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Computed Tomographic Findings in Cats with Mycobacterial Infection

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Keywords:	Feline, Mycobacteriosis, Computed tomography, Infection, Diagnosis
Abstract:	Objectives The objective of this study was to describe the imaging findings in computed tomography (CT) associated with confirmed mycobacterial infection in cats. Methods CT images from 20 cats with confirmed mycobacterial disease were retrospectively reviewed. Five cats underwent conscious full-body CT in a VetMouseTrapTM device. All other cats had thoracic CT performed under general anaesthesia, with the addition of CT investigation of the head/neck, abdomen and limbs in some cases. Results Mycobacterial infection was seen most frequently in adult (mean age 7.4 years; range 0.6-14 years) neutered male cats (11/20). The most common infections were Mycobacterium microti (6/20) and Mycobacterium bovis (6/20). CT abnormalities were most commonly seen in the thorax, consisting of bronchial (9/20), alveolar (8/20), ground glass (6/20) or structured interstitial (15/20) lung patterns, which were often mixed. Tracheobronchial, sternal and cranial mediastinal lymphadenomegaly were common (16/20). Other abnormalities included abdominal (8/13) or peripheral (10/18) lymphadenomegaly, hepatosplenomegaly (7/13), mixed osteolytic/osteoproliferative skeletal lesions (7/20), and cutaneous or subcutaneous soft tissue masses/nodules (4/20). Conclusions and relevance CT of feline mycobacteriosis shows a wide range of abnormalities often involving multiple organ systems and mimicking many other feline

diseases. Mycobacteriosis should be considered in the differential diagnosis of thoracic, abdominal and skeletal disorders in cats.



1	Computed Tomographic Findings in Cats with Mycobacterial Infection
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22	Keywords - feline, mycobacteriosis, computed tomography, infection, diagnosis
23	

26 **Objectives**

27 The objective of this study was to describe the imaging findings in computed tomography

28 (CT) associated with confirmed mycobacterial infection in cats.

29 Methods

30 CT images from 20 cats with confirmed mycobacterial disease were retrospectively 31 reviewed. Five cats underwent conscious full-body CT in a VetMouseTrapTM device. All 32 other cats had thoracic CT performed under general anaesthesia, with the addition of CT 33 investigation of the head/neck, abdomen and limbs in some cases.

34 **Results**

Mycobacterial infection was seen most frequently in adult (mean age 7.4 years; range 35 36 0.6-14 years) neutered male cats (11/20). The most common infections were 37 Mycobacterium microti (6/20) and Mycobacterium bovis (6/20). CT abnormalities were 38 most commonly seen in the thorax, consisting of bronchial (9/20), alveolar (8/20), ground 39 glass (6/20) or structured interstitial (15/20) lung patterns, which were often mixed. 40 Tracheobronchial, sternal and cranial mediastinal lymphadenomegaly were common 41 (16/20). Other abnormalities included abdominal (8/13) or peripheral (10/18)42 lymphadenomegaly, hepatosplenomegaly (7/13), mixed osteolytic/osteoproliferative 43 skeletal lesions (7/20), and cutaneous or subcutaneous soft tissue masses/nodules (4/20).

44 **Conclusions and relevance**

45 CT of feline mycobacteriosis shows a wide range of abnormalities often involving
46 multiple organ systems and mimicking many other feline diseases. Mycobacteriosis

- 47 should be considered in the differential diagnosis of thoracic, abdominal and skeletal
- 48 disorders in cats.
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- 51

52 Introduction

Feline mycobacteriosis is a worldwide veterinary health concern, and although definitive data on case numbers worldwide are lacking, mycobacterial infections in cats have been recognised with increasing frequency in the UK, as well as being seen in many other countries. Mycobacterial disease in domestic cats can result from infection by one of a number of species. The most commonly identified mycobacteria include *Mycobacterium microti* and *Mycobacterium bovis*, which are primary pathogens and members of the tuberculous complex group of mycobacteria.¹⁻³ Non-tuberculous mycobacterial species are less commonly identified within clinically affected cats.⁴

60 Clinical presentation of mycobacterial infection in cats is variable, and is dependent primarily on the species of mycobacteria involved and, importantly, the route of infection.^{2,5-7} Historically, alimentary 61 62 lesions resulting from ingestion of milk from cows infected with M. bovis were most common; however 63 with overall reduction of tuberculosis in the national bovine herd since the early 1900's and widespread 64 pasteurisation of milk this is no longer the case.⁸ Single or multiple cutaneous lesions with or without 65 lymph node involvement, and characteristically affecting the so-called 'fight and bite sites' (such as the 66 head and limbs), now represent the most common presentation of mycobacterial infection in cats: they typically result from infection acquired from prey species.^{3,9} Infection acquired through inhalation or 67 68 ingestion, resulting in respiratory or alimentary disease, is seen less frequently. The clinical presentation of 69 these forms of disease, and of disseminated disease resulting from haematogenous spread of infection, can 70 include non-specific signs such as weight loss, anorexia, coughing, anaemia, vomiting/diarrhoea, 71 hepatosplenomegaly, generalised lymphadenopathy and pyrexia.⁷

72 Definitive diagnosis of mycobacterial disease in cats can present significant problems, in part due 73 to difficulties in sample handling, and limitations in the available laboratory diagnostic techniques. As 74 such, mycobacterial infections are likely underdiagnosed within the domestic cat population. In addition to 75 significant morbidity resulting from primary infection, subclinical infection and recurrence of infection 76 following treatment are common.⁷ Since significant and potentially fatal multisystemic disease can result 77 from infection with mycobacterial species, and since there are potential zoonotic risks associated with all 78 members of the tuberculosis complex,^{7,10} identification and correct handling of potential cases is of the 79 upmost importance.

80 Previous publications detailing the diagnostic imaging findings in cats with confirmed 81 mycobacterial infection are limited to a single retrospective case series looking at survey radiographic 82 changes involving 33 cats,¹¹ and a number of isolated case reports describing the radiographic features of 83 feline mycobacteriosis.¹²⁻¹⁴ Computed tomography is increasingly available to the veterinary community, 84 and it offers significant advantages over survey radiography by eliminating superimposition of anatomy, 85 having superior contrast resolution and being able to clarify intrathoracic lesions where radiographic findings are negative or non-specific.^{15,16} In addition, the decreased scan times which are achievable with 86 87 modern multi-detector scanners make CT a valuable tool in investigation of multisystemic disease in 88 clinically compromised patients. The CT features of mycobacterial disease in cats have not been described 89 previously. The aim of this paper was to review CT images from a large number of cats with confirmed 90 mycobacteriosis and to describe the range of abnormalities that can occur.

91

92 Materials and Methods

This study comprises a descriptive, retrospective case series. CT studies carried out between August 2009 and January 2015, of cats with confirmed mycobacterial infection were submitted to one of the authors (DGM). Inclusion criteria consisted of: (i) confirmation of mycobacterial infection and (ii) a CT study of diagnostic quality. To confirm mycobacterial involvement, aspirated and/or biopsy samples of affected tissue had been stained with Ziehl-Neelson (ZN) and found to have changes indicative of mycobacteriosis.¹ Where possible, tissue culture,¹⁷ interferon-gamma release assay, or PCR testing had been used to identify which mycobacterial species was involved.^{4,18,19}

Pseudonymised CT studies of the confirmed mycobacterial cases were examined without knowledge of specific clinical information by a third year diagnostic imaging resident who was however informed about the topic of the study (AM). To prevent bias by the assumption of disease, CT studies covering the thorax and other body parts from an additional ten cats with confirmed non-mycobacterial diseases were included and also pseudonymised. Images were evaluated using dedicated DICOM viewer software (Osirix, Geneva, Switzerland, version 5.8.5-64bit) on a computer workstation (Apple Mac Pro, Apple, USA) with a calibrated LCD flat screen monitor (Apple Cinemax Display, 30 inch, Apple, USA). 107 During the course of image evaluation, multi-planar reconstructions, maximum and minimum intensity108 projections and variable windowing settings were used according to the preferences of the viewer.

109 CT studies were reviewed for the following diagnostic criteria: bronchial thickening; alveolar 110 pattern; ground glass opacity or structured interstitial lung change; evidence of pleural or pericardial 111 effusion, or pleural/mediastinal thickening; thoracic, abdominal or peripheral lymphadenomegaly, or lymph 112 node mineralisation; abdominal organomegaly, peritoneal effusion, other abdominal organ-associated 113 lesions; osteolysis or osteoproliferative changes; cutaneous/subcutaneous/oral/nasal lesions; vascular and 114 dystrophic soft tissue calcification. The extent of any abnormality was characterised as focal, multifocal, or 115 diffuse. The degree of each change was graded as absent/normal, mild, moderate or severe.

116

117 **Results**

118 Twenty cats met the inclusion criteria. After all image interpretive data had been collected the 119 additional ten non-mycobacterial cat studies were identified and their data were excluded from further 120 analysis. The most common infections were M. microti and M. bovis, confirmed in 6/20 cases each. A non-121 specified *M. tuberculosis* complex species was described in one case and in the remaining 7/20 cases the 122 species involved was not known. Eleven of the 20 cats were neutered males and 9/20 were neutered 123 females. The study group comprised 7/20 Domestic Short Hair, 4/20 Siamese, 2/20 Domestic Long Hair 124 and 1/20 of each of the following; Persian, Birman, Norwegian Forest Cat, Burmilla, British Short Hair, 125 Bengal and Maine Coon cats. The age of one cat was not known. For the remaining cats the mean age was 126 7.4 ± 3.8 years (range 0.6-14 years).

Five of the 20 cats underwent conscious full-body CT in a specific containing device (VetMouseTrap[™], University of Illinois at Urbana-Champaign, Urbana, IL).²⁰ The remaining 15 cats were scanned under general anaesthesia, with images of the following body regions obtained: thorax only (2), head/neck and thorax (3), thorax and abdomen (4), head/neck, thorax and abdomen (2), head/neck, thorax, abdomen and single forelimb (2), head, thorax, bilateral tarsi/elbows (1), thorax and single hind limb (1). Intravenous contrast medium (iopamidol or iohexol, 600-700mg I/kg) was administered to 12/20

133 cats, and post-contrast images of some or all body parts were obtained. Use of contrast medium depended

on the findings in the pre-contrast images, the clinical condition of the cat, and the preferences of theattending radiologist and primary clinician in each case.

136 Within the evaluated imaging studies, thoracic abnormalities were noted in 19/20 cases. Diffuse 137 bronchial thickening was present in 9/20 cats; being mild in eight cases and moderate in one. Eight cats 138 showed a focal alveolar pattern; mild in two cases, moderate in three cases and severe in three cases (Figure 139 1(a)). Diffuse or patchy ground glass opacity was noted in 6/20 cats; mild in three cases, moderate in two 140 cases and severe in one case. The most common pulmonary parenchymal change was a diffuse structured 141 interstitial pattern, which was present in 15/20 cats, being either nodular (7/15) or reticulonodular (8/15) in 142 nature; mild in six cases, moderate in five cases and severe in four cases (Figure 1(b,c)). Thoracic CT 143 images of 14/20 cats were considered to show a mixed pulmonary pattern, with a single pattern present in 144 4/20 cases. The appearance of the pulmonary parenchyma was normal in 2/20 cats, though one of these had 145 a thoracic lymphadenopathy despite normal lungs. Of the 20 cats, 16 had sternal, cranial mediastinal and/or 146 tracheobronchial lymphadenomegaly (Figure 2). Moderate lymphadenomegaly affecting the sternal or 147 tracheobronchial nodes was most common. One cat had moderate mineralisation of an enlarged cranial 148 mediastinal lymph node.

None of the cats had any evidence of pleural or pericardial effusion. One cat showed mild, diffuse pleural thickening. One cat showed mild mineralisation of the aortic root. Two cats had regions of cavitation within the lungs, associated in both cases with focal or multifocal nodular or alveolar changes (Figure 3(a)). Three cats had scattered foci of mineralisation within the lungs, again associated with other focal parenchymal changes (Figure 3(b)).

154 Thirteen of the 20 cats had imaging studies that included the abdomen. Abdominal 155 lymphadenomegaly was present in 8/13 cases and was typically generalised. The lymph nodes affected 156 could not always be individually identified, but included those of the celiac and cranial mesenteric centres, 157 which variably comprised the hepatic, splenic, gastric, pancreaticoduodenal, jejunal and colic nodes. 158 Lymphadenomegaly was mild in two cats, moderate in four cats and severe in two cats. In one cat with a 159 generalised moderate abdominal lymphadenomegaly, mild mineralisation of a mesenteric lymph node was 160 present (Figure 4(a)). Mild hepatomegaly was present in 3/13 cats and moderate hepatomegaly in 1/13. 161 Mild splenomegaly was present in 6/13 cats and moderate splenomegaly in 1/13. Two cats with

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splenomegaly (one mild and one moderate) were also noted to show heterogeneity within the splenic parenchyma following contrast medium administration. Additional abdominal organ changes were noted in 3/13 cats; one had a moderately enlarged pancreas, one had multiple nodules within both kidneys, and one an irregular outline to the left kidney. Peritoneal effusion was not noted in any cat.

166 The appearance of the peripheral lymph nodes was assessed in 18/20 cats. The two cats not 167 included in this assessment had CT studies of the thorax only, without inclusion of any extra-thoracic 168 lymph node group. In 10/18 cases peripheral lymphadenomegaly was present, mild in 3/18, moderate in 169 2/18 and severe in 5/18. In 8/10 cats with lymphadenomegaly, the most significant enlargement was noted 170 in the mandibular and medial retropharyngeal nodes (Figure 4(b)); however, multifocal 171 lymphadenomegaly, involving the superficial cervical (prescapular), axillary, inguinal and/or popliteal 172 nodes was variably present. In the other 2/10 cats, the head and neck were not imaged, but enlargement of 173 the superficial cervical and popliteal lymph nodes was noted respectively. Five of the eight cats in which 174 peripheral lymphadenomegaly was not noted underwent conscious CT in the VetMouseTrap[™] device and 175 three underwent CT studies which did not include the head and neck.

176 Focal osteolytic lesions were present in 7/20 cats (although it was not possible to assess the entire 177 skeleton in 15 cats as they did not have full body CT examinations); changes were mild in four cases and 178 moderate in three cases. These lesions affected the nasal bridge in three cats and the limbs in the remaining 179 four, and were predominantly characterised by regions of cortical lysis (5/7) or erosive lesions at joint 180 surfaces (2/7) (Figure 5). An associated pathological long bone fracture was present in one case. In all but 181 one of these cases osteoproliferative changes, either periosteal reaction or periarticular osteophytosis, were 182 noted in the same location as the osteolytic change. The osteoproliferation was mild in three cases and 183 severe in three cases; however, the degree of proliferative change did not necessarily correlate with the 184 degree of lytic change in each case.

Cutaneous or subcutaneous lesions were only infrequently present within the studies evaluated. Focal mass lesions over the nasal bridge were noted in 2/20 cats, graded moderate in one and severe in one. One other cat had a small amount of fluid accumulation and soft tissue thickening in the dorsal nasal chambers. Each of these lesions was adjacent to bony abnormalities. A focal, but extensive, mass lesion was noted along the ventral head and neck of one cat. One cat was found to have multiple, widely distributed, subcutaneous nodules. Diffuse extra-thoracic dystrophic soft tissue mineralisation was notnoted in any cat.

192

193 **Discussion**

Mycobacteriosis in cats is known to be a highly variable disease, and should always be considered as a possible differential diagnosis in cases which present with multisystemic signs. Mycobacterial disease is likely under-recognised, primarily due to a lack of awareness of the full spectrum of changes which can be associated with it.

Mycobacterial infection is most commonly seen in adult, neutered male cats consistent with the results of our study.⁹ Domestic Short Hair cats predominate in our study, but to a lesser degree than noted in the previous radiographic case series (36% vs 87%).¹¹ The reason for this is unknown, but may reflect a higher proportion of pedigree cats within a referral population, which are therefore more likely to undergo advanced imaging.

203 CT abnormalities of the thorax were commonly noted, being present in all but one cat. However, 204 multisystemic abnormalities were also common, with changes affecting more than one anatomical region in 205 all but five cases. Of these five, three had abnormalities detected on clinical examination which were not 206 appreciable on the CT images. In cats, systemic mycobacterial infection is most commonly caused by *M*. 207 *bovis* or *M microti*,^{3,9,21-24} and our results are consistent with this.

Previous reports of radiographic findings in cats with mycobacterial disease described a mild 208 209 predominance of a mixed lung pattern (ie, a combination of bronchial, alveolar and/or interstitial 210 changes),^{19,11-13,22-24} but distinct alveolar, bronchial or interstitial patterns in isolation were also identified.¹¹ 211 Interestingly, where cases in our study displayed mixed lung patterns, bronchial thickening and ground 212 glass opacity were most commonly graded as mild, whereas alveolar pattern and structured interstitial 213 patterns were more likely to be moderate or severe. This is interesting for two reasons. Firstly, the mild 214 bronchial and unstructured interstitial patterns are of a degree that comparable changes may not be easily 215 visible radiographically, or may be attributed to expiratory or underexposed radiographs, or 216 superimposition of other structures. In a radiographic study it is therefore possible that only a more 217 significant overlying alveolar or nodular pattern may be recognised, leading to classification as a single

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218 lung pattern. As superimposition effects are eliminated by CT, it becomes easier to identify these mild 219 changes in addition to the more marked ones. Secondly, a mild bronchial or unstructured interstitial pattern 220 may be indicative of concurrent conditions, such as low-grade allergic airway disease, rather than being 221 directly related to an active mycobacterial infection.^{25,26} Differentiation of these may not be possible.

222 Within our study, the most commonly encountered single lung pattern was structured interstitial. 223 However, these cases could be further subdivided into cases displaying a nodular pattern, comprising 224 scattered rounded hyperattenuating foci, and a reticulonodular pattern, where rounded foci and linear or 225 sickle-shaped hyperattenuating structures overlie to give a more complex overall pattern. In humans, a 226 faint, diffuse reticulonodular pattern is considered characteristic of miliary tuberculosis.²⁷ While nodular 227 and reticulonodular patterns are distinguishable on good quality radiographs in cats, the distinction is only 228 rarely made in veterinary imaging. On CT however, the difference is more easily appreciable. The 229 diagnostic and prognostic significance of the variable patterns in feline patients is currently unknown, but 230 certainly warrants further investigation, as a structured interstitial pattern is a common finding in many 231 feline lung pathologies (eg, pulmonary fibrosis, metastatic neoplasia, eosinophilic bronchopneumopathy 232 and a wide range of infectious pneumonias).

233 It is interesting to note that within our study population, two cats were found to have cavitations 234 within their lungs. While this feature is relatively common in both humans and dogs with tuberculosis²⁸⁻³⁰ it was not noted in any case in the previous radiographic study of cats,¹¹ and the only paper which describes 235 236 cavitating tubercles in cats was published in 1949.⁵ The lesions noted in the two cats in this study were 237 small (<1 cm) and were contained within regions of nodular or alveolar change. It is possible therefore that 238 they may not have been visible on radiographs, again highlighting the advantage of cross sectional imaging. 239 Alternatively, this may indeed reflect a rare occurrence in feline patients, which has occurred coincidentally 240 within our study population. In either case, it is an important characteristic to recognise, as cavitated lung masses are occasionally identified in feline patients with lung neoplasia,^{16,25} and the potential for 241 242 misdiagnosis exists in cats with mycobacteriosis which show this feature. In addition this should be 243 recognised because these cats likely pose an increased zoonotic risk compared with those showing the more 244 typical structured interstitial pattern as they may allow mycobacteria to gain access to the upper airways.

245 Thoracic lymphadenomegaly is a feature of numerous pulmonary and multisystemic conditions in 246 cats including, but not limited to, infiltrative and metastatic neoplasia, hypereosinophilic syndromes and 247 systemic mycosis/bacteriosis.³¹ As expected thoracic lymphadenomegaly was commonly noted within our 248 study population, but in contrast to the findings of the previous radiographic paper, mild and moderate 249 enlargement predominated over severe.¹¹ This may reflect the difficulty in recognising minor changes on 250 radiographs. It is also worth noting that even given the superior contrast resolution of CT, with mild 251 lymphadenomegaly, particularly in the perihilar region; changes were more easily appreciated in post-252 contrast studies when compared with pre-contrast. This suggests that there is value in performing post-253 contrast scans in all cases (unless there is a clinical contraindication), which was not standard practice 254 within our study population.

255 Mineralisation of thoracic lymph nodes and pulmonary parenchyma can result from chronic 256 inflammation associated with mycobacterial infection;^{9,13,32} it is also seen in cases of both primary and 257 metastatic pulmonary neoplasia, and chronic airways disease.²⁵ In either case, it is a finding which most 258 likely relates to the disease process that is present. In contrast, mild aortic root calcification, such as that 259 seen in one case (a seven year old cat) in our study is, in our experience, an occasional finding in middle 260 aged to older cats, and not necessarily related to clinical disease.

261 While peripheral and abdominal lymphadenomegaly were relatively common within the study 262 population, it is possible that the number of cases with mild or moderate lymphadenomegaly in the head 263 and neck was artificially low. This is because all cases recorded as having normal peripheral lymph nodes 264 on physical examination either did not undergo imaging of the head and neck, or were scanned conscious 265 within a VetMouseTrapTM device. The protocol for these scans involved a short scan time (in order to 266 minimise movement) resulting in a relatively large slice thickness and consequently a reduced longitudinal 267 resolution. This can compromise assessment of small structures so it is possible that mild or moderate 268 abnormalities of the head and neck, such as lymphadenomegaly, may have been overlooked. Other 269 abdominal changes such as hepatomegaly, splenomegaly, renal and pancreatic changes were noted 270 relatively infrequently and were mild or moderate in extent, consistent with previous reports.^{9,11,12,33}

Two distinct manifestations of skeletal disease were noted within our population. The lesions characterised by cortical lysis likely represent sites of primary bacterial inoculation and as such are

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273 consistent in location with 'fight and bite' injuries; whereas periarticular abnormalities are consistent with 274 an infectious polyarthritis resulting from haematogenous dissemination of bacteria. It is interesting to note 275 that the appendicular skeletal lesions in this study were clinically evident, and affected regions were 276 intentionally included in the imaging studies. Clinically silent skeletal lesions may have been overlooked as 277 the limbs were excluded from the majority of studies. The only studies in which the limbs were included in 278 full were those performed using the VetMouseTrapTM device and it is possible that subtle or focal regions 279 of bone lysis or proliferation may not have been recognised due to the lower resolution of these studies.

Cutaneous lesions were noted infrequently in this study. While this may initially seem surprising, given that cutaneous lesions represent a common presentation of mycobacteriosis,^{3,9} it reflects the fact that CT imaging is more likely to be employed in cases presenting with systemic disease, or used as a staging tool in cats with clinically evident focal skin lesions without requirement for imaging of the lesions themselves. The presence of intranasal changes in one cat is interesting, as these are indicative of a mycobacterial rhinitis, a manifestation of respiratory mycobacteriosis which may not be commonly recognised.

287 There are a number of limitations to this study. The most significant of these is that although 288 mycobacteriosis was confirmed in each case, histopathology on all involved tissues was not typically 289 performed; therefore, it is not possible to confirm that all changes seen were due to mycobacterial infection. 290 Due to the inherent difficulties in confirmation of mycobacterial infection, the time lapse between 291 acquisition of CT images and definitive confirmation of diagnosis was very variable; it extended to four 292 years and nine months in one case (though a lapse of one to four months was more typical). In all cases 293 however, at the time of imaging, the combination of clinical and pathological findings gave sufficient 294 confidence in the diagnosis to allow treatment to be instigated; imaging was used to stage the cases and so 295 guide the intensity and duration of treatment. The evolution of changes over time in association with 296 treatment has not been described, and will be interesting to explore in the future. Finally, given the 297 retrospective nature of this study there are inconsistencies between cases with respect to factors such as 298 regions imaged and use of contrast medium. This leads to a bias in our results, and may underestimate 299 subclinical disease, particularly affecting the peripheral structures. As mentioned, the limited resolution of 300 smaller structures on VetMouseTrap[™] scans contributes to this. However, the advantages of this technique

- 301 for disease screening, particularly in clinically compromised patients should not be ignored, and as faster
- 302 scanners become more commonplace many of the resolution difficulties will be eliminated.
- 303

304 Conclusions

305	As expected, the majority of CT changes noted in this study represent multisystemic disease,
306	typically with combinations of pulmonary infiltration, lymphadenomegaly and organomegaly. These
307	changes are strongly suggestive of infiltrative disease, differentials for which can include neoplasia (such as
308	lymphoma or mast cell disease), chronic inflammation/infectious processes (mycobacteriosis, feline
309	infectious peritonitis or systemic mycosis), hypereosinophilic syndrome and amyloidosis. ²⁴ While no
310	abnormality has been recognised that is specific for mycobacteriosis, it is important that the potential for
311	mycobacterial infection is considered when these types of changes are identified in feline patients,
312	especially if they have non-specific clinical signs. In addition, when managing patients with a diagnosis of
313	mycobacteriosis, the potential for widespread clinical and sub-clinical abnormalities must be considered
314	and investigated in full.
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316	
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319	
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321	
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- 326
- 327 References

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328	1.	Snider WR. Tuberculosis in canine and feline populations. Review of the literature. Am Rev Respir
329	<i>Dis</i> 197	1; 104: 877-87.
330	2.	Snider WR, Cohen D, Reif JS, et al. Tuberculosis in canine and feline populations. Study of high
331	risk pop	oulations in Pennsylvania, 1966-1968. Am Rev Respir Dis 1971; 104: 866-76.
332	3.	Gunn-Moore DA, McFarland SE, Brewer JI, et al. Mycobacterial disease in cats in Great Britain:
333	I. Cultu	re results, geographical distribution and clinical presentation of 339 cases. J Feline Med Surg 2011;
334	13: 934-44.	
335	4.	Malik R, Smits B, Reppas G, et al. Ulcerated and nonulcerated nontuberculous cutaneous
336	mycoba	cterial granulomas in cats and dogs. Vet Dermatol 2013; 24: 146-53.
337	5.	Jennings AR. The distribution of tuberculous lesions in the dog and cat, with reference to the
338	pathoge	nesis. Vet Rec 1949; 61: 380-84.
339	6.	Huitema H and van Vloten J. Murine tuberculosis in a cat. Antonie Van Leeuwenhoek 1960; 26:
340	235-40.	
341	7.	Gunn-Moore D. Mycobacterial infections in cats and dogs. In: Ettinger S and Feldman E (eds).
342	Textboo	ok of Veterinary Internal Medicine. 7th ed. Philadelphia: Saunders, 2010: 875-81.
343	8.	de la Rua-Domenech R. Human Mycobacterium bovis infection in the United Kingdom:
344	Incident	ce, risks, control measures and review of the zoonotic aspects of bovine tuberculosis. Tuberculosis
345	(Edinb)	2006; 86: 77-109.
346	9.	Gunn-Moore D, Dean R and Shaw S. Mycobacterial infections in cats and dogs. In Pract 2010;
347	32: 444-	-52.
348	10.	Cima G. Cat transmits TB to humans in UK. J Am Vet Med Assoc 2014; 244: 1116.
349	11.	Bennett AD, Lalor S, Schwarz T, et al. Radiographic findings in cats with mycobacterial
350	infection	ns. J Feline Med Surg 2011; 13: 718-24.
351	12.	Baral RM, Metcalfe SS, Krockenberger MB, et al. Disseminated Mycobacterium avium infection
352	in young	g cats: overrepresentation of Abyssinian cats. J Feline Med Surg 2006; 8: 23-44.

353 13. Foster SF, Martin P, Davis W, et al. Chronic pneumonia caused by *Mycobacterium*354 *thermoresistibile* in a cat. *J Small Anim Pract* 1999; 40: 433-8.

355 14. Paltrinieri S. Tuberculosis in the dog and cat. *Nuova Vet* 1930; 6: 7.

356	15.	Prather AB, Berry CR and Thrall DE. Use of Radiography in Combination with Computed
357	Tomog	graphy for the Assessment of Noncardiac Thoracic Disease in the Dog and Cat. Vet Radiol
358	Ultrase	ound 2005; 46: 114-21.
359	16.	Henninger W. Use of computed tomography in the diseased feline thorax. J Small Anim Pract
360	2003; 4	44: 56-64.
361	17.	Daniel R, Evans H, Rolfe S, et al. Outbreak of tuberculosis caused by Mycobacterium bovis in
362	golden	Guernsey goats in Great Britain. Vet Rec 2009; 165: 335-42.
363	18.	Rhodes SG, Gruffydd-Jones T, Gunn-Moore D, et al. Interferon-gamma test for feline
364	tubercu	alosis. Vet Rec 2008; 162: 453-5.
365	19.	Rhodes SG, Gruffydd-Jones T, Gunn-Moore D, et al. Adaptation of IFN-gamma ELISA and
366	ELISP	OT tests for feline tuberculosis. Vet Immunol Immunopathol 2008; 124: 379-84.
367	20.	Oliveira CR, Ranallo FN, Pijanowski GJ, et al. The Vetmousetrap™: A Device for Computed
368	Tomog	graphic Imaging of the Thorax of Awake Cats. Vet Radiol Ultrasound 2011; 52: 41-52.
369	21.	Greene CE and Gunn-Moore D. Mycobacterial Infections. In: Greene CE (ed). Infectious Diseases
370	of the 1	Dog and Cat. 4th ed. Philadelphia: Elsevier Health Services, 2012: 495-510.
371	22.	Gunn-Moore DA, Jenkins PA and Lucke VM. Feline tuberculosis: a literature review and
372	discuss	sion of 19 cases caused by an unusual mycobacterial variant. Vet Rec 1996; 138: 53-8.
373	23.	Barry M, Taylor J and Woods JP. Disseminated Mycobacterium avium infection in a cat. Can Vet
374	J 2002	; 43: 369-71.
375	24.	Gow AG. What is your diagnosis? Mycobacterial infection. J Small Anim Pract 2006; 47: 484-5.
376	25.	Thrall DE. The Canine and Feline Lung. In: Thrall DE (ed). Textbook of Veterinary Diagnostic
377	Imagin	g. 6th ed. St. Louis: Elsevier Saunders, 2013: 608-31.
378	26.	Schwarz T and Johnson V. Lungs and Bronchi. In: Schwarz T and Saunders J (eds). Veterinary
379	Compu	ited Tomography. Chichester: Wiley-Blackwell, 2011: 261-78.

- 380 27. Velez MG and Velez VJ, Jr. Diffuse reticulonodular infiltrates. *Cleve Clin J Med* 2012; 79: 16-7.
- 381 28. Krysl J, Korzeniewska-Kosela M, Muller NL, et al. Radiologic features of pulmonary
- tuberculosis: an assessment of 188 cases. *Can Assoc Radiol J* 1994; 45: 101-7.
- 383 29. Gadkowski LB and Stout JE. Cavitary pulmonary disease. *Clin Microbiol Rev* 2008; 21: 305-33.

384 30. Olsson SE. On tuberculosis in the dog; a study with special reference to x-ray diagnosis. Cornell 385 Vet 1957; 47: 193-219.

386 31. Baines E. The mediastinum. In: Schwarz T and Johnson V (eds). BSAVA manual of canine and

387 feline thoracic imaging. Gloucester: BSAVA, 2008: 177-99.

- 388 32. Hix JW, Jones TC and Karlson AG. Avian tubercle bacillus infection in the cat. J Am Vet Med
- 389 Assoc 1961; 138: 641-7.
- 390 33. Knippel A, Hetzel U and Baumgartner W. Disseminated Mycobacterium avium-intracellulare
- 391 Infection in a Persian cat. J Vet Med 2004; 51: 464-6.

1	Computed Tomographic Findings in Cats with Mycobacterial Infection
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22	Keywords – feline, mycobacteriosis, computed tomography, infection, diagnosis
23	

26 **Objectives**

27 The objective of this study was to describe the imaging findings in computed tomography

28 (CT) associated with confirmed mycobacterial infection in cats.

29 Methods

30 CT images from 20 cats with confirmed mycobacterial disease were retrospectively 31 reviewed. Five cats underwent conscious full-body CT in a VetMouseTrapTM device. All 32 other cats had thoracic CT performed under general anaesthesia, with the addition of CT 33 investigation of the head/neck, abdomen and limbs in some cases.

34 **Results**

Mycobacterial infection was seen most frequently in adult (mean age 7.4 years; range 35 36 0.6-14 years) neutered male cats (11/20). The most common infections were 37 Mycobacterium microti (6/20) and Mycobacterium bovis (6/20). CT abnormalities were 38 most commonly seen in the thorax, consisting of bronchial (9/20), alveolar (8/20), ground 39 glass (6/20) or structured interstitial (15/20) lung patterns, which were often mixed. 40 Tracheobronchial, sternal and cranial mediastinal lymphadenomegaly were common 41 (16/20). Other abnormalities included abdominal (8/13) or peripheral (10/18)42 lymphadenomegaly, hepatosplenomegaly (7/13), mixed osteolytic/osteoproliferative 43 skeletal lesions (7/20), and cutaneous or subcutaneous soft tissue masses/nodules (4/20).

44 **Conclusions and relevance**

45 CT of feline mycobacteriosis shows a wide range of abnormalities often involving
46 multiple organ systems and mimicking many other feline diseases. Mycobacteriosis

- 47 should be considered in the differential diagnosis of thoracic, abdominal and skeletal
- 48 disorders in cats.
- 49
- 50
- 51

52 Introduction

Feline mycobacteriosis is a worldwide veterinary health concern, and although definitive data on case numbers worldwide are lacking, mycobacterial infections in cats have been recognised with increasing frequency in the UK, as well as being seen in many other countries. Mycobacterial disease in domestic cats can result from infection by one of a number of species. The most commonly identified mycobacteria include *Mycobacterium microti* and *Mycobacterium bovis*, which are primary pathogens and members of the tuberculous complex group of mycobacteria.¹⁻³ Non-tuberculous mycobacterial species are less commonly identified within clinically affected cats.⁴

60 Clinical presentation of mycobacterial infection in cats is variable, and is dependent primarily on the species of mycobacteria involved and, importantly, the route of infection.^{2,5-7} Historically, alimentary 61 62 lesions resulting from ingestion of milk from cows infected with M. bovis were most common; however 63 with overall reduction of tuberculosis in the national bovine herd since the early 1900's and widespread 64 pasteurisation of milk this is no longer the case.⁸ Single or multiple cutaneous lesions with or without 65 lymph node involvement, and characteristically affecting the so-called 'fight and bite sites' (such as the 66 head and limbs), now represent the most common presentation of mycobacterial infection in cats: they typically result from infection acquired from prey species.^{3,9} Infection acquired through inhalation or 67 68 ingestion, resulting in respiratory or alimentary disease, is seen less frequently. The clinical presentation of 69 these forms of disease, and of disseminated disease resulting from haematogenous spread of infection, can 70 include non-specific signs such as weight loss, anorexia, coughing, anaemia, vomiting/diarrhoea, 71 hepatosplenomegaly, generalised lymphadenopathy and pyrexia.⁷

72 Definitive diagnosis of mycobacterial disease in cats can present significant problems, in part due 73 to difficulties in sample handling, and limitations in the available laboratory diagnostic techniques. As 74 such, mycobacterial infections are likely underdiagnosed within the domestic cat population. In addition to 75 significant morbidity resulting from primary infection, subclinical infection and recurrence of infection 76 following treatment are common.⁷ Since significant and potentially fatal multisystemic disease can result 77 from infection with mycobacterial species, and since there are potential zoonotic risks associated with all 78 members of the tuberculosis complex,^{7,10} identification and correct handling of potential cases is of the 79 upmost importance.

80 Previous publications detailing the diagnostic imaging findings in cats with confirmed 81 mycobacterial infection are limited to a single retrospective case series looking at survey radiographic 82 changes involving 33 cats,¹¹ and a number of isolated case reports describing the radiographic features of 83 feline mycobacteriosis.¹²⁻¹⁴ Computed tomography is increasingly available to the veterinary community, 84 and it offers significant advantages over survey radiography by eliminating superimposition of anatomy, 85 having superior contrast resolution and being able to clarify intrathoracic lesions where radiographic findings are negative or non-specific.^{15,16} In addition, the decreased scan times which are achievable with 86 87 modern multi-detector scanners make CT a valuable tool in investigation of multisystemic disease in 88 clinically compromised patients. The CT features of mycobacterial disease in cats have not been described 89 previously. The aim of this paper was to review CT images from a large number of cats with confirmed 90 mycobacteriosis and to describe the range of abnormalities that can occur.

91

92 Materials and Methods

This study comprises a descriptive, retrospective case series. CT studies carried out between August 2009 and January 2015, of cats with confirmed mycobacterial infection were submitted to one of the authors (DGM). Inclusion criteria consisted of: (i) confirmation of mycobacterial infection and (ii) a CT study of diagnostic quality. To confirm mycobacterial involvement, aspirated and/or biopsy samples of affected tissue had been stained with Ziehl-Neelson (ZN) and found to have changes indicative of mycobacteriosis.¹ Where possible, tissue culture,¹⁷ interferon-gamma release assay, or PCR testing had been used to identify which mycobacterial species was involved.^{4,18,19}

Pseudonymised CT studies of the confirmed mycobacterial cases were examined without knowledge of specific clinical information by a third year diagnostic imaging resident who was however informed about the topic of the study (AM). To prevent bias by the assumption of disease, CT studies covering the thorax and other body parts from an additional ten cats with confirmed non-mycobacterial diseases were included and also pseudonymised. Images were evaluated using dedicated DICOM viewer software (Osirix, Geneva, Switzerland, version 5.8.5-64bit) on a computer workstation (Apple Mac Pro, Apple, USA) with a calibrated LCD flat screen monitor (Apple Cinemax Display, 30 inch, Apple, USA). During the course of image evaluation, multi-planar reconstructions, maximum and minimum intensity
 projections and variable windowing settings were used according to the preferences of the viewer.

109 CT studies were reviewed for the following diagnostic criteria: bronchial thickening; alveolar 110 pattern; ground glass opacity or structured interstitial lung change; evidence of pleural or pericardial 111 effusion, or pleural/mediastinal thickening; thoracic, abdominal or peripheral lymphadenomegaly, or lymph 112 node mineralisation; abdominal organomegaly, peritoneal effusion, other abdominal organ-associated 113 lesions; osteolysis or osteoproliferative changes; cutaneous/subcutaneous/oral/nasal lesions; or vascular and 114 dystrophic soft tissue calcification. The extent of any abnormality was characterised as focal, multifocal, or 115 diffuse. The degree of each change was graded as absent/normal, mild, moderate or severe.

116

117 **Results**

118 Twenty cats met the inclusion criteria. After all image interpretive data had been collected the 119 additional ten non-mycobacterial cat studies were identified and their data were excluded from further 120 analysis. The most common infections were M. microti and M. bovis, confirmed in 6/20 cases each. A non-121 specified *M. tuberculosis* complex species was described in one case and in the remaining 7/20 cases the 122 species involved was not known. Eleven of the 20 cats were neutered males and 9/20 were neutered 123 females. The study group comprised 7/20 Domestic Short Hair, 4/20 Siamese, 2/20 Domestic Long Hair 124 and 1/20 of each of the following; Persian, Birman, Norwegian Forest Cat, Burmilla, British Short Hair, 125 Bengal and Maine Coon cats. The age of one cat was not known. For the remaining cats the mean age was 126 7.4 ± 3.8 years (range 0.6-14 years).

Five of the 20 cats underwent conscious full-body CT in a specific containing device
(VetMouseTrap[™], University of Illinois at Urbana-Champaign, Urbana, IL).²⁰ The remaining 15 cats were
scanned under general anaesthesia, with images of the following body regions obtained: thorax only (2),
head/neck and thorax (3), thorax and abdomen (4), head/neck, thorax and abdomen (2), head/neck, thorax,
abdomen and single forelimb (2), head, thorax, bilateral tarsi/elbows (1), thorax and single hind limb (1).
Intravenous contrast medium (iopamidol or iohexol, 600-700mg I/kg) was administered to 12/20

133 cats, and post-contrast images of some or all body parts were obtained. Use of contrast medium depended

on the findings in the pre-contrast images, the clinical condition of the cat, and the preferences of theattending radiologist and primary clinician in each case.

136 Within the evaluated imaging studies, thoracic abnormalities were noted in 19/20 cases. Diffuse 137 bronchial thickening was present in 9/20 cats; being mild in eight cases and moderate in one. Eight cats 138 showed a focal alveolar pattern; mild in two cases, moderate in three cases and severe in three cases (Figure 139 1(a)). Diffuse or patchy ground glass opacity was noted in 6/20 cats; mild in three cases, moderate in two 140 cases and severe in one case. The most common pulmonary parenchymal change was a diffuse structured 141 interstitial pattern, which was present in 15/20 cats, being either nodular (7/15) or reticulonodular (8/15) in 142 nature; mild in six cases, moderate in five cases and severe in four cases (Figure 1(b,c)). Thoracic CT 143 images of 14/20 cats were considered to show a mixed pulmonary pattern, with a single pattern present in 144 4/20 cases. The appearance of the pulmonary parenchyma was normal in 2/20 cats, though one of these had 145 a thoracic lymphadenopathy despite normal lungs. Of the 20 cats, 16 had sternal, cranial mediastinal and/or 146 tracheobronchial lymphadenomegaly (Figure 2). Moderate lymphadenomegaly affecting the sternal or 147 tracheobronchial nodes was most common. One cat had moderate mineralisation of an enlarged cranial 148 mediastinal lymph node.

None of the cats had any evidence of pleural or pericardial effusion. One cat showed mild, diffuse pleural thickening. One cat showed mild mineralisation of the aortic root. Two cats had regions of cavitation within the lungs, associated in both cases with focal or multifocal nodular or alveolar changes (Figure 3(a)). Three cats had scattered foci of mineralisation within the lungs, again associated with other focal parenchymal changes (Figure 3(b)).

154 Thirteen of the 20 cats had imaging studies that included the abdomen. Abdominal 155 lymphadenomegaly was present in 8/13 cases and was typically generalised diffuse. The lymph nodes 156 affected could not always be individually identified, but included those of the celiac and cranial mesenteric 157 centres, which variably comprised the hepatic, splenic, gastric, pancreaticoduodenal, jejunal and colic 158 nodes. Lymphadenomegaly was mild in two cats, moderate in four cats and severe in two cats. In one cat 159 with a generalised moderate abdominal lymphadenomegaly, mild mineralisation of a mesenteric lymph 160 node was present (Figure 4(a)). Mild hepatomegaly was present in 3/13 cats and moderate hepatomegaly in 161 1/13. Mild splenomegaly was present in 6/13 cats and moderate splenomegaly in 1/13. Two cats with

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splenomegaly (one mild and one moderate) were also noted to show heterogeneity within the splenic parenchyma following contrast medium administration. Additional abdominal organ changes were noted in 3/13 cats; one had a moderately enlarged pancreas, one had multiple nodules within both kidneys, and one an irregular outline to the left kidney. Peritoneal effusion was not noted in any cat.

166 The appearance of the peripheral lymph nodes was assessed in 18/20 cats. The two cats not 167 included in this assessment had CT studies of the thorax only, without inclusion of any extra-thoracic 168 lymph node group. In 10/18 cases peripheral lymphadenomegaly was present, mild in 3/18, moderate in 169 2/18 and severe in 5/18. In 8/10 cats with lymphadenomegaly, the most significant enlargement was noted 170 in the mandibular and medial retropharyngeal nodes (Figure 4(b)); however, multifocal more diffuse 171 lymphadenomegaly, involving the superficial cervical (prescapular), axillary, inguinal and/or popliteal 172 nodes was variably present. In the other 2/10 cats, the head and neck were not imaged, but enlargement of 173 the superficial cervical and popliteal lymph nodes was noted respectively. Five of the eight cats in which 174 peripheral lymphadenomegaly was not noted underwent conscious CT in the VetMouseTrap[™] device and 175 three underwent CT studies which did not include the head and neck.

176 Focal osteolytic lesions were present in 7/20 cats (although it was not possible to assess the entire 177 skeleton in 15 cats as they did not have full body CT examinations); changes were mild in four cases and 178 moderate in three cases. These lesions affected the nasal bridge in three cats and the limbs in the remaining 179 four, and were predominantly characterised by regions of cortical lysis (5/7) or erosive lesions at joint 180 surfaces (2/7) (Figure 5). An associated pathological long bone fracture was present in one case. In all but 181 one of these cases osteoproliferative changes, either periosteal reaction or periarticular osteophytosis, were 182 noted in the same location as the osteolytic change. The osteoproliferation was mild in three cases and 183 severe in three cases; however, the degree of proliferative change did not necessarily correlate with the 184 degree of lytic change in each case.

Cutaneous or subcutaneous lesions were only infrequently present within the studies evaluated. Focal mass lesions over the nasal bridge were noted in 2/20 cats, graded moderate in one and severe in one. One other cat had a small amount of fluid accumulation and soft tissue thickening in the dorsal nasal chambers. Each of these lesions was adjacent to bony abnormalities. A focal, but extensive, mass lesion was noted along the ventral head and neck of one cat. One cat was found to have multiple, widely distributed, subcutaneous nodules. Diffuse extra-thoracic dystrophic soft tissue mineralisation was notnoted in any cat.

192

193 **Discussion**

Mycobacteriosis in cats is known to be a highly variable disease, and should always be considered as a possible differential diagnosis in cases which present with multisystemic signs. Mycobacterial disease is likely under-recognised, primarily due to a lack of awareness of the full spectrum of changes which can be associated with it.

Mycobacterial infection is most commonly seen in adult, neutered male cats consistent with the results of our study.⁹ Domestic Short Hair cats predominate in our study, but to a lesser degree than noted in the previous radiographic case series (36% vs 87%).¹¹ The reason for this is unknown, but may reflect a higher proportion of pedigree cats within a referral population, which are therefore more likely to undergo advanced imaging.

203 CT abnormalities of the thorax were commonly noted, being present in all but one cat. However, 204 multisystemic abnormalities were also common, with changes affecting more than one anatomical region in 205 all but five cases. Of these five, three had abnormalities detected on clinical examination which were not 206 appreciable on the CT images. In cats, systemic mycobacterial infection is most commonly caused by *M*. 207 *bovis* or *M microti*,^{3,9,21-24} and our results are consistent with this.

Previous reports of radiographic findings in cats with mycobacterial disease described a mild 208 209 predominance of a mixed lung pattern (ie, a combination of bronchial, alveolar and/or interstitial 210 changes),^{19,11-13,22-24} but distinct alveolar, bronchial or interstitial patterns in isolation were also identified.¹¹ 211 Interestingly, where cases in our study displayed mixed lung patterns, bronchial thickening and ground 212 glass opacity were most commonly graded as mild, whereas alveolar pattern and structured interstitial 213 patterns were more likely to be moderate or severe. This is interesting for two reasons. Firstly, the mild 214 bronchial and unstructured interstitial patterns are of a degree that comparable changes may not be easily 215 visible radiographically, or may be attributed to expiratory or underexposed radiographs, or 216 superimposition of other structures. In a radiographic study it is therefore possible that only a more 217 significant overlying alveolar or nodular pattern may be recognised, leading to classification as a single

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218 lung pattern. As superimposition effects are eliminated by CT, it becomes easier to identify these mild 219 changes in addition to the more marked ones. Secondly, a mild bronchial or unstructured interstitial pattern 220 may be indicative of concurrent conditions, such as low-grade allergic airway disease, rather than being 221 directly related to an active mycobacterial infection.^{25,26} Differentiation of these may not be possible.

222 Within our study, the most commonly encountered single lung pattern was structured interstitial. 223 However, these cases could be further subdivided into cases displaying a nodular pattern, comprising 224 scattered rounded hyperattenuating foci, and a reticulonodular pattern, where rounded foci and linear or 225 sickle-shaped hyperattenuating structures overlie to give a more complex overall pattern. In humans, a 226 faint, diffuse reticulonodular pattern is considered characteristic of miliary tuberculosis.²⁷ While nodular 227 and reticulonodular patterns are distinguishable on good quality radiographs in cats, the distinction is only 228 rarely made in veterinary imaging. On CT however, the difference is more easily appreciable. The 229 diagnostic and prognostic significance of the variable patterns in feline patients is currently unknown, but 230 certainly warrants further investigation, as a structured interstitial pattern is a common finding in many 231 feline lung pathologies (eg, pulmonary fibrosis, metastatic neoplasia, eosinophilic bronchopneumopathy 232 and a wide range of infectious pneumonias).

233 It is interesting to note that within our study population, two cats were found to have cavitations 234 within their lungs. While this feature is relatively common in both humans and dogs with tuberculosis²⁸⁻³⁰ it was not noted in any case in the previous radiographic study of cats,¹¹ and the only paper which describes 235 236 cavitating tubercles in cats was published in 1949.⁵ The lesions noted in the two cats in this study were 237 small (<1 cm) and were contained within regions of nodular or alveolar change. It is possible therefore that 238 they may not have been visible on radiographs, again highlighting the advantage of cross sectional imaging. 239 Alternatively, this may indeed reflect a rare occurrence in feline patients, which has occurred coincidentally 240 within our study population. In either case, it is an important characteristic to recognise, as cavitated lung masses are occasionally identified in feline patients with lung neoplasia,^{16,25} and the potential for 241 242 misdiagnosis exists in cats with mycobacteriosis which show this feature. In addition this should be 243 recognised because these cats likely pose an increased zoonotic risk compared with those showing the more 244 typical structured interstitial pattern as they may allow mycobacteria to gain access to the upper airways.

245 Thoracic lymphadenomegaly is a feature of numerous pulmonary and multisystemic conditions in 246 cats including, but not limited to, infiltrative and metastatic neoplasia, hypereosinophilic syndromes and 247 systemic mycosis/bacteriosis.³¹ As expected thoracic lymphadenomegaly was commonly noted within our 248 study population, but in contrast to the findings of the previous radiographic paper, mild and moderate 249 enlargement predominated over severe.¹¹ This may reflect the difficulty in recognising minor changes on 250 radiographs. It is also worth noting that even given the superior contrast resolution of CT, with mild 251 lymphadenomegaly, particularly in the perihilar region; changes were more easily appreciated in post-252 contrast studies when compared with pre-contrast. This suggests that there is value in performing post-253 contrast scans in all cases (unless there is a clinical contraindication), which was not standard practice 254 within our study population.

255 Mineralisation of thoracic lymph nodes and pulmonary parenchyma can result from chronic 256 inflammation associated with mycobacterial infection;^{9,13,32} it is also seen in cases of both primary and 257 metastatic pulmonary neoplasia, and chronic airways disease.²⁵ In either case, it is a finding which most 258 likely relates to the disease process that is present. In contrast, mild aortic root calcification, such as that 259 seen in one case (a seven year old cat) in our study is, in our experience, an occasional finding in middle 260 aged to older cats, and not necessarily related to clinical disease.

261 While peripheral and abdominal lymphadenomegaly were relatively common within the study 262 population, it is possible that the number of cases with mild or moderate lymphadenomegaly in the head 263 and neck was artificially low. This is because all cases recorded as having normal peripheral lymph nodes 264 on physical examination either did not undergo imaging of the head and neck, or were scanned conscious 265 within a VetMouseTrapTM device. The protocol for these scans involved a short scan time (in order to 266 minimise movement) resulting in a relatively large slice thickness and consequently a reduced longitudinal 267 resolution. This can compromise assessment of small structures so it is possible that mild or moderate 268 abnormalities of the head and neck, such as lymphadenomegaly, may have been overlooked. Other 269 abdominal changes such as hepatomegaly, splenomegaly, renal and pancreatic changes were noted 270 relatively infrequently and were mild or moderate in extent, consistent with previous reports.^{9,11,12,33}

Two distinct manifestations of skeletal disease were noted within our population. The lesions characterised by cortical lysis likely represent sites of primary bacterial inoculation and as such are

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273 consistent in location with 'fight and bite' injuries; whereas periarticular abnormalities are consistent with 274 an infectious polyarthritis resulting from haematogenous dissemination of bacteria. It is interesting to note 275 that the appendicular skeletal lesions in this study were clinically evident, and affected regions were 276 intentionally included in the imaging studies. Clinically silent skeletal lesions may have been overlooked as 277 the limbs were excluded from the majority of studies. The only studies in which the limbs were included in 278 full were those performed using the VetMouseTrapTM device and it is possible that subtle or focal regions 279 of bone lysis or proliferation may not have been recognised due to the lower resolution of these studies.

Cutaneous lesions were noted infrequently in this study. While this may initially seem surprising, given that cutaneous lesions represent a common presentation of mycobacteriosis,^{3,9} it reflects the fact that CT imaging is more likely to be employed in cases presenting with systemic disease, or used as a staging tool in cats with clinically evident focal skin lesions without requirement for imaging of the lesions themselves. The presence of intranasal changes in one cat is interesting, as these are indicative of a mycobacterial rhinitis, a manifestation of respiratory mycobacteriosis which may not be commonly recognised.

287 There are a number of limitations to this study. The most significant of these is that although 288 mycobacteriosis was confirmed in each case, histopathology on all involved tissues was not typically 289 performed; therefore, it is not possible to confirm that all changes seen were due to mycobacterial infection. 290 Due to the inherent difficulties in confirmation of mycobacterial infection, the time lapse between 291 acquisition of CT images and definitive confirmation of diagnosis was very variable; it extended to four 292 years and nine months in one case (though a lapse of one to four months was more typical). In all cases 293 however, at the time of imaging, the combination of clinical and pathological findings gave sufficient 294 confidence in the diagnosis to allow treatment to be instigated; imaging was used to stage the cases and so 295 guide the intensity and duration of treatment. The evolution of changes over time in association with 296 treatment has not been described, and will be interesting to explore in the future. Finally, given the 297 retrospective nature of this study there are inconsistencies between cases with respect to factors such as 298 regions imaged and use of contrast medium. This leads to a bias in our results, and may underestimate 299 subclinical disease, particularly affecting the peripheral structures. As mentioned, the limited resolution of 300 smaller structures on VetMouseTrap[™] scans contributes to this. However, the advantages of this technique

- 301 for disease screening, particularly in clinically compromised patients should not be ignored, and as faster
- 302 scanners become more commonplace many of the resolution difficulties will be eliminated.
- 303

304 Conclusions

305	As expected, the majority of CT changes noted in this study represent multisystemic disease,
306	typically with combinations of pulmonary infiltration, lymphadenomegaly and organomegaly. These
307	changes are strongly suggestive of infiltrative disease, differentials for which can include neoplasia (such as
308	lymphoma or mast cell disease), chronic inflammation/infectious processes (mycobacteriosis, feline
309	infectious peritonitis or systemic mycosis), hypereosinophilic syndrome and amyloidosis. ²⁴ While no
310	abnormality has been recognised that is specific for mycobacteriosis, it is important that the potential for
311	mycobacterial infection is considered when these types of changes are identified in feline patients,
312	especially if they have non-specific clinical signs. In addition, when managing patients with a diagnosis of
313	mycobacteriosis, the potential for widespread clinical and sub-clinical abnormalities must be considered
314	and investigated in full.
315	

316

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- 319
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- 321

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- 326
- 327 References

Journal of Feline Medicine and Surgery

328	1. Snider WR. Tuberculosis in canine and feline populations. Review of the literature. <i>Am Rev Respir</i>
329	Dis 1971; 104: 877-87.
330	2. Snider WR, Cohen D, Reif JS, et al. Tuberculosis in canine and feline populations. Study of high
331	risk populations in Pennsylvania, 1966-1968. Am Rev Respir Dis 1971; 104: 866-76.
332	3. Gunn-Moore DA, McFarland SE, Brewer JI, et al. Mycobacterial disease in cats in Great Britain:
333	I. Culture results, geographical distribution and clinical presentation of 339 cases. J Feline Med Surg 2011;
334	13: 934-44.
335	4. Malik R, Smits B, Reppas G, et al. Ulcerated and nonulcerated nontuberculous cutaneous
336	mycobacterial granulomas in cats and dogs. Vet Dermatol 2013; 24: 146-53.
337	5. Jennings AR. The distribution of tuberculous lesions in the dog and cat, with reference to the
338	pathogenesis. Vet Rec 1949; 6127: 380-84.
339	6. Huitema H and van Vloten J. Murine tuberculosis in a cat. Antonie Van Leeuwenhoek 1960; 26:
340	235-40.
341	7. Gunn-Moore D. Mycobacterial infections in cats and dogs. In: Ettinger S and Feldman E (eds).
342	Textbook of Veterinary Internal Medicine. 7th ed. Philadelphia: Saunders, 2010: 875-81.
343	8. de la Rua-Domenech R. Human Mycobacterium bovis infection in the United Kingdom:
344	Incidence, risks, control measures and review of the zoonotic aspects of bovine tuberculosis. Tuberculosis
345	(Edinb) 2006; 86: 77-109.
346	9. Gunn-Moore D, Dean R and Shaw S. Mycobacterial infections in cats and dogs. <i>In Pract</i> 2010;
347	32: 444-52.
348	10. Cima G. Cat transmits TB to humans in UK. <i>J Am Vet Med Assoc</i> 2014; 244: 1116.
349	11. Bennett AD, Lalor S, Schwarz T, et al. Radiographic findings in cats with mycobacterial
350	infections. J Feline Med Surg 2011; 13: 718-24.
351	12. Baral RM, Metcalfe SS, Krockenberger MB, et al. Disseminated Mycobacterium avium infection
352	in young cats: overrepresentation of Abyssinian cats. J Feline Med Surg 2006; 8: 23-44.
352 353	 in young cats: overrepresentation of Abyssinian cats. <i>J Feline Med Surg</i> 2006; 8: 23-44. 13. Foster SF, Martin P, Davis W, et al. Chronic pneumonia caused by <i>Mycobacterium</i>

- 354 *thermoresistibile* in a cat. J Small Anim Pract 1999; 40: 433-8.
- 355 14. Paltrinieri S. Tuberculosis in the dog and cat. *Nuova Vet* 1930; 6: 7.

356	15.	Prather AB, Berry CR and Thrall DE. Use of Radiography in Combination with Computed
357	Tomogr	raphy for the Assessment of Noncardiac Thoracic Disease in the Dog and Cat. Vet Radiol
358	Ultraso	und 2005; 46: 114-21.
359	16.	Henninger W. Use of computed tomography in the diseased feline thorax. J Small Anim Pract
360	2003; 44	4: 56-64.
361	17.	Daniel R, Evans H, Rolfe S, et al. Outbreak of tuberculosis caused by Mycobacterium bovis in
362	golden (Guernsey goats in Great Britain. Vet Rec 2009; 165: 335-42.
363	18.	Rhodes SG, Gruffydd-Jones T, Gunn-Moore D, et al. Interferon-gamma test for feline
364	tubercul	losis. Vet Rec 2008; 162: 453-5.
365	19.	Rhodes SG, Gruffydd-Jones T, Gunn-Moore D, et al. Adaptation of IFN-gamma ELISA and
366	ELISPC	OT tests for feline tuberculosis. Vet Immunol Immunopathol 2008; 124: 379-84.
367	20.	Oliveira CR, Ranallo FN, Pijanowski GJ, et al. The Vetmousetrap [™] : A Device for Computed
368	Tomogr	raphic Imaging of the Thorax of Awake Cats. Vet Radiol Ultrasound 201011; 52: 41-52.
369	21.	Greene CE and Gunn-Moore D. Mycobacterial Infections. In: Greene CE (ed). Infectious Diseases
370	of the D	og and Cat. 4th ed. Philadelphia: Elsevier Health Services, 2012: 495-510.
371	22.	Gunn-Moore DA, Jenkins PA and Lucke VM. Feline tuberculosis: a literature review and
372	discussi	on of 19 cases caused by an unusual mycobacterial variant. Vet Rec 1996; 138: 53-8.
373	23.	Barry M, Taylor J and Woods JP. Disseminated Mycobacterium avium infection in a cat. Can Vet
374	J 2002;	43: 369-71.
375	24.	Gow AG. What is your diagnosis? Mycobacterial infection. J Small Anim Pract 2006; 47: 484-5.
376	25.	Thrall DE. The Canine and Feline Lung. In: Thrall DE (ed). Textbook of Veterinary Diagnostic
377	Imaging	g. 6th ed. St. Louis: Elsevier Saunders, 2013: 608-31.
378	26.	Schwarz T and Johnson V. Lungs and Bronchi. In: Schwarz T and Saunders J (eds). Veterinary
379	Comput	ed Tomography. Chichester: Wiley-Blackwell, 2011: 261-78.

- 380 27. Velez MG and Velez VJ, Jr. Diffuse reticulonodular infiltrates. *Cleve Clin J Med* 2012; 79: 16-7.
- 381 28. Krysl J, Korzeniewska-Kosela M, Muller NL, et al. Radiologic features of pulmonary
- tuberculosis: an assessment of 188 cases. *Can Assoc Radiol J* 1994; 45: 101-7.
- 383 29. Gadkowski LB and Stout JE. Cavitary pulmonary disease. *Clin Microbiol Rev* 2008; 21: 305-33.

384 30. Olsson SE. On tuberculosis in the dog; a study with special reference to x-ray diagnosis. Cornell 385 Vet 1957; 47: 193-219.

386 31. Baines E. The mediastinum. In: Schwarz T and Johnson V (eds). BSAVA manual of canine and

387 feline thoracic imaging. Gloucester: BSAVA, 2008: 177-99.

- 388 32. Hix JW, Jones TC and Karlson AG. Avian tubercle bacillus infection in the cat. J Am Vet Med
- 389 Assoc 1961; 138: 641-7.
- 390 33. Knippel A, Hetzel U and Baumgartner W. Disseminated Mycobacterium avium-intracellulare
- 391 Infection in a Persian cat. J Vet Med 2004; 51: 464-6.

JUIN ad 2004; 51.



Figure 1. CT appearance of lung parenchyma in three cats with mycobacteriosis, at the level of the accessory lung lobe. (a) Alveolar pattern affecting multiple lung lobes. (b) Diffuse structured interstitial pattern comprising multiple relatively well defined nodules (arrows). (c) Diffuse structured interstitial pattern comprising mixed nodular and linear structures, characteristic of a reticulonodular pattern 91x84mm (300 x 300 DPI)



Figure 2. Thoracic lymphadenopathy in two cats with mycobacteriosis. (a) Transverse thoracic CT image at the level of the third sternebra showing an enlarged cranial mediastinal lymph node (arrowheads) containing a focus of mineralisation (arrow). (b) Post-contrast transverse CT image at the level of the cardiac base showing bilaterally enlarged tracheobronchial lymph nodes (LN) surrounding the trachea (T). The use of contrast medium allows differentiation from the cardiac and vascular structures 134x84mm (300 x 300 DPI)



Figure 3. Less common thoracic abnormalities in three cats with mycobacteriosis. (a(i)) Transverse thoracic CT image at the level of the caudal mainstem bronchi showing a partially cavitated nodule (arrow). (a(ii)) Transverse thoracic CT image at the level of the accessory lung lobe showing more extensive parenchymal cavitation (b) Transverse thoracic CT image at the level of the third thoracic vertebra showing a region of alveolar pattern containing mineralised foci (*)

98x84mm (300 x 300 DPI)



Figure 4. Extrathoracic lymphadenomegaly in two cats with mycobacteriosis. (a) Dorsal plane CT reconstruction of the abdomen at the level of the descending colon in a cat with a partially mineralised colic lymph node (arrow). (b) Dorsal plane CT reconstruction of the ventral neck in a cat with marked bilateral medial retropharyngeal lymphadenomegaly (LN) 128x84mm (300 x 300 DPI)



Figure 5. Bony lesions in four cats with mycobacteriosis. (a) Transverse CT image at the level of the canine teeth, demonstrating focal lysis (arrowhead) and moderate, irregular periosteal reaction (arrow). These is a soft tissue mass lesion associated with the bony change (*). (b) Transverse CT image at the level of the proximal radius and ulna which shows focal cortical lysis of the ulna (arrows) and marked smooth periosteal reaction. The adjacent radius is normal. (c) Sagittal plane CT reconstruction of the radius and ulna in a cat with a pathological fracture secondary to a tuberculous lesion. Lytic foci are visible in the distal radius (arrows), and there is a moderate smooth periosteal reaction (arrowhead). The adjacent ulna is normal. (d) Transverse CT image at the level of the humeral condyle showing marked periarticular new bone formation in a cat with a mycobacterial polyarthritis

152x140mm (300 x 300 DPI)

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1	Computed Tomographic Findings in Cats with Mycobacterial Infection
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26 **Objectives**

27 The objective of this study was to describe the imaging findings in computed tomography

28 (CT) associated with confirmed mycobacterial infection in cats.

29 Methods

30 CT images from 20 cats with confirmed mycobacterial disease were retrospectively 31 reviewed. Five cats underwent conscious full-body CT in a VetMouseTrapTM device. All 32 other cats had thoracic CT performed under general anaesthesia, with the addition of CT 33 investigation of the head/neck, abdomen and limbs in some cases.

34 **Results**

35 Mycobacterial infection was seen most frequently in adult (mean age 7.4 years; range 36 0.6-14 years) neutered male cats (11/20). The most common infections were 37 Mycobacterium microti (6/20) and Mycobacterium bovis (6/20). CT abnormalities were 38 most commonly seen in the thorax, consisting of bronchial (9/20), alveolar (8/20), ground 39 glass (6/20) or structured interstitial (15/20) lung patterns, which were often mixed. 40 Tracheobronchial, sternal and cranial mediastinal lymphadenomegaly were common 41 (16/20). Other abnormalities included abdominal (8/13) or peripheral (10/18)42 lymphadenomegaly, hepatosplenomegaly (7/13), mixed osteolytic/osteoproliferative 43 skeletal lesions (7/20), and cutaneous or subcutaneous soft tissue masses/nodules (4/20).

44 Conclusions and relevance

45 CT of feline mycobacteriosis shows a wide range of abnormalities often involving46 multiple organ systems and mimicking many other feline diseases. Mycobacteriosis

- 47 should be considered in the differential diagnosis of thoracic, abdominal and skeletal
- 48 disorders in cats.
- 49
- 50
- 51

52 Introduction

Feline mycobacteriosis is a worldwide veterinary health concern, and although definitive data on case numbers worldwide are lacking, mycobacterial infections in cats have been recognised with increasing frequency in the UK, as well as being seen in many other countries. Mycobacterial disease in domestic cats can result from infection by one of a number of species. The most commonly identified mycobacteria include *Mycobacterium microti* and *Mycobacterium bovis*, which are primary pathogens and members of the tuberculous complex group of mycobacteria.¹⁻³ Non-tuberculous mycobacterial species are less commonly identified within clinically affected cats.⁴

60 Clinical presentation of mycobacterial infection in cats is variable, and is dependent primarily on the species of mycobacteria involved and, importantly, the route of infection.^{2,5-7} Historically, alimentary 61 62 lesions resulting from ingestion of milk from cows infected with M. bovis were most common; however 63 with overall reduction of tuberculosis in the national bovine herd since the early 1900's and widespread 64 pasteurisation of milk this is no longer the case.⁸ Single or multiple cutaneous lesions with or without 65 lymph node involvement, and characteristically affecting the so-called 'fight and bite sites' (such as the 66 head and limbs), now represent the most common presentation of mycobacterial infection in cats: they typically result from infection acquired from prey species.^{3,9} Infection acquired through inhalation or 67 68 ingestion, resulting in respiratory or alimentary disease, is seen less frequently. The clinical presentation of 69 these forms of disease, and of disseminated disease resulting from haematogenous spread of infection, can 70 include non-specific signs such as weight loss, anorexia, coughing, anaemia, vomiting/diarrhoea, 71 hepatosplenomegaly, generalised lymphadenopathy and pyrexia.⁷

72 Definitive diagnosis of mycobacterial disease in cats can present significant problems, in part due 73 to difficulties in sample handling, and limitations in the available laboratory diagnostic techniques. As 74 such, mycobacterial infections are likely underdiagnosed within the domestic cat population. In addition to 75 significant morbidity resulting from primary infection, subclinical infection and recurrence of infection 76 following treatment are common.⁷ Since significant and potentially fatal multisystemic disease can result 77 from infection with mycobacterial species, and since there are potential zoonotic risks associated with all 78 members of the tuberculosis complex,^{7,10} identification and correct handling of potential cases is of the 79 upmost importance.

80 Previous publications detailing the diagnostic imaging findings in cats with confirmed 81 mycobacterial infection are limited to a single retrospective case series looking at survey radiographic 82 changes involving 33 cats,¹¹ and a number of isolated case reports describing the radiographic features of feline mycobacteriosis.¹²⁻¹⁴ Computed tomography is increasingly available to the veterinary community, 83 84 and it offers significant advantages over survey radiography by eliminating superimposition of anatomy, 85 having superior contrast resolution and being able to clarify intrathoracic lesions where radiographic findings are negative or non-specific.^{15,16} In addition, the decreased scan times which are achievable with 86 87 modern multi-detector scanners make CT a valuable tool in investigation of multisystemic disease in 88 clinically compromised patients. The CT features of mycobacterial disease in cats have not been described 89 previously. The aim of this paper was to review CT images from a large number of cats with confirmed 90 mycobacteriosis and to describe the range of abnormalities that can occur.

91

92 Materials and Methods

This study comprises a descriptive, retrospective case series. CT studies carried out between August 2009 and January 2015, of cats with confirmed mycobacterial infection were submitted to one of the authors (DGM). Inclusion criteria consisted of: (i) confirmation of mycobacterial infection and (ii) a CT study of diagnostic quality. To confirm mycobacterial involvement, aspirated and/or biopsy samples of affected tissue had been stained with Ziehl-Neelson (ZN) and found to have changes indicative of mycobacteriosis.¹ Where possible, tissue culture,¹⁷ interferon-gamma release assay, or PCR testing had been used to identify which mycobacterial species was involved.^{4,18,19}

Pseudonymised CT studies of the confirmed mycobacterial cases were examined without knowledge of specific clinical information by a third year diagnostic imaging resident who was however informed about the topic of the study (AM). To prevent bias by the assumption of disease, CT studies covering the thorax and other body parts from an additional ten cats with confirmed non-mycobacterial diseases were included and also pseudonymised. Images were evaluated using dedicated DICOM viewer software (Osirix, Geneva, Switzerland, version 5.8.5-64bit) on a computer workstation (Apple Mac Pro, Apple, USA) with a calibrated LCD flat screen monitor (Apple Cinemax Display, 30 inch, Apple, USA). 107 During the course of image evaluation, multi-planar reconstructions, maximum and minimum intensity108 projections and variable windowing settings were used according to the preferences of the viewer.

109 CT studies were reviewed for the following diagnostic criteria: bronchial thickening; alveolar 110 pattern; ground glass opacity or structured interstitial lung change; evidence of pleural or pericardial 111 effusion, or pleural/mediastinal thickening; thoracic, abdominal or peripheral lymphadenomegaly, or lymph 112 node mineralisation; abdominal organomegaly, peritoneal effusion, other abdominal organ-associated 113 lesions; osteolysis or osteoproliferative changes; cutaneous/subcutaneous/oral/nasal lesions; vascular and 114 dystrophic soft tissue calcification. The extent of any abnormality was characterised as focal, multifocal, or 115 diffuse. The degree of each change was graded as absent/normal, mild, moderate or severe.

116

117 **Results**

118 Twenty cats met the inclusion criteria. After all image interpretive data had been collected the 119 additional ten non-mycobacterial cat studies were identified and their data were excluded from further 120 analysis. The most common infections were M. microti and M. bovis, confirmed in 6/20 cases each. A non-121 specified *M. tuberculosis* complex species was described in one case and in the remaining 7/20 cases the 122 species involved was not known. Eleven of the 20 cats were neutered males and 9/20 were neutered 123 females. The study group comprised 7/20 Domestic Short Hair, 4/20 Siamese, 2/20 Domestic Long Hair 124 and 1/20 of each of the following; Persian, Birman, Norwegian Forest Cat, Burmilla, British Short Hair, 125 Bengal and Maine Coon cats. The age of one cat was not known. For the remaining cats the mean age was 126 7.4 ± 3.8 years (range 0.6-14 years).

Five of the 20 cats underwent conscious full-body CT in a specific containing device (VetMouseTrap[™], University of Illinois at Urbana-Champaign, Urbana, IL).²⁰ The remaining 15 cats were scanned under general anaesthesia, with images of the following body regions obtained: thorax only (2), head/neck and thorax (3), thorax and abdomen (4), head/neck, thorax and abdomen (2), head/neck, thorax, abdomen and single forelimb (2), head, thorax, bilateral tarsi/elbows (1), thorax and single hind limb (1). Intravenous contrast medium (iopamidol or iohexol, 600-700mg I/kg) was administered to 12/20

133 cats, and post-contrast images of some or all body parts were obtained. Use of contrast medium depended

on the findings in the pre-contrast images, the clinical condition of the cat, and the preferences of theattending radiologist and primary clinician in each case.

136 Within the evaluated imaging studies, thoracic abnormalities were noted in 19/20 cases. Diffuse 137 bronchial thickening was present in 9/20 cats; being mild in eight cases and moderate in one. Eight cats 138 showed a focal alveolar pattern; mild in two cases, moderate in three cases and severe in three cases (Figure 139 1(a)). Diffuse or patchy ground glass opacity was noted in 6/20 cats; mild in three cases, moderate in two 140 cases and severe in one case. The most common pulmonary parenchymal change was a diffuse structured 141 interstitial pattern, which was present in 15/20 cats, being either nodular (7/15) or reticulonodular (8/15) in 142 nature; mild in six cases, moderate in five cases and severe in four cases (Figure 1(b,c)). Thoracic CT 143 images of 14/20 cats were considered to show a mixed pulmonary pattern, with a single pattern present in 144 4/20 cases. The appearance of the pulmonary parenchyma was normal in 2/20 cats, though one of these had 145 a thoracic lymphadenopathy despite normal lungs. Of the 20 cats, 16 had sternal, cranial mediastinal and/or 146 tracheobronchial lymphadenomegaly (Figure 2). Moderate lymphadenomegaly affecting the sternal or 147 tracheobronchial nodes was most common. One cat had moderate mineralisation of an enlarged cranial 148 mediastinal lymph node.

None of the cats had any evidence of pleural or pericardial effusion. One cat showed mild, diffuse pleural thickening. One cat showed mild mineralisation of the aortic root. Two cats had regions of cavitation within the lungs, associated in both cases with focal or multifocal nodular or alveolar changes (Figure 3(a)). Three cats had scattered foci of mineralisation within the lungs, again associated with other focal parenchymal changes (Figure 3(b)).

154 Thirteen of the 20 cats had imaging studies that included the abdomen. Abdominal 155 lymphadenomegaly was present in 8/13 cases and was typically generalised. The lymph nodes affected 156 could not always be individually identified, but included those of the celiac and cranial mesenteric centres, 157 which variably comprised the hepatic, splenic, gastric, pancreaticoduodenal, jejunal and colic nodes. 158 Lymphadenomegaly was mild in two cats, moderate in four cats and severe in two cats. In one cat with a 159 generalised moderate abdominal lymphadenomegaly, mild mineralisation of a mesenteric lymph node was 160 present (Figure 4(a)). Mild hepatomegaly was present in 3/13 cats and moderate hepatomegaly in 1/13. 161 Mild splenomegaly was present in 6/13 cats and moderate splenomegaly in 1/13. Two cats with

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splenomegaly (one mild and one moderate) were also noted to show heterogeneity within the splenic parenchyma following contrast medium administration. Additional abdominal organ changes were noted in 3/13 cats; one had a moderately enlarged pancreas, one had multiple nodules within both kidneys, and one an irregular outline to the left kidney. Peritoneal effusion was not noted in any cat.

166 The appearance of the peripheral lymph nodes was assessed in 18/20 cats. The two cats not 167 included in this assessment had CT studies of the thorax only, without inclusion of any extra-thoracic 168 lymph node group. In 10/18 cases peripheral lymphadenomegaly was present, mild in 3/18, moderate in 169 2/18 and severe in 5/18. In 8/10 cats with lymphadenomegaly, the most significant enlargement was noted 170 in the mandibular and medial retropharyngeal nodes (Figure 4(b)); however, multifocal 171 lymphadenomegaly, involving the superficial cervical (prescapular), axillary, inguinal and/or popliteal 172 nodes was variably present. In the other 2/10 cats, the head and neck were not imaged, but enlargement of 173 the superficial cervical and popliteal lymph nodes was noted respectively. Five of the eight cats in which 174 peripheral lymphadenomegaly was not noted underwent conscious CT in the VetMouseTrapTM device and 175 three underwent CT studies which did not include the head and neck.

176 Focal osteolytic lesions were present in 7/20 cats (although it was not possible to assess the entire 177 skeleton in 15 cats as they did not have full body CT examinations); changes were mild in four cases and 178 moderate in three cases. These lesions affected the nasal bridge in three cats and the limbs in the remaining 179 four, and were predominantly characterised by regions of cortical lysis (5/7) or erosive lesions at joint 180 surfaces (2/7) (Figure 5). An associated pathological long bone fracture was present in one case. In all but 181 one of these cases osteoproliferative changes, either periosteal reaction or periarticular osteophytosis, were 182 noted in the same location as the osteolytic change. The osteoproliferation was mild in three cases and 183 severe in three cases; however, the degree of proliferative change did not necessarily correlate with the 184 degree of lytic change in each case.

Cutaneous or subcutaneous lesions were only infrequently present within the studies evaluated. Focal mass lesions over the nasal bridge were noted in 2/20 cats, graded moderate in one and severe in one. One other cat had a small amount of fluid accumulation and soft tissue thickening in the dorsal nasal chambers. Each of these lesions was adjacent to bony abnormalities. A focal, but extensive, mass lesion was noted along the ventral head and neck of one cat. One cat was found to have multiple, widely distributed, subcutaneous nodules. Diffuse extra-thoracic dystrophic soft tissue mineralisation was notnoted in any cat.

192

193 **Discussion**

Mycobacteriosis in cats is known to be a highly variable disease, and should always be considered as a possible differential diagnosis in cases which present with multisystemic signs. Mycobacterial disease is likely under-recognised, primarily due to a lack of awareness of the full spectrum of changes which can be associated with it.

Mycobacterial infection is most commonly seen in adult, neutered male cats consistent with the results of our study.⁹ Domestic Short Hair cats predominate in our study, but to a lesser degree than noted in the previous radiographic case series (36% vs 87%).¹¹ The reason for this is unknown, but may reflect a higher proportion of pedigree cats within a referral population, which are therefore more likely to undergo advanced imaging.

CT abnormalities of the thorax were commonly noted, being present in all but one cat. However, multisystemic abnormalities were also common, with changes affecting more than one anatomical region in all but five cases. Of these five, three had abnormalities detected on clinical examination which were not appreciable on the CT images. In cats, systemic mycobacterial infection is most commonly caused by *M*. *bovis* or *M microti*,^{3,9,21-24} and our results are consistent with this.

Previous reports of radiographic findings in cats with mycobacterial disease described a mild 208 209 predominance of a mixed lung pattern (ie, a combination of bronchial, alveolar and/or interstitial 210 changes),^{19,11-13,22-24} but distinct alveolar, bronchial or interstitial patterns in isolation were also identified.¹¹ 211 Interestingly, where cases in our study displayed mixed lung patterns, bronchial thickening and ground 212 glass opacity were most commonly graded as mild, whereas alveolar pattern and structured interstitial 213 patterns were more likely to be moderate or severe. This is interesting for two reasons. Firstly, the mild 214 bronchial and unstructured interstitial patterns are of a degree that comparable changes may not be easily 215 visible radiographically, or may be attributed to expiratory or underexposed radiographs, or 216 superimposition of other structures. In a radiographic study it is therefore possible that only a more 217 significant overlying alveolar or nodular pattern may be recognised, leading to classification as a single

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218 lung pattern. As superimposition effects are eliminated by CT, it becomes easier to identify these mild 219 changes in addition to the more marked ones. Secondly, a mild bronchial or unstructured interstitial pattern 220 may be indicative of concurrent conditions, such as low-grade allergic airway disease, rather than being 221 directly related to an active mycobacterial infection.^{25,26} Differentiation of these may not be possible.

222 Within our study, the most commonly encountered single lung pattern was structured interstitial. 223 However, these cases could be further subdivided into cases displaying a nodular pattern, comprising 224 scattered rounded hyperattenuating foci, and a reticulonodular pattern, where rounded foci and linear or 225 sickle-shaped hyperattenuating structures overlie to give a more complex overall pattern. In humans, a 226 faint, diffuse reticulonodular pattern is considered characteristic of miliary tuberculosis.²⁷ While nodular 227 and reticulonodular patterns are distinguishable on good quality radiographs in cats, the distinction is only 228 rarely made in veterinary imaging. On CT however, the difference is more easily appreciable. The 229 diagnostic and prognostic significance of the variable patterns in feline patients is currently unknown, but 230 certainly warrants further investigation, as a structured interstitial pattern is a common finding in many 231 feline lung pathologies (eg, pulmonary fibrosis, metastatic neoplasia, eosinophilic bronchopneumopathy 232 and a wide range of infectious pneumonias).

233 It is interesting to note that within our study population, two cats were found to have cavitations 234 within their lungs. While this feature is relatively common in both humans and dogs with tuberculosis²⁸⁻³⁰ it was not noted in any case in the previous radiographic study of cats,¹¹ and the only paper which describes 235 236 cavitating tubercles in cats was published in 1949.⁵ The lesions noted in the two cats in this study were 237 small (<1 cm) and were contained within regions of nodular or alveolar change. It is possible therefore that 238 they may not have been visible on radiographs, again highlighting the advantage of cross sectional imaging. 239 Alternatively, this may indeed reflect a rare occurrence in feline patients, which has occurred coincidentally 240 within our study population. In either case, it is an important characteristic to recognise, as cavitated lung masses are occasionally identified in feline patients with lung neoplasia,^{16,25} and the potential for 241 242 misdiagnosis exists in cats with mycobacteriosis which show this feature. In addition this should be 243 recognised because these cats likely pose an increased zoonotic risk compared with those showing the more 244 typical structured interstitial pattern as they may allow mycobacteria to gain access to the upper airways.

245 Thoracic lymphadenomegaly is a feature of numerous pulmonary and multisystemic conditions in 246 cats including, but not limited to, infiltrative and metastatic neoplasia, hypereosinophilic syndromes and 247 systemic mycosis/bacteriosis.³¹ As expected thoracic lymphadenomegaly was commonly noted within our 248 study population, but in contrast to the findings of the previous radiographic paper, mild and moderate 249 enlargement predominated over severe.¹¹ This may reflect the difficulty in recognising minor changes on 250 radiographs. It is also worth noting that even given the superior contrast resolution of CT, with mild 251 lymphadenomegaly, particularly in the perihilar region; changes were more easily appreciated in post-252 contrast studies when compared with pre-contrast. This suggests that there is value in performing post-253 contrast scans in all cases (unless there is a clinical contraindication), which was not standard practice 254 within our study population.

255 Mineralisation of thoracic lymph nodes and pulmonary parenchyma can result from chronic 256 inflammation associated with mycobacterial infection;^{9,13,32} it is also seen in cases of both primary and 257 metastatic pulmonary neoplasia, and chronic airways disease.²⁵ In either case, it is a finding which most 258 likely relates to the disease process that is present. In contrast, mild aortic root calcification, such as that 259 seen in one case (a seven year old cat) in our study is, in our experience, an occasional finding in middle 260 aged to older cats, and not necessarily related to clinical disease.

261 While peripheral and abdominal lymphadenomegaly were relatively common within the study 262 population, it is possible that the number of cases with mild or moderate lymphadenomegaly in the head 263 and neck was artificially low. This is because all cases recorded as having normal peripheral lymph nodes 264 on physical examination either did not undergo imaging of the head and neck, or were scanned conscious 265 within a VetMouseTrapTM device. The protocol for these scans involved a short scan time (in order to 266 minimise movement) resulting in a relatively large slice thickness and consequently a reduced longitudinal 267 resolution. This can compromise assessment of small structures so it is possible that mild or moderate 268 abnormalities of the head and neck, such as lymphadenomegaly, may have been overlooked. Other 269 abdominal changes such as hepatomegaly, splenomegaly, renal and pancreatic changes were noted 270 relatively infrequently and were mild or moderate in extent, consistent with previous reports.^{9,11,12,33}

Two distinct manifestations of skeletal disease were noted within our population. The lesions characterised by cortical lysis likely represent sites of primary bacterial inoculation and as such are 273 consistent in location with 'fight and bite' injuries; whereas periarticular abnormalities are consistent with 274 an infectious polyarthritis resulting from haematogenous dissemination of bacteria. It is interesting to note 275 that the appendicular skeletal lesions in this study were clinically evident, and affected regions were 276 intentionally included in the imaging studies. Clinically silent skeletal lesions may have been overlooked as 277 the limbs were excluded from the majority of studies. The only studies in which the limbs were included in 278 full were those performed using the VetMouseTrapTM device and it is possible that subtle or focal regions 279 of bone lysis or proliferation may not have been recognised due to the lower resolution of these studies.

Cutaneous lesions were noted infrequently in this study. While this may initially seem surprising, given that cutaneous lesions represent a common presentation of mycobacteriosis,^{3,9} it reflects the fact that CT imaging is more likely to be employed in cases presenting with systemic disease, or used as a staging tool in cats with clinically evident focal skin lesions without requirement for imaging of the lesions themselves. The presence of intranasal changes in one cat is interesting, as these are indicative of a mycobacterial rhinitis, a manifestation of respiratory mycobacteriosis which may not be commonly recognised.

287 There are a number of limitations to this study. The most significant of these is that although 288 mycobacteriosis was confirmed in each case, histopathology on all involved tissues was not typically 289 performed; therefore, it is not possible to confirm that all changes seen were due to mycobacterial infection. 290 Due to the inherent difficulties in confirmation of mycobacterial infection, the time lapse between 291 acquisition of CT images and definitive confirmation of diagnosis was very variable; it extended to four 292 years and nine months in one case (though a lapse of one to four months was more typical). In all cases 293 however, at the time of imaging, the combination of clinical and pathological findings gave sufficient 294 confidence in the diagnosis to allow treatment to be instigated; imaging was used to stage the cases and so 295 guide the intensity and duration of treatment. The evolution of changes over time in association with 296 treatment has not been described, and will be interesting to explore in the future. Finally, given the 297 retrospective nature of this study there are inconsistencies between cases with respect to factors such as 298 regions imaged and use of contrast medium. This leads to a bias in our results, and may underestimate 299 subclinical disease, particularly affecting the peripheral structures. As mentioned, the limited resolution of 300 smaller structures on VetMouseTrapTM scans contributes to this. However, the advantages of this technique

- 301 for disease screening, particularly in clinically compromised patients should not be ignored, and as faster
- 302 scanners become more commonplace many of the resolution difficulties will be eliminated.
- 303

304 Conclusions

305 As expected, the majority of CT changes noted in this study represent multisystemic disease, 306 typically with combinations of pulmonary infiltration, lymphadenomegaly and organomegaly. These 307 changes are strongly suggestive of infiltrative disease, differentials for which can include neoplasia (such as 308 lymphoma or mast cell disease), chronic inflammation/infectious processes (mycobacteriosis, feline 309 infectious peritonitis or systemic mycosis), hypereosinophilic syndrome and amyloidosis.²⁴ While no 310 abnormality has been recognised that is specific for mycobacteriosis, it is important that the potential for 311 mycobacterial infection is considered when these types of changes are identified in feline patients, 312 especially if they have non-specific clinical signs. In addition, when managing patients with a diagnosis of 313 mycobacteriosis, the potential for widespread clinical and sub-clinical abnormalities must be considered 314 and investigated in full. 315 316 317 This research received no specific grant from any funding agency in the public, commercial, or

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- 326
- 327 References

328	1.	Snider WR. Tuberculosis in canine and feline populations. Review of the literature. Am Rev Respir
329	Dis 1971; 104: 877-87.	
330	2.	Snider WR, Cohen D, Reif JS, et al. Tuberculosis in canine and feline populations. Study of high
331	risk po	pulations in Pennsylvania, 1966-1968. Am Rev Respir Dis 1971; 104: 866-76.
332	3.	Gunn-Moore DA, McFarland SE, Brewer JI, et al. Mycobacterial disease in cats in Great Britain:
333	I. Cult	ure results, geographical distribution and clinical presentation of 339 cases. J Feline Med Surg 2011;
334	13: 934-44.	
335	4.	Malik R, Smits B, Reppas G, et al. Ulcerated and nonulcerated nontuberculous cutaneous
336	mycob	acterial granulomas in cats and dogs. Vet Dermatol 2013; 24: 146-53.
337	5.	Jennings AR. The distribution of tuberculous lesions in the dog and cat, with reference to the
338	pathogenesis. Vet Rec 1949; 61: 380-84.	
339	6.	Huitema H and van Vloten J. Murine tuberculosis in a cat. Antonie Van Leeuwenhoek 1960; 26:
340	235-40).
340 341	235-4(7.). Gunn-Moore D. Mycobacterial infections in cats and dogs. In: Ettinger S and Feldman E (eds).
340 341 342	235-40 7. Textbo). Gunn-Moore D. Mycobacterial infections in cats and dogs. In: Ettinger S and Feldman E (eds). bok of Veterinary Internal Medicine. 7th ed. Philadelphia: Saunders, 2010: 875-81.
340 341 342 343	235-4(7. Textbo 8.). Gunn-Moore D. Mycobacterial infections in cats and dogs. In: Ettinger S and Feldman E (eds). bok of Veterinary Internal Medicine. 7th ed. Philadelphia: Saunders, 2010: 875-81. de la Rua-Domenech R. Human Mycobacterium bovis infection in the United Kingdom:
 340 341 342 343 344 	235-4(7. Textbo 8. Incide	 Gunn-Moore D. Mycobacterial infections in cats and dogs. In: Ettinger S and Feldman E (eds). bok of Veterinary Internal Medicine. 7th ed. Philadelphia: Saunders, 2010: 875-81. de la Rua-Domenech R. Human Mycobacterium bovis infection in the United Kingdom: nce, risks, control measures and review of the zoonotic aspects of bovine tuberculosis. <i>Tuberculosis</i>
340 341 342 343 344 345	235-4(7. Textbo 8. Incide: <i>(Edinb</i>	 Gunn-Moore D. Mycobacterial infections in cats and dogs. In: Ettinger S and Feldman E (eds). bok of Veterinary Internal Medicine. 7th ed. Philadelphia: Saunders, 2010: 875-81. de la Rua-Domenech R. Human Mycobacterium bovis infection in the United Kingdom: ince, risks, control measures and review of the zoonotic aspects of bovine tuberculosis. <i>Tuberculosis</i> (2006; 86: 77-109.)
 340 341 342 343 344 345 346 	235-4(7. Textbo 8. Incide: (<i>Edinb</i> 9.	 Gunn-Moore D. Mycobacterial infections in cats and dogs. In: Ettinger S and Feldman E (eds). bok of Veterinary Internal Medicine. 7th ed. Philadelphia: Saunders, 2010: 875-81. de la Rua-Domenech R. Human Mycobacterium bovis infection in the United Kingdom: nce, risks, control measures and review of the zoonotic aspects of bovine tuberculosis. <i>Tuberculosis</i> c) 2006; 86: 77-109. Gunn-Moore D, Dean R and Shaw S. Mycobacterial infections in cats and dogs. <i>In Pract</i> 2010;
340 341 342 343 344 345 346 347	235-4(7. Textbo 8. Incide (<i>Edinb</i> 9. 32: 44	 Gunn-Moore D. Mycobacterial infections in cats and dogs. In: Ettinger S and Feldman E (eds). bok of Veterinary Internal Medicine. 7th ed. Philadelphia: Saunders, 2010: 875-81. de la Rua-Domenech R. Human Mycobacterium bovis infection in the United Kingdom: nce, risks, control measures and review of the zoonotic aspects of bovine tuberculosis. <i>Tuberculosis</i> 2006; 86: 77-109. Gunn-Moore D, Dean R and Shaw S. Mycobacterial infections in cats and dogs. <i>In Pract</i> 2010; 4-52.
 340 341 342 343 344 345 346 347 348 	235-4(7. Textbo 8. Incide: (<i>Edinb</i> 9. 32: 44	 Gunn-Moore D. Mycobacterial infections in cats and dogs. In: Ettinger S and Feldman E (eds). book of Veterinary Internal Medicine. 7th ed. Philadelphia: Saunders, 2010: 875-81. de la Rua-Domenech R. Human Mycobacterium bovis infection in the United Kingdom: nce, risks, control measures and review of the zoonotic aspects of bovine tuberculosis. <i>Tuberculosis</i> c) 2006; 86: 77-109. Gunn-Moore D, Dean R and Shaw S. Mycobacterial infections in cats and dogs. <i>In Pract</i> 2010; 4-52. Cima G. Cat transmits TB to humans in UK. <i>J Am Vet Med Assoc</i> 2014; 244: 1116.
 340 341 342 343 344 345 346 347 348 349 	 235-40 7. Textbox 8. Incider (<i>Edinb</i> 9. 32: 44 10. 11. 	 Gunn-Moore D. Mycobacterial infections in cats and dogs. In: Ettinger S and Feldman E (eds). bok of Veterinary Internal Medicine. 7th ed. Philadelphia: Saunders, 2010: 875-81. de la Rua-Domenech R. Human Mycobacterium bovis infection in the United Kingdom: nce, risks, control measures and review of the zoonotic aspects of bovine tuberculosis. <i>Tuberculosis</i> 2006; 86: 77-109. Gunn-Moore D, Dean R and Shaw S. Mycobacterial infections in cats and dogs. <i>In Pract</i> 2010; 4-52. Cima G. Cat transmits TB to humans in UK. <i>J Am Vet Med Assoc</i> 2014; 244: 1116. Bennett AD, Lalor S, Schwarz T, et al. Radiographic findings in cats with mycobacterial
 340 341 342 343 344 345 346 347 348 349 350 	235-4(7. Textbo 8. Incide: (<i>Edinb</i> 9. 32: 44 10. 11. infecti	 Gunn-Moore D. Mycobacterial infections in cats and dogs. In: Ettinger S and Feldman E (eds). book of Veterinary Internal Medicine. 7th ed. Philadelphia: Saunders, 2010: 875-81. de la Rua-Domenech R. Human Mycobacterium bovis infection in the United Kingdom: nce, risks, control measures and review of the zoonotic aspects of bovine tuberculosis. <i>Tuberculosis</i> c) 2006; 86: 77-109. Gunn-Moore D, Dean R and Shaw S. Mycobacterial infections in cats and dogs. <i>In Pract</i> 2010; 4-52. Cima G. Cat transmits TB to humans in UK. <i>J Am Vet Med Assoc</i> 2014; 244: 1116. Bennett AD, Lalor S, Schwarz T, et al. Radiographic findings in cats with mycobacterial ons. <i>J Feline Med Surg</i> 2011; 13: 718-24.

- in young cats: overrepresentation of Abyssinian cats. *J Feline Med Surg* 2006; 8: 23-44.
- 353 13. Foster SF, Martin P, Davis W, et al. Chronic pneumonia caused by Mycobacterium
- 354 *thermoresistibile* in a cat. J Small Anim Pract 1999; 40: 433-8.
- 355 14. Paltrinieri S. Tuberculosis in the dog and cat. *Nuova Vet* 1930; 6: 7.

- 356 15. Prather AB, Berry CR and Thrall DE. Use of Radiography in Combination with Computed 357 Tomography for the Assessment of Noncardiac Thoracic Disease in the Dog and Cat. Vet Radiol 358 Ultrasound 2005; 46: 114-21.
- 359 16. Henninger W. Use of computed tomography in the diseased feline thorax. J Small Anim Pract 360 2003; 44: 56-64.
- 361 17. Daniel R, Evans H, Rolfe S, et al. Outbreak of tuberculosis caused by Mycobacterium bovis in
- 362 golden Guernsey goats in Great Britain. Vet Rec 2009; 165: 335-42.
- 363 18. Rhodes SG, Gruffydd-Jones T, Gunn-Moore D, et al. Interferon-gamma test for feline 364 tuberculosis. Vet Rec 2008; 162: 453-5.
- 365 19. Rhodes SG, Gruffydd-Jones T, Gunn-Moore D, et al. Adaptation of IFN-gamma ELISA and
- 366 ELISPOT tests for feline tuberculosis. Vet Immunol Immunopathol 2008; 124: 379-84.
- 367 20. Oliveira CR, Ranallo FN, Pijanowski GJ, et al. The VetmousetrapTM: A Device for Computed
- 368 Tomographic Imaging of the Thorax of Awake Cats. Vet Radiol Ultrasound 2011; 52: 41-52.
- 369 21. Greene CE and Gunn-Moore D. Mycobacterial Infections. In: Greene CE (ed). Infectious Diseases
- 370 of the Dog and Cat. 4th ed. Philadelphia: Elsevier Health Services, 2012: 495-510.
- 371 22. Gunn-Moore DA, Jenkins PA and Lucke VM. Feline tuberculosis: a literature review and 372 discussion of 19 cases caused by an unusual mycobacterial variant. Vet Rec 1996; 138: 53-8.
- 373 23. Barry M, Taylor J and Woods JP. Disseminated Mycobacterium avium infection in a cat. Can Vet 374 J 2002; 43: 369-71.
- 375 Gow AG. What is your diagnosis? Mycobacterial infection. J Small Anim Pract 2006; 47: 484-5. 24.
- 376 25. Thrall DE. The Canine and Feline Lung. In: Thrall DE (ed). Textbook of Veterinary Diagnostic
- 377 Imaging. 6th ed. St. Louis: Elsevier Saunders, 2013: 608-31.
- 378 26. Schwarz T and Johnson V. Lungs and Bronchi. In: Schwarz T and Saunders J (eds). Veterinary 379
- Computed Tomography. Chichester: Wiley-Blackwell, 2011: 261-78.
- 380 27. Velez MG and Velez VJ, Jr. Diffuse reticulonodular infiltrates. Cleve Clin J Med 2012; 79: 16-7.
- 381 28. Krysl J, Korzeniewska-Kosela M, Muller NL, et al. Radiologic features of pulmonary
- 382 tuberculosis: an assessment of 188 cases. Can Assoc Radiol J 1994; 45: 101-7.
- 383 29. Gadkowski LB and Stout JE. Cavitary pulmonary disease. Clin Microbiol Rev 2008; 21: 305-33.

384 30. Olsson SE. On tuberculosis in the dog; a study with special reference to x-ray diagnosis. Cornell 385 Vet 1957; 47: 193-219. 386 Baines E. The mediastinum. In: Schwarz T and Johnson V (eds). BSAVA manual of canine and 31. 387 feline thoracic imaging. Gloucester: BSAVA, 2008: 177-99. 388 32. Hix JW, Jones TC and Karlson AG. Avian tubercle bacillus infection in the cat. J Am Vet Med 389 Assoc 1961; 138: 641-7. 390 33. Knippel A, Hetzel U and Baumgartner W. Disseminated Mycobacterium avium-intracellulare 391 Infection in a Persian cat. J Vet Med 2004; 51: 464-6. 392 393 Figure 1. CT appearance of lung parenchyma in three cats with mycobacteriosis, at the level of the 394 accessory lung lobe. (a) Alveolar pattern affecting multiple lung lobes. (b) Diffuse structured 395 interstitial pattern comprising multiple relatively well defined nodules (arrows). (c) Diffuse 396 structured interstitial pattern comprising mixed nodular and linear structures, characteristic of a 397 reticulonodular pattern 398 399 Figure 2. Thoracic lymphadenopathy in two cats with mycobacteriosis. (a) Transverse thoracic CT 400 image at the level of the third sternebra showing an enlarged cranial mediastinal lymph node 401 (arrowheads) containing a focus of mineralisation (arrow). (b) Post-contrast transverse CT image at 402 the level of the cardiac base showing bilaterally enlarged tracheobronchial lymph nodes (LN) 403 surrounding the trachea (T). The use of contrast medium allows differentiation from the cardiac and 404 vascular structures 405 406 Figure 3. Less common thoracic abnormalities in three cats with mycobacteriosis. (a(i)) Transverse 407 thoracic CT image at the level of the caudal mainstem bronchi showing a partially cavitated nodule

408 (arrow). (a(ii)) Transverse thoracic CT image at the level of the accessory lung lobe showing more

409 extensive parenchymal cavitation (b) Transverse thoracic CT image at the level of the third thoracic

410 vertebra showing a region of alveolar pattern containing mineralised foci (*)

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Figure 4. Extrathoracic lymphadenomegaly in two cats with mycobacteriosis. (a) Dorsal plane CT reconstruction of the abdomen at the level of the descending colon in a cat with a partially mineralised colic lymph node (arrow). (b) Dorsal plane CT reconstruction of the ventral neck in a cat with marked bilateral medial retropharyngeal lymphadenomegaly (LN)

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417 Figure 5. Bony lesions in four cats with mycobacteriosis. (a) Transverse CT image at the level of the 418 canine teeth, demonstrating focal lysis (arrowhead) and moderate, irregular periosteal reaction 419 (arrow). These is a soft tissue mass lesion associated with the bony change (*). (b) Transverse CT 420 image at the level of the proximal radius and ulna which shows focal cortical lysis of the ulna 421 (arrows) and marked smooth periosteal reaction. The adjacent radius is normal. (c) Sagittal plane 422 CT reconstruction of the radius and ulna in a cat with a pathological fracture secondary to a 423 tuberculous lesion. Lytic foci are visible in the distal radius (arrows), and there is a moderate smooth 424 periosteal reaction (arrowhead). The adjacent ulna is normal. (d) Transverse CT image at the level of 425 the humeral condyle showing marked periarticular new bone formation in a cat with a mycobacterial 426 polyarthritis

PL. QL