there is no general agreement 75 regarding standardised criteria to diagnose laminitis or to classify affected animals based on the phase of disease progression and/or disease aetiology (Parks and Mair 2009; Rohrbach and others 1995). The 76 debilitating consequences of laminitis do, however, require prompt veterinary intervention and accurate 77 78 diagnosis is therefore essential.

All the factors outlined above complicate the overall challenge of diagnostic reasoning based on clinical signs, presenting the veterinary clinician with a challenge to diagnose laminitis differentially from other forms of orthopaedic disorder. Therefore, the aim of this study was to compare the prevalence of selected clinical signs in laminitis and non-laminitis lameness cases in order to evaluate the capabilities of clinical signs to differentially diagnose laminitis from other causes of lameness. The study is presented considering recommendations

85 of the Strengthening the Reporting of Observational studies in Epidemiology

- 86 (STROBE) statement (von Elm and others 2007).
- 87

88 Materials and Methods

- 89 Data were collected from two groups:
- 90 *Group A*

A convenience sample of five veterinary institutions (two referral centres, two large first-opinion and referral equine hospitals and a first-opinion mixed practice) were visited and invited to provide data for this study. In addition, veterinary practices (n=93) that were interested in participating in a parallel epidemiological investigation of equine laminitis, were contacted by telephone or email and invited to provide data on clinical signs of lameness (of any origin) for the study reported here. 96 A literature review was conducted to identify previously suggested clinical signs of laminitis and differential 97 diagnoses. The resultant list was reviewed by expert equine clinicians in selected referral hospitals and 98 laminitis researchers, and a 'lameness reporting form' (LM) (Supplementary Information Item 1) was 99 designed to gather information on laminitis-relevant clinical signs from both laminitic (cases) and non-100 laminitic lame (controls) horses.

Part one of the LM gathered case identifying information with five subsequent sections recording whether clinical signs pertaining to the foot, stance and lameness irregularities (clinical signs) were present, absent or had not been assessed. Part two of the LM allowed practitioners to record their diagnosis as free text and to select specific diagnostic techniques used to confirm the diagnosis from six tick-box options. A free-text comments section was also included for any additional information pertinent to confirmation of the diagnosis.

Participating practitioners were asked to complete a LM for equine lameness of any origin seen between February-April 2009, and January 2010-May 2011, with the second phase of data collection initiated to increase numbers for analysis. Completed forms were returned by post using supplied reply-paid envelopes. Upon arrival LMs were divided into two groups for analysis: one group containing reported laminitis cases and another containing all animals for which the primary cause of lameness was not laminitis (controls).

111 *Group B*

Following this development phase, a 'laminitis reporting form' (LRF) was finalised (Supplementary Information Item 2) as previously described (Wylie and others 2013a). As for the LM, the LRF consisted of five distinct sections on lameness, stance characteristics, feet affected and observed laminitis-related acute and chronic clinical signs. Based on the data collected from animals in Group A, some modifications to the form were made, hence for the purposes of this study only those clinical signs which were reported for both groups were compared. No further clinical data were recorded for the purposes of this study.

118

A LRF was completed for any case of laminitis, defined as a horse or pony with veterinary-diagnosed,
clinically apparent laminitis (i.e. an active episode of laminitis), attended by one of the participating
practitioners (Wylie and others 2013a). In animals with recurring laminitis, an episode of veterinary-

122	diagnosed active laminitis was defined as new if the animal had returned to its previous/normal level of
123	soundness and had not received analgesic medication for 14 days or more between episodes (Wylie and
124	others 2013a). However, for the purposes of this study only the first episode of laminitis was included.
125	Practices were asked to complete the LRF for all eligible cases occurring from May 2009 to April 2011.
126	Statistical analysis
127	To increase the numbers for data analysis, Groups A and B were combined. Multiple different clinical signs
128	were categorised (present, not present or not assessed) under the following five sections:
129	(1) <u>Lameness</u> : recumbency, refusal to move unless forced, reluctance to walk, lame at walk, lame at trot,
130	short stilted gait at walk, short stilted gait at trot, difficulty turning
131	(2) <u>Stance</u> : shifting weight, front feet placed in front of body, reluctance to lift foot
132	(3) <u>Feet affected</u> : bilateral front feet, bilateral hind feet or all four feet
133	(4) <u>Acute clinical signs</u> : increased digital pulse, increased hoof temperature, pain on sole pressure
134	(5) <u>Chronic clinical signs</u> : Coronary band swelling, coronary band depression, divergent growth rings,
135	change in hoof wall angle, wall separation, flat/convex sole, widened white line, pink crescent dorsal
136	to frog, sole prolapse
137	Initial examination, coding of data and descriptive analyses were conducted using Microsoft Excel (Excel
138	2003, Microsoft). The prevalence (including corresponding 95% confidence intervals [CI]) of each clinical
139	sign, excluding records where the sign was not assessed, in both case and control animals and the between-
140	group differences in prevalence of presence of clinical sign were determined. Associations between each
141	clinical sign and case or control status were tested using logistic regression models reporting adjusted odds
142	ratios (OR) taking into account veterinary practice as a fixed effect, with 95% confidence intervals (CI), and
143	Wald test P-values. All analyses were conducted in R Statistical Package (version 3.1.2 [©] 2014 The R

Foundation for Statistical Computing) using the 'epicalc' and 'tree' packages. Statistical significance was set at a value of P<0.05.

146 Multivariable analysis was carried out using a multi-factorial classification - tree-based statistical models (hereafter 'tree models') (Clark and Pregibon 1997). This analytical technique was chosen due to the 147 unbalanced dataset with potentially different combinations of factors present in different horses. The analysis 148 149 consisted of determining a binary division of the clinical signs prevalence data (laminitis vs. non-laminitis 150 lameness), such that there is the largest difference in terms of prevalence of laminitis vs. non-laminitis lameness for those two subsets of data. One subset of animals with a specific clinical sign is first considered 151 (e.g. those with 'bilateral forelimb lameness') and the binary division in terms of any of the other clinical 152 signs resulting in the largest difference in prevalence of laminitis is determined. The other subset is then 153 154 considered (e.g. those with no 'bilateral forelimb lameness') and again the clinical signs for which binary division gives the largest difference in prevalence of laminitis vs. non-laminitis lameness is determined. The 155 different "branches" of the tree are independent of each other in terms of what binary partitions are presented. 156 This binary partitioning is continued for smaller and smaller subsets of data until no differentiation in terms 157 of prevalence is possible. The trees are then 'pruned' to exclude very small differentiations based on a few 158 horses. The analysis is presented in graphical form allowing easy comprehension of the grouping of clinical 159 signs giving the largest differences in prevalence in the data. Univariable comparisons of the distribution of 160 clinical signs for particular subsets identified in the trees were then carried out as per the association between 161 162 clinical signs and case/controls status described above.

Five separate preliminary tree models were produced for the following characteristics to represent the features of clinically active laminitis recorded: i) lameness, ii) stance, iii) feet affected, iv) acute signs only and iv) acute and chronic signs. 'Lame at trot' and 'short stilted gait at trot' were excluded from the lameness tree model due to large numbers of missing data where these signs had not been assessed (missing for 55.0% and 49.4% of observations, respectively).

After consideration of the five preliminary trees, those variables identified in each preliminary tree as being the greatest differentiators in terms of laminitis were analysed together to form two combined tree models: (i) a combined model of lameness, stance characteristics, feet affected and observed laminitis-related acute clinical signs to reflect active episodes of laminitis in horses with no evidence of chronic laminitis, and (ii) a combined model of lameness, stance characteristics, feet affected and observed laminitis-related acute and 173 chronic clinical signs to reflect active episodes of laminitis in horses with evidence of previous SADP failure

174 (chronic laminitis).

175

176 **Results**

177 Recruitment

178 *Group A*

All five veterinary establishments visited agreed to provide data for this study. In addition, 25 first-opinion 179 veterinary practices agreed to participate, of which 14 (46.7%) contributed data to the study. Lameness forms 180 were provided for 238 unique horses/ponies: 89 (37.4%) from referral practices and 149 (62.6%) from first-181 182 opinion practices. Thirty-seven animals (15.5%) were diagnosed by veterinary practitioners as laminitis cases 183 and 201 (84.5%) were diagnosed with non-laminitis lameness. Other causes of lameness included, but were not restricted to, proximal suspensory desmitis (n=40, 17.3%), foot abscesses (n=22, 9.5%) and fractures 184 (n=16, 6.9%). Overall, 73 (30.7%: CI 24.8, 36.5) Group A animals were diagnosed on the basis of clinical 185 186 signs without further diagnostic procedures (cases 32.4%: CI 17.3, 47.5, controls 30.3%: CI 24.0, 36.7) and 155 (65.1%: CI 59.1, 71.2) animals were diagnosed using multiple diagnostic modalities (cases 62.2%: CI 187 188 46.5, 77.8, controls 65.7%: CI 59.1, 72.2). Stated diagnostic techniques used to investigate lameness in the laminitic cases included clinical examination (94.6%: CI 87.3, 100), radiography (64.9%: CI 49.5, 80.2), 189 190 regional anaesthesia (nerve blocks) (13.5%: CI 2.5, 24.5), surgical/post-mortem findings (13.5%: CI 2.5, 24.5) and blood testing for concurrent predisposing metabolic conditions (8.1%: CI 0.01, 16.9). 191

192 *Group B*

The recruitment of cases is described in detail in Wylie et al. (2013a). In brief, LRFs were received for 551
unique horses/ponies from 30 first-opinion veterinary practices over the two-year period.

195 Clinical signs

196 The prevalence of the presence of each clinical sign in laminitis cases and non-laminitis lame controls, 197 excluding records where the sign was not assessed, and difference in prevalence between the two groups are provided in Table 1. The overall prevalence of specific clinical signs ranged from 2.7% (CI 1.5, 3.9) for 'sole 198 199 prolapse' (number assessed = 706) to 85.0% (CI 81.4, 88.7) for 'lame at trot' (number assessed = 367). The 200 difference in prevalence between cases and controls ranged from -14.1% for 'lame at trot' (sign more common in controls) to +71.9% for 'short stilted gait at walk' (found more often in cases than controls). 201 There were five clinical signs with a difference in prevalence of greater than +50%: three lameness-related 202 signs ('reluctance to walk', 'short, stilted gait at walk' and 'difficulty turning'), one stance-related sign 203 ('shifting weight') and one acute clinical sign ('increased digital pulse'). 204

The logistic regression results are provided in Table 2. For each clinical sign there was a statistically significant increase in the odds of occurrence in the laminitis (cases) group, with the exception of 'recumbent', 'lame at trot' and 'coronary band swelling' for which there was no significant difference (P>0.05). No odds ratio could be calculated for 'coronary band depression' or 'sole prolapse' because no animals in the control group showed these clinical signs.

The preliminary tree models are provided in Supplementary Information Item 3. Consideration of the lameness tree identified the best discriminator as 'short stilted gait at walk'; 93.1% (CI 90.6, 95.5) of animals with that clinical sign had laminitis; 94.1% (CI 91.6, 96.5) of animals with both 'short stilted gait at walk' and 'difficulty turning' had laminitis. Of the 219 animals that did not have a 'short stilted gait at walk', only 27.9% (CI 21.9, 33.8) had laminitis – however, if they had 'difficulty turning' 59.7% (CI 48.0, 71.5) had laminitis. For animals where both these clinical signs were absent, if they were 'reluctant to walk' 40.0% (CI 15.2, 64.8) had laminitis.

The best discriminator in the stance tree was 'shifting weight'; 98.1% (CI 96.6, 99.6) of animals with that clinical sign had laminitis. In animals that were not 'shifting weight', 'front feet placed in front of the body' identified 94.2% (CI 89.2, 99.1) as laminitis cases. In the 'acute clinical signs' tree, 91.0% (CI 88.5, 93.5) of animals with 'increased digital pulses' had
laminitis, and 'pain on sole pressure' in the absence of 'increased digital pulses' identified 69.0% (CI 52.1,
85.8) as cases of laminitis.

The best discriminator in the 'acute and chronic clinical signs' tree was 'increased digital pulses'; 91.0% (CI 88.4, 93.5) of animals with that clinical sign had laminitis, and the additional presence of 'divergent growth rings' identified 100% as laminitis cases.

226 The tree diagram combining categories of clinical signs for acute laminitis with lameness, stance and feet is provided in Figure 1. Presence of 'lameness in both forelimbs' was the best discriminator, with 93.1% (CI 227 90.7, 95.5) of animals with this clinical sign belonging to the laminitis group. Additional presence of an 228 229 'increased digital pulse' improved diagnostic accuracy to 99% (CI 97.9, 100) (P<0.001). A 'bilateral forelimb lameness' with no 'increase in digital pulse', yet presence of a 'short stilted gait at walk' identified 230 231 100% of animals as laminitis cases, however statistical analysis of this sub-group and the presence of 'shifting weight' was not possible due to small numbers of animals with these signs. The presence of 'pain 232 233 on sole pressure' was not statistically associated with improved clinical discrimination (P=0.30).

The overall tree diagram considering both acute and chronic laminitis clinical signs with lameness, stance and feet is provided in Figure 2. Presence of 'lameness in both forelimbs' was again the best discriminator; 92% of animals with this clinical sign had laminitis (P<0.001). The additional presence of 'increased digital pulses' improved this to 99% of cases (P<0.001). Presence of a 'flat/convex sole' also provided improved clinical discrimination (P=0.002). It was not possible to assess statistical significance for 'short stilted gait at walk', or 'shifting weight', again because of the small numbers of animals with these signs.

240

241 Discussion

This is the first study comparing the prevalence of veterinary-recognised clinical signs in laminitis and othercauses of lameness to evaluate the capabilities of discrimination for differential diagnosis.

244 A wide range of clinical signs were displayed by the laminitic cases, in agreement with previous reviews 245 (Baxter 1994; Eustace 2010; Hunt and Wharton 2010; Swanson 1999). There were no individual, or combinations of, clinical signs present in every case. The clinical signs that were considered to be the most 246 247 useful on the basis of this work were three features of lameness investigation ('reluctance to walk', 'short, 248 stilted gait at walk' and 'difficulty turning'), one feature of stance ('shifting weight') and an 'increased digital pulse'. All these signs had a difference in prevalence of over 50% between active laminitis cases (signs more 249 prevalent) and non-laminitic lame horses (signs less prevalent). As the clinical details forms were designed to 250 gather information on laminitis, it may be expected there was a statistically significant difference in the 251 252 distribution of many of the clinical signs between laminitis cases and non-laminitis lameness controls. For the purposes of this study it was considered important to focus only on the lameness-associated clinical signs 253 for two main reasons. Firstly, because regardless of the underlying pathological process of laminitis, the 254 255 common feature of all cases of laminitis is the induction of pathological changes within the SADP, resulting 256 in overt foot pain and clinical signs related to lameness (Baxter 1994; Budras and others 2009a; Budras and others 2009b; Eades 2010), and as a consequence previous epidemiological studies of laminitis have used 257 258 only lameness-associated clinical signs as their case inclusion/exclusion criteria (Alford and others 2001; 259 Dorn and others 1975; Hood and others 1994; Menzies-Gow and others 2010a; Parsons and others 2007; 260 Slater and others 1995). Secondly, to keep the amount of work required by the veterinary surgeons to a minimum to enhance compliance. Collection of data regarding systemic clinical signs would have increased 261 the amount of work required by the participating veterinary practitioners, and it was considered that their 262 presence would aid the diagnosis of the underlying, predisposing condition rather than laminitis directly. 263 264 Nevertheless, it is acknowledged that as part of the diagnostic process veterinarians will use the animal's 265 history and other clinical features in making their diagnosis. As such, collection of additional clinical data in 266 future studies would be useful to improve the current decision trees, as well as to generate further trees 267 pertaining to, for example, signs of colic.

Currently, visual assessment of lameness is a highly subjective process. Many kinetic and kinematic methods for objectively assessing lameness have been reviewed previously (Hood and others 2001; Keegan 2010), and it is possible that these may prove to be more reliable than visual assessment alone in the future (Dyson 271 2011). Further evaluation of techniques to evaluate stance and gait characteristics of lame animals may result 272 in a more objective method of diagnosing and/or scoring laminitis, as well as other reasons for lameness. Recently developed techniques allow assessment of horse movement without impeding the use of the animal, 273 274 and may have a role in evidence-based assessment of lameness in horses in veterinary practice in the future 275 (Dyson 2011; Keegan 2010; Pfau and others 2007). There was no statistically significant difference in prevalence of 'lameness at trot' between cases and controls, and this variable was not included in the tree 276 analysis due to large number of laminitic cases that were not assessed at trot. The high level of missing data 277 is likely to reflect the appropriate reluctance of veterinary surgeons to trot suspect laminitis cases on welfare 278 279 grounds and so as not to exacerbate lamellar pathology, and the common use of intrasynovial anaesthesia for 280 diagnosis of other lamenesses commonly evaluated at the trot.

281 Two clinical signs – 'coronary band depression' and 'prolapsed sole' - were pathognomonic for laminitis in 282 this study, . were only found in 13.6% and 3.7% of cases, respectively. Both these signs can indicate disease progression to chronic phase laminitis (i.e. SADP failure and distal phalanx dislocation within the hoof); 283 284 therefore these signs would not be expected to be present in acute cases, unless they were also suffering from 285 concurrent pathology such as chronic seedy toe/white line disease or severe club feet (Kuwano and others 286 1999). These results may help veterinary practitioners prioritise where to begin their clinical examination of 287 an active laminitis case, as primary inspection of the sole and coronary band would prevent the animal undergoing lameness evaluation which could precipitate further SADP damage/failure. 288

Two overall combined trees were generated to reflect the two clinical scenarios of active laminitis, one consisting of clinical signs considered to occur in the acute phase of the disease, and one that also contained data reflective of lamellar damage and displacement of the SADP. In both scenarios, the presence of a bilateral lameness was the most useful discriminator, followed by the presence of increased digital pulses. Whilst these clinical presentations are not specific for laminitis, this work provides an evidence-base for case diagnosis and future epidemiological case definitions.

This work did not provide evidence for some commonly cited clinical signs of diagnostic importance. In particular, 'front feet in front of the body', taken to represent the classic 'laminitis stance', was found in less than half of the diagnosed active laminitis cases, and did not prove to be a useful discriminator. Therefore, despite much anecdotal publicity of this visibly apparent clinical sign (Stashak 2002; Swanson 1999),
veterinarians, researchers and owners should be careful to avoid relying on its presence for making a
diagnosis of laminitis [40].

The use of clinical recording forms based on evidence-based recommendations may help veterinary practitioners structure their clinical examination of an active laminitis case. However, in medical practice well-validated diagnostic algorithms tools are underused (Pearson and others 1994). For example, a simple predictor based on seven clinical signs for ischaemia in humans was only used in 2.8% of cases (Corey and Merenstein 1987). The clinical usefulness of developing such a technique would need to be established by a survey of first-opinion practitioners to decide whether such a tool would provide useful assistance in laminitis diagnosis in the field.

308 The limitations of this study include diagnosis by a number of different veterinary clinicians, which may have 309 different levels of experience. To take this into account veterinary practice was included in the generation of the odds ratio estimates, however, misclassification bias may still occur, although this would have tended to 310 311 shift the odds ratios towards non-significant. Similarly, as it is not possible to obtain a definitive diagnosis of active laminitis in an observational epidemiological study there was the potential for misclassification of 312 cases and controls. For this reason, veterinary recordings of the clinical signs observed was used, as 313 described in Wylie et al., (Wylie and others 2013a, b) and misclassification would have again reduced the 314 315 ability to detect significant differences rather than produce anomalous significant differences. Inclusion of 316 data in the tree models required the animals to have data for each included variable, resulting in smaller numbers of contributing individuals as the trees became more complex. Consequently, although the variables 317 retained high statistical significance, smaller contributing sample sizes led to larger confidence intervals 318 319 around prevalence point estimates and the need therefore for some caution in their interpretation.

It is acknowledged that there may be some bias in the data if veterinary practitioners did not accurately detail the clinical signs which they observed and perhaps listed clinical signs that they anticipated to reflect their diagnosis. Furthermore, it would be interesting to collect greater numbers of control animals to conduct the analyses between specific control lamenesses, such as forelimb foot pain only, to highlight more subtle differences between presenting pathologies. 325 In conclusion, separate clinical signs were compared between laminitis and non-laminitis cases of lameness, 326 and no individual sign was present in every case of laminitis. The clinical signs which best indicated a case of laminitis were characteristic of the chronic phase of the disease only. Improved evaluation of the clinical 327 signs displayed by laminitic animals examined by first-opinion practitioners will lead to equine welfare 328 329 improvements, as the best recoveries occur in animals undergoing intensive treatment within several hours of 330 the appearance of the disease (Redden 1986). Future consensus on a basic disease definition may permit future systematic review and meta-analysis of epidemiological investigations collecting similar information in 331 different locations worldwide. 332

333

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426 Table 1: Prevalence and 95% confidence intervals (CI) for each clinical sign in both laminitis cases and non-laminitis lameness controls, excluding records where the

427 sign was not assessed, and the percentage of horses that were assessed with corresponding difference in prevalence.

Clinical signs		Cases (n=		Controls	(n=201)		Overall (n=789)							
		Present (n)	Absent (n)	Prevalence (%)	LCI (%)	UCI (%)	Present (n)	Absent (n)	Prevalence (%)	LCI (%)	UCI (%)	Number assessed	Percentage assessed (%)	Difference in prevalence (%)
Lameness	Recumbent	24	479	4.8	2.9	6.6	1	191	0.5	0.0	1.5	695	88.1	+4.3
	Refusal to move unless forced	148	361	29.1	25.1	33.0	14	180	7.2	3.6	10.9	703	89.1	+21.9
	Reluctance walk	395	155	71.8	68.1	75.6	38	157	19.5	13.9	25.1	745	94.4	+52.3
	Lame walk	409	95	81.2	77.7	84.6	76	122	38.4	31.6	45.2	702	89.0	+42.8
	Lame trot	152	42	78.4	72.6	84.2	160	13	92.5	88.6	96.4	367	46.5	-14.1
	Short stilted walk	446	66	87.1	84.2	90.0	29	162	15.2	10.1	20.3	703	89.1	+71.9
	Short stilted trot	125	55	69.4	62.7	76.2	53	119	30.8	23.9	37.7	352	44.6	+38.6
	Difficulty turning	456	47	90.7	88.1	93.2	52	137	27.5	21.2	33.9	692	87.7	+63.1
Stance	Shifting weight	316	256	55.2	51.2	59.3	7	188	3.6	1.0	6.2	767	97.2	+51.7
	Front feet in front	250	317	44.1	40.0	48.2	6	190	3.1	0.7	5.5	763	96.7	+41.0
	Reluctance lift foot	300	269	52.7	48.6	56.8	24	169	12.4	7.8	17.1	762	96.6	+40.3
Feet Affected	Bilateral fore	538	44	92.4	90.3	94.6	32	152	17.4	11.9	22.9	766	97.1	+71.7
	Bilateral hind All four feet	244 234	323 348	43.0 40.2	39.0 36.2	47.1 44.2	25 5	156 193	13.8 2.5	8.8 0.3	18.8 4.7	748 780	94.8 98.9	+28.3 +39.5
Acute	Increased digital pulse	520	50	91.2	88.9	93.6	45	150	23.1	17.2	29.0	765	97.0	+68.2
	Increased hoof temperature	324	218	59.8	55.7	63.9	30	164	15.5	10.4	20.6	736	93.3	+44.3
	Pain sole pressure	263	271	49.3	45.0	53.5	35	149	19.0	13.4	24.7	718	91.0	+30.2
Chronic	Coronary band swelling	27	505	5.1	3.2	6.9	6	186	3.1	0.7	5.6	724	91.8	+2.0
	Coronary band depression	73	462	13.6	10.7	16.6	0	192	0.0	0.0	0.0	727	92.1	+13.6
	Divergent growth rings	148	378	28.1	24.3	32.0	3	190	1.6	0.0	3.3	719	91.1	+26.6

Change hoof wall angle	129	383	25.2	21.4	29.0	7	186	3.6	1.0	6.3	705	89.4	+21.6
Wall separation	71	445	13.8	10.8	16.7	2	184	1.1	0.0	2.6	702	89.0	+12.7
Flat/convex sole	232	291	44.4	40.1	48.6	9	180	4.8	1.7	7.8	712	90.2	+39.6
Widened white line	133	368	26.6	22.7	30.4	8	176	4.4	1.4	7.3	685	86.8	+22.2
Pink crescent	46	464	9.0	6.5	11.5	1	189	0.5	0.0	1.6	700	88.7	+8.5
Sole prolapse	19	498	3.7	2.1	5.3	0	189	0.0	0.0	0.0	706	89.5	+3.7

429 Table 2: Odds ratios and 95% confidence intervals (CI), with corresponding Wald *P*-values,

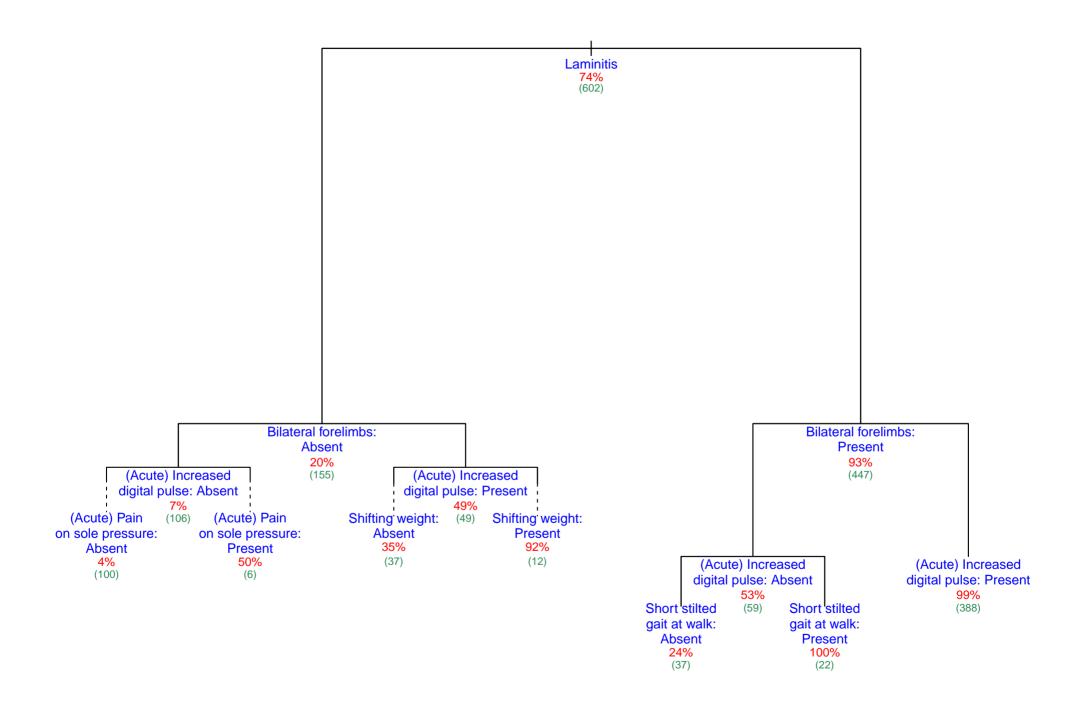
430 for each clinical sign in laminitis cases compared to non-laminitis lameness controls. ORs are

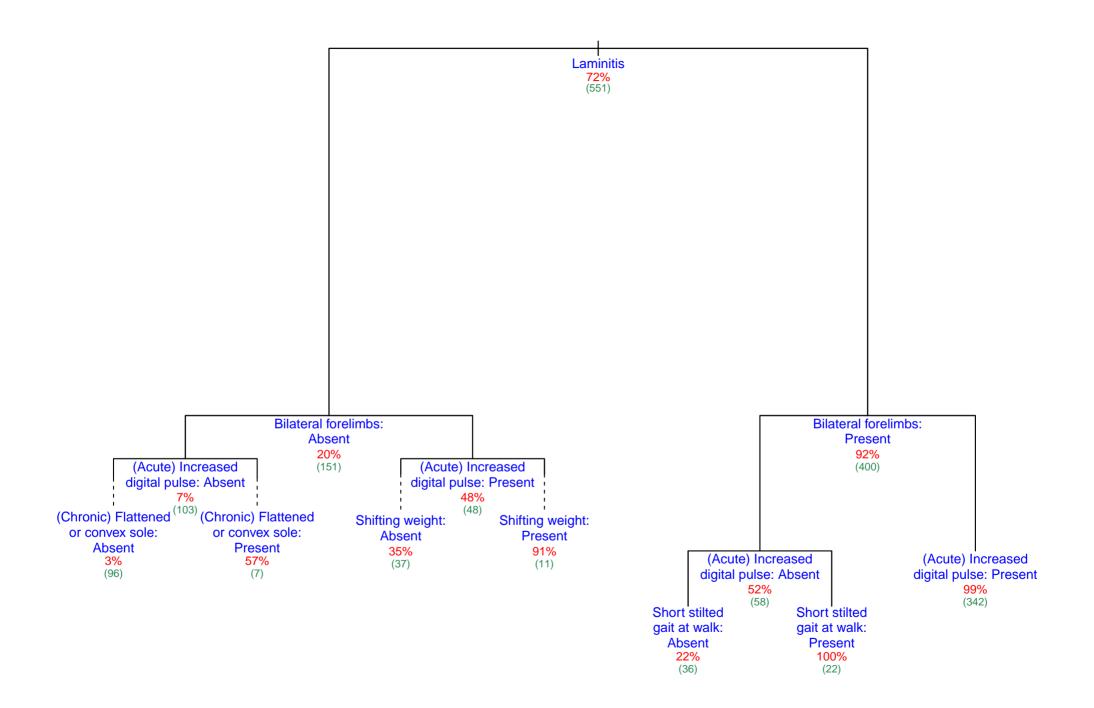
431 adjusted for the effect of veterinary practice.

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Clinical Signs		Number	Adjusted Odds Ratio	95% Confidence Interval	Wald P-value
Lameness	Recumbent	695	5.1	0.5, 51.4	0.17
	Refusal to move unless forced	703	3.5	1.6, 7.7	0.002
	Reluctance walk	745	4.4	2.2, 8.6	<0.001
	Lame walk	702	2.2	1.0, 4.7	0.04
	Lame trot	367	0.3	0.0, 2.6	0.29
	Short stilted walk	703	9.4	4.5, 19.6	<0.001
	Short stilted trot	352	3.9	1.6, 9.6	0.003
	Difficulty turning	692	16.9	7.0, 40.8	<0.001
Stance	Shifting weight	767	17.7	6.8, 45.6	<0.001
	Front feet in front	763	24.5	7.9, 75.9	<0.001
	Reluctance lift foot	762	4.0	1.9, 8.1	<0.001
Feet Affected	Bilateral fore	766	40.5	16.3, 100.9	<0.001
	Bilateral hind	748	21.3	7.7, 59.1	<0.001
	All four feet	780	96.3	22.1, 419.8	<0.001
Acute	Increased digital pulse	765	13.2	6.0, 29.3	<0.001
	Increased hoof temperature	736	5.7	2.8, 11.5	<0.001
	Pain sole pressure	718	2.7	1.4, 5.3	0.005
Chronic	Coronary band swelling	727	1.1	0.3, 3.9	0.88
	Coronary band depression	724	NA	NA	NA
	Divergent growth rings	719	96.3	17.1, 542.8	<0.001
	Change hoof wall angle	705	21.1	6.3, 71.0	<0.001
	Wall separation	702	58.5	5.1, 672.8	<0.001
	Flat/convex sole	712	15.5	5.9, 40.5	<0.001
	Widened white line	685	17.3	5.5, 54.5	<0.001
	Pink crescent	700	16.5	2.0, 136.5	0.009
	Sole prolapse	706	NA	NA	NA

433	Figure 1: Tree diagram of the occurrence of laminitis for combinations of lameness, stance,
434	feet affected, and acute laminitis clinical signs. Data were from 586 horses/ponies for which
435	information on each clinical sign was described, of which 74% had laminitis. The percentage
436	at the end of each branch are the occurrence rates of laminitis in those horses/ponies with that
437	particular combination of clinical signs, and the value in brackets the number of
438	horses/ponies of that particular combination of clinical signs.
439	Figure 2: Overall tree diagram of the occurrence of laminitis for combinations of lameness,
440	stance, feet affected, acute and chronic laminitis clinical signs. Data were from 551
441	horses/ponies for which information on each clinical sign was described, of which 72% had
442	laminitis. The percentage at the end of each branch are the occurrence rates of laminitis in
443	those horses/ponies with that particular combination of clinical signs, and the value in
444	brackets the number of horses/ponies of that particular combination of clinical signs.
445	
445 446	
446	
446 447	Supplementary Information Item 1: Lameness reporting form (LM) used to investigate the
446 447 448	Supplementary Information Item 1: Lameness reporting form (LM) used to investigate the clinical signs of laminitis in Group A recruiting both cases and controls.
446 447 448 449	
446 447 448 449 450	clinical signs of laminitis in Group A recruiting both cases and controls.
446 447 448 449 450 451	clinical signs of laminitis in Group A recruiting both cases and controls. Supplementary Information Item 2: Laminitis reporting form (LRF) used to investigate the





- Supplementary Information Item 1: Lameness reporting form (LM) used to investigate the clinical signs of laminitis in Group A recruiting both cases and controls.

	L	MENE	ESS R	EPO	RTIN	IG FO	RM	1		AHT
Name of ho	orse/pony: _									격변형
Surname of	f owner/Case	e I.D:								Animal Health Trust
Date of clin	ical examina	tion:			/			/		
Lameness					Δεσ	eccm	ent	(nle:	250 (circle 1 option per line)
Recumbent					AS	Yes		No		Didn't assess
	nove unless	forced	1			Yes		No		Didn't assess
Reluctance		Torcea				Yes		No		Didn't assess
Lame at wa						Yes		No		Didn't assess
Lame at tro						Yes		No	-	Didn't assess
	ed gait at wa	lk				Yes		No		Didn't assess
	ed gait at tro					Yes		No		Didn't assess
Difficulty tu		-				Yes		No		Didn't assess
							,		'	
Stance					Ace	sessm	ent	(ple:	ase o	circle 1 option per line)
Shifting we	iaht		_	_	-	Yes		No		Didn't assess
	placed in from	nt of b	odv		<u> </u>	Yes				Didn't assess
	laced under					Yes		No		Didn't assess
Square star						Yes		No		Didn't assess
	for a foot to	be lift	ed			Yes				Didn't assess
										bran e abbobb
Feet	Assessme	nt			Mo	st sev	ore	lv at	ffor	cted foot/feet
affected	(please circle		n per li	ne)		ase tick				
Right fore		/ Did	-						1.11	
Left fore	Yes / No		n't as							
Right hind		1	n't as							
Left hind	Yes / No		n't as							
Lore mild	105 / 110	/ 214	in e as	0000						
Clinical sig	gns of the <u>r</u> oot/feet	<u>nost</u> s	ever	ely	Ass	sessm	ent	(plea	ase o	circle 1 option per line)
	ligital pulse					Yes	1	No	1	Didn't assess
	noof tempera	ture				Yes		No		Didn't assess
	hoof temper					Yes	1	No	1	Didn't assess
Pain on sole						Yes	1	No	1	Didn't assess
	and swelling					Yes	1	No	1	Didn't assess
	and depress					Yes	1	No	1	Didn't assess
Clinical sig	gns of the <u>r</u> pot/feet	<u>nost</u> s	ever	ely	Ass	sessm	ent	(plea	ase o	circle 1 option per line)
Divergent o	rowth rings	(wider	r at he	eels)		Yes	1	No	1	Didn't assess
	dorsal hoof v					Yes		No	1	Didn't assess
Wall separa			-		1	Yes		No	1	Didn't assess
	r convex sol	e				Yes	- <u>-</u>	No	1	Didn't assess
Widened w						Yes		No	1	Didn't assess
	nt dorsal to	frog			1	Yes		No	1	Didn't assess
Prolapsed s						Yes		No	1	Didn't assess
fice use only:							<i>.</i>			
the use only:		An	imal H	lealth	Trus	1361				
AHT	Reference:							7		

- 7 Supplementary Information Item 2: Laminitis reporting form (LRF) used to investigate the
- 8 clinical signs of laminitis in Group B (cases only).

344	ANIMAL HI LAMINITIS REI	EALTH TRUST PORTING FORM	1 wa	Ind Horse Welfare
1. Name of horse/por	ıy:			
2. Surname of owner,	/Case ID:			
3. Date of clinical exa	mination: /		y	
4. Lameness		Assessment (<i>please cross o</i> NO	ne option per line) DIDN'T ASSESS
Recumbent				
Refusal to move u	nless forced			
Reluctance to wall	k			
Lame at walk				
Lame at trot				
Short, stilted gait	at walk			
Short, stilted gait	at trot			
Difficulty turning				
5. Stance		Assessment (YES	please cross o NO	<i>ne option per line)</i> DIDN'T ASSESS
Shifting weight				
Leg trembling				
Front feet placed i	in front of body			
Hind feet placed u				
Reluctance for a for				
5 Feet affected At	ffected (nlease cross one	ontion per line)		rely affected foot
Right fore	ffected (please cross one ES NO	option per line) DIDN'T ASSESS	(or feet if	rely affected foot bilaterally affected) ss all that apply)
Right fore [Left fore [(or feet if	bilaterally affected)
Right fore [Left fore [Right hind [(or feet if	bilaterally affected)
Right fore [Left fore [(or feet if	bilaterally affected)
Right fore [Left fore [Right hind [Left hind [ES NO		(or feet if <i>(please cro</i>	bilaterally affected)
Right fore [Left fore [Right hind [Left hind [Clinical signs of the affected foot/feet	e most severely	DIDN'T ASSESS	(or feet if) (please cro [[[[]]]]]]]]]]]]]]]	bilaterally affected) sss all that apply)
Right fore [Left fore [Right hind [Left hind [7. Clinical signs of th affected foot/feet [Increased digital p	e most severely	DIDN'T ASSESS	(or feet if) (please cro [[[[]]]]]]]]]]]]]]]	bilaterally affected) sss all that apply)
Right fore [Left fore [Right hind [Left hind [7. Clinical signs of th affected foot/feet [Increased digital p [ES NO	DIDN'T ASSESS	(or feet if) (please cro [[[[]]]]]]]]]]]]]]]	bilaterally affected) sss all that apply)
Right fore [Left fore [Right hind [Left hind [7. Clinical signs of th affected foot/feet [Increased digital p	ES NO	DIDN'T ASSESS	(or feet if) (please cro [[[[]]]]]]]]]]]]]]]	bilaterally affected) sss all that apply)
Right fore [Left fore [Right hind [Left hind [7. Clinical signs of th affected foot/feet [Increased digital p [Increased hoof ter [Focal sole pain in [ES NO	DIDN'T ASSESS	(or feet if) (please cro [[[[]]]]]]]]]]]]]]]	bilaterally affected) sss all that apply)
Right fore [Left fore [Right hind [Left hind [7. Clinical signs of th affected foot/feet [Increased digital p [Increased hoof ter [Focal sole pain in [ES NO	DIDN'T ASSESS	(or feet if) (please cro [[[[]]]]]]]]]]]]]]]	bilaterally affected) sss all that apply)
Right fore [Left fore [Right hind [Left hind [Clinical signs of the affected foot/feet [Increased digital p [Increased hoof ter [Focal sole pain in [Generalised dorsa [Coronary band sw [ES NO	DIDN'T ASSESS	(or feet if) (please cro please cross o NO	bilaterally affected) sss all that apply)
Right fore [Left fore [Right hind [Right hind [Left hind [7. Clinical signs of th affected foot/feet [Increased digital p [Focal sole pain in [Generalised dorsa [Coronary band sw [Coronary band de [8. Clinical signs of th affected foot/feet [ES NO	DIDN'T ASSESS	(or feet if) (please cro please cross o NO	bilaterally affected)
Right fore [Left fore [Right hind [Right hind [Right hind [Output [Right hind [Increased doot/feet [Increased hoot feet [Focal sole pain in [Generalised dorsa [Coronary band sw [Coronary band de [S. Clinical signs of th [affected foot/feet [Divergent growth	ES NO Set NO Set Severely Nouse Set Severely Nouse Set Severely S	DIDN'T ASSESS	(or feet if) (please cro please cross o NO	bilaterally affected)
Yi Right fore [Left fore [Right hind [Right hind [Right hind [Clinical signs of the affected foot/feet [Increased digital p [Focal sole pain in Generalised dorsa Coronary band sw Coronary band de 8. Clinical signs of the affected foot/feet Divergent growth Deviation in dorsa [ES NO Service NO	DIDN'T ASSESS	(or feet if) (please cro please cross o NO	bilaterally affected)
Right fore [Left fore [Right hind [Increased doot/feet Increased hoot/feet Focal sole pain in Generalised dorsa Coronary band sw Coronary band de 8. Clinical signs of th affected foot/feet Divergent growth Deviation in dorsa Wall separation [ES NO Service NO	DIDN'T ASSESS	(or feet if) (please cro please cross o NO	bilaterally affected)
Right fore [Left fore [Right hind [Increased doot/feet [Increased hoof tent Focal sole pain in Generalised dorsa Coronary band sw Coronary band de [Clinical signs of th affected foot/feet Divergent growth Deviation in dorsa	ES NO Service NO	DIDN'T ASSESS	(or feet if) (please cro please cross o NO	bilaterally affected)
Right fore [Left fore [Right hind [Right hind [Right hind [Left hind [7. Clinical signs of the affected foot/feet Increased digital p Increased hoof ter Focal sole pain in Generalised dorsa Coronary band sw Coronary band de 8. Clinical signs of the affected foot/feet Divergent growth Deviation in dorsa Wall separation Flat sole Convex sole	ES NO	DIDN'T ASSESS	(or feet if) (please cro please cross o NO	bilaterally affected)
Right fore [Left fore [Right hind [Right hind [Left hind [7. Clinical signs of th affected foot/feet [Increased digital p [Focal sole pain in [Generalised dorsa [Coronary band sw [Coronary band de [8. Clinical signs of th affected foot/feet [Divergent growth [Deviation in dorsa [Wall separation [Flat sole [Convex sole [Abnormally wide w [ES NO	DIDN'T ASSESS	(or feet if) (please cro please cross o NO	bilaterally affected)
Right fore [Left fore [Right hind [Right hind [Left hind [7. Clinical signs of th affected foot/feet [Increased digital p [Focal sole pain in [Generalised dorsa Coronary band sw Coronary band de [8. Clinical signs of th affected foot/feet [Divergent growth [Deviation in dorsa [Wall separation [Flat sole [Convex sole [Abnormally wide w [ES NO	DIDN'T ASSESS	(or feet if) (please cro please cross o NO	bilaterally affected)

