Ventral cervical subcutaneous Aspergillus species fungal granuloma in a cat

Courtney Bartels1,2, Alejandro Alvarez-Sanchez1, Bharadhwaj Ranganathan1,2, T William O’Neill3,4 and Katy L Townsend1

Abstract

Case summary An 8-year-old domestic longhair cat was evaluated for a right ventral subcutaneous cervical mass. Serial bloodwork and contrast-enhanced cranial and thoracic CT initially lacked ethmoturbinate lysis and showed a progressive, vascularized, right ventral cervical mass involving local lymph nodes. The mass was removed surgically on two occasions. Histopathology and fungal culture were diagnostic for a recurring sclerosing fungal granuloma and pyogranulomatous and eosinophilic lymphadenitis, consistent with Aspergillus species. The cat was treated with oral itraconazole; however, owing to the owner’s non-compliance in administering the medication and disease progression, the cat was humanely euthanized 3 years after diagnosis.

Relevance and novel information The development of a cervical subcutaneous fungal granuloma of Aspergillus species in a domestic longhair cat before obvious maxillary, orbital or ethmoturbinate lysis on initial diagnostics is rare and suggests an early onset of lymphatic or hematogenous spread from a suspected nidus of infection within the sinonasal cavity.

Keywords: Aspergillus; granuloma; fungal; subcutaneous

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Introduction

Aspergillus species are a ubiquitous opportunistic fungus found in the environment. It is considered to be an emerging infectious disease that affects mammals worldwide. Aspergillosis can be classified as invasive, semi-invasive or non-invasive, and it is frequently caused by Aspergillus fumigatus, followed by Aspergillus niger, Aspergillus felis and Aspergillus fischeri. Common sites of infection are the sinus cavity, orbit and thorax, with focal infections reported in the ear canal, cornea, gastrointestinal tract and urinary bladder. Aspergillus species infection typically occurs through inhalation. Rarely, spread can occur via lymphatic or hematogenous routes. This report documents the development of a cervical subcutaneous Aspergillus species granuloma in a cat prior to obvious maxillary or ethmoturbinate lysis, suggesting early-onset lymphatic or hematogenous spread.

Case description

An 8-year-old male neutered domestic longhair cat weighing 6.54 kg with an unknown vaccination status was referred for a 3-week history of a right ventral cervical subcutaneous fungal granuloma of Aspergillus species. The cat was treated with oral itraconazole; however, owing to the owner’s non-compliance in administering the medication and disease progression, the cat was humanely euthanized 3 years after diagnosis.

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cervical mass. Primary care complete blood count (CBC) and serum biochemistry revealed hyperglobulinemia (8.7 g/dl; reference interval [RI] 2.9–4.7) as the only clinically significant finding.

On presentation, physical examination revealed a large, subcutaneous, mobile mass (30 mm × 25 mm) in the right ventral cervical region. CBC showed no clinically significant findings. Serum biochemistry revealed mild hypoalbuminemia (2.4 g/dl; RI 2.6–4.0) and hyperglobulinemia (7.4 g/dl; RI 2.9–4.7). Feline leukemia virus antigen and feline immunodeficiency antibody SNAP tests (IDEXX Laboratory) were negative. The patient was sedated with dexmedetomidine 0.005 mg/kg (Dexdomitor; Orion Pharma) and butorphanol 0.3 mg/kg intravenously (Torbutrol; Bristol-Myers). Cranial, cervical and thoracic CT scans were acquired (Aquilion 64 slice, 36-5 [Toshiba] and Tsurumichuo 4-chrome [Tsurumi-Ku]) before and after intravenous injection of iodinated contrast medium (1.83 ml/kg iodixanol [Isovue 300; General Electric Healthcare]). Images revealed lymphadenopathy of the right medial retroparotygeal lymph node (RMRLN; 39 mm × 18 mm × 19 mm) and right mandibular lymph node (RMLN; 12 mm × 14 mm × 7 mm), bilateral non-destructive rhinitis with soft tissue attenuating and non-contrast-enhancing material in the frontal sinus and cervical mass, left mandibular lymphadenopathy, decreased lung lobe sizes and tortuosity of the cranial mediastinal blood vessels (Figure 1). Cytology of the mass revealed mixed inflammation with intralesional fungi. Marginal surgical excision of the recurring fungal granuloma and RSLCN were performed and submitted for histopathological analysis and fungal culture. Histopathology yielded similar results: pyogranulomatous lymphadenitis with intralesional fungal hyphae (Figure 3). Fungal culture revealed *Aspergillus* species characterized by branching fungal hyphae with roughly parallel walls and Grocott's methenamine silver (GMS) and periodic acid–Schiff (PAS) confirmed recurrence. Speciation was not performed. The patient was discharged with gabapentin (10 mg/kg PO q8h [Neurontin; Pfizer]), robenacoxib (1 mg/kg PO [Onsior; Elanco]) once after discharge and discharged with buprenorphine (0.01 mg/kg oral transmucosally [OTM] q8h [Buprenex; Indivior]) and liquid suspension itraconazole (5.05 mg/kg PO q24h [Sporanox; Janssen Pharmaceutical]). However, owing to difficulty administering at home, the patient only received one dose of itraconazole.

The patient re-presented 17 months postoperatively for suspected recurrence of the cervical fungal granuloma. Physical examination revealed a firm, lobulated right subcutaneous ventral cervical mass (50 mm × 30 mm) near the scar of the previous excision site. The right superficial cervical lymph node (RSCLN) was firm and enlarged on palpation (15 mm in length). CBC, serum biochemistry and CT were repeated and compared with previous visits. Anemia (hematocrit 27%; RI 30–45%), lymphopenia (679/µl; RI 1500–7000), hypoalbuminemia (2.5 g/dl; RI 2.6–4.0) and hyperglobulinemia (9.2 g/dl; 2.9–4.7) were identified. Imaging revealed an enlarged right cervical mass, left mandibular lymphadenopathy, decreased lung lobe sizes and tortuosity of the cranial mediastinal blood vessels (Figure 1). Cytology of the mass revealed mixed inflammation with intralesional fungi. Marginal surgical excision of the recurring fungal granuloma and RSLCN were performed and submitted for histopathological analysis and fungal culture.

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The cat re-presented 11 months after the second surgery for acute respiratory distress. Liquid suspension itraconazole (5.05 mg/kg PO q24h) had not been administered as instructed. Physical examination revealed a firm, lobulated, subcutaneous mass (120 mm × 40 mm) in the left ventral cervical region, immediately caudal to the mandible. A gallop rhythm with a grade 4/6 left parasternal systolic murmur was auscultated. CBC revealed a non-regenerative anemia (hematocrit 15.2%; RI 30–45%); reticulocytes 6800/µl [RI 0–50,000]). Doppler blood pressure was 112 mmHg. An echocardiogram showed left ventricle thickening with mitral regurgitation and turbulent outflow, consistent with hypertrophic cardiomyopathy without signs of heart failure. Contrast-enhanced cranial and thoracic CT was repeated, showing a large mass (56 mm × 38 mm × 34 mm) extending from the right retrobulbar region to the nasopharynx, caudally to the thoracic inlet, causing rightward deviation of the hyoid apparatus, larynx, trachea, thyroid...
Figure 1 (a) Sagittal, soft tissue window, contrast-enhanced CT of the cervical region 5 months after initial presentation. Note the severely enlarged, progressive, well-vascularized right ventral cervical mass (asterisk) with involvement of the right medial retropharyngeal lymph node (arrow). (b) Transverse, soft tissue window, contrast enhanced CT of nasal cavity. Note the minimal soft tissue opacity in the nasal cavity and no bony lysis (arrow). (c) Transverse, soft tissue window, contrast-enhanced CT of the cranial cervical region. Note the mass (asterisk) and the right medial retropharyngeal lymph node (arrow). (d) Sagittal, soft tissue window CT of the cervical region with iodinated contrast 22 months after initial presentation. Note the mass (asterisk). (e) Transverse, soft tissue window CT of the cervical region with iodinated contrast. Note the enlarged right ventral cervical mass (asterisk). (f) Sagittal, soft tissue window CT of the cervical region with iodinated contrast 33 months after initial presentation. Note the retrobulbar mass effect, as well as the cervical mass dorsal and ventral to the trachea (asterisks). (g) Transverse, soft tissue window CT of the cervical region with iodinated contrast. Noted the mass effect displacing the right globe (arrow)
glands and cervical esophagus, as well as displacement of the right globe (Figure 1). The left lateral mandibular lymph node was enlarged (12 mm × 7 mm). The left common carotid artery coursed through the mass, and there was sternal and cranial mediastinal lymphadenopathy, in addition to a pulmonary nodule within the left caudal lung lobe. There was marked loss of endoturbinate structures of the right and left nasal cavity, with soft tissue-attenuating material within the lateroventral aspect of the right nasal cavity. Given the extent of the fungal granuloma and recurring nature, surgical resection was not advised, and the cat was humanely euthanized without necropsy.

Discussion
This report is of an Aspergillus species granuloma forming in the subcutaneous ventral cervical region of a cat suspected to be secondary to hematogenous or lymphatic dissemination from the nasal cavity. Initial diagnosis of Aspergillus species was achieved based on surgical excision and histopathology. Recurrence was diagnosed based on repeat surgical excision, histopathology, histochemical stains (GMS and PAS; Figures 2 and 3) and fungal culture. Histopathology revealed severe pyogranulomatous and eosinophilic lymphadenitis with numerous intralesional, branching and fungal hyphae with parallel walls, findings consistent with previously described aspergillosis cases.2–4

Serial chemistries showed persistent hyperglobulinemia, the most frequently reported laboratory abnormality associated with aspergillosis.9–12 Similar changes can be found with neoplastic processes; therefore, SPE was pursued. These results were consistent with a polyclonal gammopathy, signifying an underlying inflammatory or infectious disease rather than neoplasm. Cytology and histopathology were used to identify the presence of fungal hyphae.2 These results, in conjunction with results from the SPE, confirmed Aspergillus species infection.

Cranial CT showed evidence of nasal disease in this patient at presentation. Given the presence of soft tissue-attenuating material within the nasal cavity without associated bone lysis, it is possible that the cat initially had rhinosinusitis or allergic rhinitis. Similar ventral distribution of fluid and soft tissue material was seen in a series of cats with rhinitis. Sinonasal neoplasia and fungal infections are also frequently associated with paranasal bone lysis at the time of diagnosis, yet there was no evidence of bone lysis in this cat at presentation.14,15 Additionally, hyperattenuating soft tissues were commonly seen in a study of cats with sinonasal fungal disease yet similar enhancement following contrast injection was not seen in this cat.15 Had bone and/or nasal turbinate destruction been noticed at the time of the initial CT, rhinoscopy, biopsy and culture sample acquisition would have been recommended for further evaluation.

Without bone lysis, we suspect the fungus metastasized to the cervical integument and lymph nodes via hematogenous or lymphatic routes from the nasal cavity. There are reports of rare focal infections within the urinary bladder and gastrointestinal tract. However, these dissemination routes could have resulted from an ascending infection through the urinary system or oral ingestion, respectively. It appears that this route of dissemination in cats may occur more frequently than initially thought in the absence of ethmoturbinate lysis. In addition, the feline report by Cormack and colleagues had primary dissemination to adjacent tissue planes through the naso-orbital structures and respiratory system. These routes account for the most common dissemination paths and are classically associated with upper respiratory signs and/or epistaxis.2 Orbital bone lysis was not documented in our cat until 22 months after presentation. Albeit rare, we caution readers that dissemination to the integument and locoregional lymph nodes can occur without apparent sinonasal symptomatology.
Figure 3  (a) Hematoxylin and eosin (HE) stain (×40) of the granuloma. Note the frequent eosinophils and degenerate neutrophils surrounded by concentric rings of epithelioid macrophages, lymphocytes and a fibrous capsule (yellow arrow). (b) HE stain (×400) of the edge of granuloma with rare fungal hyphae. Note that eosinophils and necrotic debris dominate this area with rare eosinophilic fungal hyphae. The right side of the image shows the internal edge of the granuloma, with foamy macrophages and scattered small lymphocytes (red arrow). (c) Grocott’s methenamine silver stain (×40) of the same granuloma as (a). Note the abundance of fungal hyphae within the center of the granuloma (green arrow). (d) Grocott’s methenamine silver stain (×400) of the same granuloma as (b). Note the fungal morphology, which consists of fungal hyphae of 3–6 µm width with roughly parallel walls, regular branching and variable angles ranging from approximately 45º to 90º (blue arrow). (e) Periodic acid–Schiff stain (×400) of the same granuloma as (b) and (d). Note the fungal hyphae, in addition to the previously described features (black arrow). (f) Ventral cervical fungal granuloma. On removal, the granuloma had coalescing pyogranulomas, eosinophilic infiltrates, supported by mature fibrocollagenous tissue, necrotic tissue, hemorrhage and collagen degeneration, as well as numerous branching hyphae with inflammatory infiltrates. A diagnosis of severe sclerosing pyogranulomatous and eosinophilic lymphadenitis associated with intralesional fungi was determined.
Although no prospective case-controlled studies exist on the treatment of feline aspergillosis cases, the prognosis for non-invasive aspergillosis is relatively good when appropriately treated, whereas the prognosis of the invasive form is relatively poor, despite treatment. Aspergillus species infections are typically treated with topical or systemic antifungals. Correct identification via molecular testing and antifungal susceptibility is instrumental in guiding therapy and limit azole cross-resistance. Molecular testing and early fungus speciation could have been used to guide antifungal selection; however, neither were performed. Most cryptic species of Aspergillus are resistant to itraconazole, yet this antifungal was selected as it is usually effective against Aspergillus fumigatus.

Conclusions
Feline aspergillosis is an emerging disease, with a growing number of cases and novel tissue involvement. Species identification is crucial to tailor treatment protocols. This report documents a case of suspected lymphatic or pulmonary dissemination of sinonasal aspergillosis to cervical lymph nodes and subcutaneous tissues in a cat years before suffering from parasanal bone destruction, warranting consideration as a differential.

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Ethical approval The work described in this manuscript involved the use of non-experimental (owned or unowned) animals. Established internationally recognized high standards ('best practice') of veterinary clinical care for the individual patient were always followed and/or this work involved the use of cadavers. Ethical approval from a committee was therefore not specifically required for publication in JFMS Open Reports. Although not required, where ethical approval was still obtained, it is stated in the manuscript.

Informed consent Informed consent (verbal or written) was obtained from the owner or legal custodian of the animal(s) described in this work (experimental or non-experimental animals, including cadavers) for all procedure(s) undertaken (prospective or retrospective studies). No animals or people are identifiable within this publication, and therefore additional informed consent for publication was not required.

ORCID iD Courtney Bartels https://orcid.org/0000-0002-3145-3278
Katy L. Townsend https://orcid.org/0000-0001-6405-0793

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