Subcutaneous Atypical Mycobacteriosis in Captive Tiger Quolls
(Dasyurus maculatus)


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Abstract. From July 1989 to October 1998, 9/37 (24%) adult captive tiger quolls (Dasyurus maculatus) were diagnosed with atypical mycobacterial infection involving the subcutis and skin. Females were more often affected than males (seven females, two males). Grossly, lesions presented as focal thickenings, plaques, and abscesses within the subcutis, often with fistulous tracts. The subcutis and skin overlying cervical and thoracic regions were the primary sites of infection. Cytology of subcutaneous impression smears from all nine affected tiger quolls revealed pyogranulomatous inflammation admixed with several acid-fast bacilli. Histologically, all tiger quolls had nodular to diffuse pyogranulomatous panniculitis and cellulitis. Small numbers of acid-fast bacilli were noted histologically in 7/9 (78%) animals. Skin cultures from seven tiger quolls were positive for one or more different Runyon group IV mycobacteria. The disease described in these tiger quolls is similar to subcutaneous atypical mycobacteriosis of humans and domestic animals.

Key words: Atypical mycobacteriosis; cellulitis; group IV mycobacteria; panniculitis; skin; tiger quolls.

The tiger quoll (Dasyurus maculatus) is a member of the family Dasyuridae, carnivorous marsupials indigenous to the Australian continent and adjacent islands. Because of the diminishing numbers of this species in the wild, a breeding program was established in 1984 at the US National Zoological Park’s Conservation and Research Center (CRC). In 1989, an adult female tiger quoll presented with a subcutaneous abscess from which a Runyon group IV mycobacterium was isolated. During the next 9 years, eight additional tiger quolls were diagnosed with similar atypical mycobacterial infections that involved the subcutis. Here, we describe the clinical and pathologic findings in tiger quolls afflicted with atypical mycobacterial infection of the subcutis and skin.

Materials and Methods

Study population

In 1984, a breeding program was established at the CRC (Front Royal, VA) based on two pairs of wild-caught tiger quolls. The majority of tiger quolls were maintained at the CRC, and a small number were kept on exhibit at the National Zoological Park (Washington, DC). Their diet consisted of Toronto Feline Diet (Milliken Wholesale Meat, Toronto, ON, Canada) supplemented with apple biscuits, live and dead mice (2–3 times/week), thawed fish (once weekly), and a multivitamin supplement. Tiger quolls were housed singly except when males were introduced into the females’ enclosures for breeding encounters. Animals were housed in large indoor enclosures, either $3.1 \times 6.2 \times 3.4$ m or $6.2 \times 3.4$ m, with access to outdoor cages of equal size. The flooring substrate was composed of pine bark, solite, and peat moss.

From 1989 to 1998, the collection had a total of 37 adult tiger quolls. During this 9-year period, skin biopsies and necropsies were preformed on nine tiger quolls with similar cutaneous disease. Table 1 contains information on gender, age, and lesion distribution for these nine animals. Six of the nine tiger quolls were born in captivity at the CRC, and three were from the original wild-caught group. The affected population included seven female and two male tiger quolls, ranging from 2.5 to 5.4 years of age at the time of diagnosis (mean = 4.2 years; median = 4.6 years).

Histopathology

Samples of skin were collected either as biopsies or during postmortem examination. Tissues were fixed in 10% neutral buffered formalin, embedded in paraffin, sectioned at 5.0 µm, and stained with hematoxylin and eosin (HE). For visualization of mycobacteria in tissue sections, Kinyoun stain was used according to a previously published protocol.

Microbiology

For mycobacterial culturing, samples of skin, lymph node, and thoracic abscesses were plated onto blood agar and Lowenstein-Jensen medium (Becton Dickinson, Cockeysville, MD) and incubated at 35 C and 7% CO₂ for up to 1 week. Mycobacterial isolates were identified by high-performance liquid chromatography and conventional methods (MRL Reference Laboratory, Cypress, CA).
Table 1. Gender, age, and distribution of lesions for nine tiger quolls with subcutaneous atypical mycobacteriosis.

<table>
<thead>
<tr>
<th>Animal No.</th>
<th>Gender</th>
<th>Age (years)*</th>
<th>Distribution of Lesion(s)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>F</td>
<td>3.75</td>
<td>Subcutis and skin, right/left lateral neck; axillary lymph node</td>
</tr>
<tr>
<td>2</td>
<td>F</td>
<td>2.8</td>
<td>Subcutis and skin, dorsal neck, dorsal/ventral thorax, dorsal lumbar, dorsal pelvis, lateral abdomen; intercostal muscles; costal pleura</td>
</tr>
<tr>
<td>3</td>
<td>F</td>
<td>5.4</td>
<td>Subcutis and skin, ventral neck, left lateral thorax, ventral abdomen</td>
</tr>
<tr>
<td>4</td>
<td>F</td>
<td>4.75</td>
<td>Subcutis and skin, right lateral/ventral thorax, right axilla</td>
</tr>
<tr>
<td>5</td>
<td>M</td>
<td>2.5</td>
<td>Subcutis, right axilla</td>
</tr>
<tr>
<td>6</td>
<td>F</td>
<td>5.2</td>
<td>Subcutis and skin, dorsal neck</td>
</tr>
<tr>
<td>7</td>
<td>F</td>
<td>4.3</td>
<td>Subcutis, ventral and right/left neck, right/left axilla; axillary, submandibular lymph nodes</td>
</tr>
<tr>
<td>8</td>
<td>M</td>
<td>4.6</td>
<td>Subcutis, dorsal neck, dorsal thorax</td>
</tr>
<tr>
<td>9</td>
<td>F</td>
<td>4.7</td>
<td>Subcutis and skin, right lateral neck</td>
</tr>
</tbody>
</table>

* At time of initial diagnosis.

Cytology

Imprints and smears of subcutaneous lesions were air dried and stained with Wright’s stain and modified Kinyoun acid-fast stain according to a previously published protocol.¹

Results

Clinical findings

The cutaneous lesions presented as single to multiple thickened areas of skin of various sizes that were sometimes hyperemic and partially alopecic (Fig. 1). The subcutis was multifocally thickened with gray-tan plaques that contained variable amounts of thick, moist to dry, tan-yellow purulent material (Fig. 2). Multifocally, the subcutaneous tissue was edematous and hemorrhagic. The subcutaneous lesions sometimes presented as small draining abscesses or fistulous tracts that communicated with the surface of the skin. The distribution of the lesions was primarily along the dorsal, lateral, and ventral aspects of the neck and thorax (Table 1). Lesions were occasionally evident in the axillary region and rarely in the lumbar, abdominal, or pelvic regions. Two tiger quolls had enlargement of regional lymph nodes (two had enlarged axillary and one had enlarged submandibular lymph nodes). One tiger quoll had suppurative tracts that extended from the dermis and subcutis deep into the underlying in-

Fig. 1. Skin; tiger quoll. Gross appearance of thickened skin with alopecia from a tiger quoll with subcutaneous atypical mycobacteriosis. Bar = 5.0 cm.

Fig. 2. Skin; tiger quoll. Cross section of skin and subcutis from a tiger quoll with subcutaneous atypical mycobacteriosis. Subcutis is edematous and hemorrhagic (arrows) and contains exudative plaques that extend into the dermis (arrowhead). Bar = 5.0 cm.
tercostal skeletal musculature and communicated with an abscess adhered to the costal pleura.

Two tiger quolls were euthanatized following chronic subcutaneous atypical mycobacteriosis that was unresponsive to treatment. One tiger quoll died during the skin biopsy procedure. One tiger quoll died from spread of subcutaneous mycobacterial infection into the thoracic cavity. The remaining five tiger quolls died or were euthanatized because of other conditions unrelated to the mycobacteriosis.

Cytology

Impression smears of subcutaneous abscesses and plaques were made for all nine tiger quolls. Serial cutaneous smears were examined during the course of the disease for several tiger quolls. In all cases, there was an inflammatory cell population characterized by neutrophils, histiocytes, and few epithelioid macrophages, Langhans giant cells, lymphocytes, and plasma cells. Several extracellular beaded to filamentous acid-fast bacilli (AFB) were noted cytologically in all nine tiger quolls.

Histopathologic findings

Histologic lesions in the subcutis were similar in all nine tiger quolls. The predominant lesion was nodular to diffuse pyogranulomatous panniculitis and cellulitis. The inflammatory infiltrate was characterized by histiocytes, neutrophils, epithelioid macrophages, lymphocytes, plasma cells, and few Langhans giant cells and mast cells combined with fibrous connective tissue (Fig. 3). Multifocally within the pyogranulomatous inflammation were regions of necrosis, mineralization, and hemorrhage of various sizes. The inflammatory infiltrate expanded and effaced normal subcutaneous adipose tissue. In six animals, the inflammation extended into the dermis, forming fistulous tracts. The overlying epidermis was ulcerated in four animals. One tiger quoll (No. 2) had an extension of the pyogranulomatous inflammation into the underlying intercostal skeletal musculature and costal pleura, and another tiger quoll had involvement of dermal sebaceous glands. In two tiger quolls (Nos. 1, 7), there were neutrophils, histiocytes, and few epithelioid macrophages within subcapsular and medullary sinusoids of regional axillary (two animals) and submandibular (one animal) lymph nodes. In 7/9 (78%) tiger quolls, few aggregates of extracellular beaded to filamentous AFB were evident within large, clear lipid vacuoles embedded within the pyogranulomatous inflammation (Fig. 4). In two tiger quolls (Nos. 1, 9), AFB were not demonstrated histologically but were demonstrated by cytology of abscess exudate, and atypical mycobacteria were cultured from respective skin samples.

Microbiologic findings

Seven tiger quolls were positive for group IV mycobacteria by aerobic culture. Table 2 contains a list of the species of mycobacterium isolated. Mycobacterium smegmatis, M. fortuitum, M. chitae, and a type IV (not identified) mycobacterium collectively were cultured from seven skin samples. One tiger quoll (No. 3) was positive by culture for both M. fortuitum and M. smegmatis. M. fortuitum was cultured from an axillary lymph node from one tiger quoll (No. 7), and M. smegmatis was cultured from a thoracic abscess.
from another tiger quoll (No. 2). Attempts to isolate group IV mycobacterium from two tiger quolls (Nos. 5, 8) were unsuccessful.

**Treatement**

Eight tiger quolls were treated empirically for atypical mycobacterial infection of the subcutis and skin. The duration of treatment varied from several weeks to >4 months. Antimicrobial agents used were amikacin (Fort Dodge Laboratories, Fort Dodge, IA), enrofloxacin (Bayer Corp., Shawnee Mission, KS), cephalixin (Teva Pharmaceuticals, Sellersville, PA), ciprofloxacin (Bayer Corp.), chloramphenicol (Sanofi, Overland Park, KS), rifabutin (Pharmacia and Upjohn Corp., Kalamazoo, MI), and rifampin (Hoechst Marion Roussel, Kansas City, MO) at appropriate dosages. In a majority of cases, affected areas were surgically debrided prior to antibiotic therapy. Some animals responded temporarily to therapy, but most were refractory to treatment. In addition, several tiger quolls had recurrence of atypical mycobacterial infection in new cutaneous locations. Only two tiger quolls (Nos. 6, 9) that responded to treatment had apparent resolution of skin lesions until their death; however, skin from previously affected areas from one of these tiger quolls (No. 9) was lost to follow-up examination.

**Discussion**

From 1989 to 1998, 24% of captive adult tiger quolls in the National Zoological Park collection were diagnosed with atypical mycobacterial infection that involved primarily the subcutis and overlying dermis. This condition was the most commonly diagnosed disease in the collection. The disease had a similar clinical presentation in all nine tiger quolls, with thickened areas of skin and subcutis primarily along the neck and thorax. Histologic lesions were similar in all tiger quolls and consisted of pyogranulomatous panniculitis and cellulitis. One or more different species of Runyon group IV mycobacteria were cultured from the skin of seven tiger quolls. The disease in these tiger quolls is similar to subcutaneous atypical mycobacteriosis (SAM) in other animals and humans.

In humans, cutaneous infection with rapidly growing (atypical) mycobacteria is usually an opportunistic disease, particularly in patients that are immunocompromised. It is caused by infection with one or more different Runyon group IV mycobacteria. *M. fortuitum, M. smegmatis, M. chelonei, M. thermoresistible, and M. phlei* have been associated with SAM in domestic felids. Infection in cats mainly resides within the subcutis and skin and rarely spreads systemically. The same appears to be true in tiger quolls; in only one tiger quoll (No. 2) did infection spread beyond the subcutis and skin and deep into the thoracic wall, and in another two tiger quolls (Nos. 1, 7) infection extended to regional lymph node(s).

In cats, lesions due to infection with atypical mycobacteria are usually within the dorsal lumbar or ventral abdominal and inguinal regions. Typically, the clinical presentation is firm, plaquelike, warm, and sometimes painful swellings within the subcutis and skin. Draining fistulous tracts frequently are associated with the masses. The distribution of the skin infection in the tiger quolls was different from that in domestic cats in that it was concentrated primarily in the dorsolateral cervical and thoracic regions. Only one tiger quoll had lesions along the ventral abdomen. However, the clinical presentation and gross lesions of infection in tiger quolls were similar to that seen in domestic cats.

**Histologically**, SAM in cats manifests as nodular to diffuse pyogranulomatous panniculitis and dermatitis. The inflammation can extend into the superficial dermis and subjacent skeletal muscle. Usually, within the pyogranulomatous inflammation there are smooth-bordered extracellular lipid vacuoles that contain fine filamentous to beaded extracellular AFB. The histologic lesions in the tiger quolls of this study were identical to those described in cats diagnosed with SAM. AFB were not observed histologically in two animals, but all tiger quolls were positive for AFB by cytology. It appears that cytology is a more sensitive method than histology for detecting AFB in tiger quolls with SAM.

An apparent association exists between cutaneous trauma and SAM in domestic cats. In one study, all...
cats infected with atypical mycobacteria had previously sustained trauma to the skin via bite wounds or road accidents. Similar cutaneous injuries have been described in dogs with atypical mycobacterial infections of the skin. A delay of months to years can occur between the initial skin injury and clinical presentation. Tiger quolls reportedly have protracted bacteremia that sometimes involves the infection of bite wounds by males into the females’ necks. Approximately 3 months after copulation, females dig nests in the substrate, which they frequent until parturition. Because group IV mycobacteria are saprophytes found typically as soil and water contaminants, the substrate used in the tiger quoll enclosures was suspected as a source of infection. Open skin wounds acquired during mating may have become infected via contaminated substrate. However, repeated attempts to culture atypical mycobacteria from the substrate were unsuccessful. Also, not all cases could be easily described by this scenario. Long intervals from the time of bite wounds to clinical presentation were observed in tiger quolls with SAM; some of these intervals were >2 years. One female and both male tiger quolls had no recorded history of cutaneous injury; however, unnoticed skin trauma cannot be discounted in these cases.

Marsupials are more susceptible to mycobacterial infections than are eutherian mammals. Most case reports of mycobacteriosis in marsupials are from captive populations. The prevalence and susceptibility of wild brushtail possums (Trichosurus vulpecula) to Mycobacterium bovis infection is well documented, but the prevalence of atypical mycobacteriosis in wild marsupials is currently unknown. The role of effective cell-mediated immunity (CMI) in the clearance of mycobacteria is well known. Lymphocyte proliferative assays revealed that primary systemic infections with Mycobacterium avium complex were associated with lowered cellular immune reactivity in Matschie’s tree kangaroos (Dendrolagus matschiei) when compared with eutherian mammals. Using the same lymphocyte proliferative assays, similar depressed cellular immune reactivity was noted in tiger quolls (unpublished data), possibly making them more susceptible to developing mycobacterial infections. Some cats infected with atypical mycobacteria have difficulty eliciting an effective CMI response, and several of these cats are concurrently infected with feline immunodeficiency virus. The roles of a possible retrovirus, genetic factors (i.e., abnormal breeding coefficient), or physiological stress as causes for possible depressed CMI in these tiger quolls are currently unknown.

Antibiotic and surgical treatment of cats infected subcutaneously with atypical mycobacteria is usually unsuccessful. Surgical resection of mycobacterial skin lesions usually results in wound dehiscence and possible occurrence of new lesions. However, there are a few reported cases of remission in cats with or without treatment. Most (75%) tiger quolls treated for SAM failed to respond to any type of antimicrobial or surgical therapy. Two of eight tiger quolls treated for atypical mycobacteriosis had apparent clinical resolution of infection.

Tiger quolls are particularly susceptible to atypical mycobacterial infections, which manifest primarily as pyogranulomatous panniculitis. Diagnosis is best made by histopathology and cytology combined with mycobacterial culture of skin lesions. The substrate likely served as a source of infection, and mycobacteria may well have entered the skin through cutaneous wounds acquired during mating interactions. The delay in onset of clinical signs can sometimes be very long, and not all tiger quolls had historic evidence of cutaneous trauma. Atypical mycobacterial infection is an important differential diagnosis for pyogranulomatous skin disease in captive tiger quolls.

Acknowledgements

We thank D. Fischer, A. Brathbauer, and V. Bonshock (NZP, Smithsonian) for laboratory assistance, Dr. Phillips (George Washington University) for lymphocyte proliferative assay results, R.-A. V. Ferris (Armed Forces Institute of Pathology) for photography, and Dr. Collins, Dr. Citino, and the tiger quoll keeper staff at CRC. The project was supported by FONZ SR. Fellowship 98-3545-A.

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