Suppurative granulomatous sinorhinitis associated with *Nocardia* spp. infection in a cat

Ataru NAKANISHI1)*, Tadahisa MASHITA1,2), Kyoko AKIYAMA1), Wakana NAKANISHI1), Takashi MORI3,4), Masaki YANO4), Tetsuo ASAI2), Rui KANO5), Syunsuke SHIMAMURA2) and Jun YASUDA2)

1)Maizuru Animal Medical Center, 625–0037 Maizuru, Japan. e-mail: atarutare@gmail.com
2)The United Graduate School of Veterinary Science, Gifu University, 501–1193 Gifu, Japan
3)Laboratory of Veterinary Clinical Oncology, Department of Veterinary Medicine, Gifu University, 501–1193 Gifu, Japan
4)Animal Medical Center, Gifu University, 501-1193 Gifu, Japan
5)Department of Pathobiology, School of Veterinary Medicine, Nihon University, 1866 Kameino, 252–8510 Fujisawa, Japan

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**ABSTRACT.** A 9-year-old spayed female cat was examined for cheek skin drainage. The skin lesion did not respond to medical therapy; thereafter, facial deformity developed. A computed tomography revealed an intranasal mass and maxillary osteolysis. The mass was histopathologically diagnosed as suppurative granulomatous inflammation caused by filamentous bacteria. The lesion responded well to radiation therapy. Although actinomycosis was suspected histopathologically, no actinomycetes were detected in the nasal lesion by a bacterial culture conducted at a commercial laboratory. The submandibular lymph node and subcutaneous tissue exhibited swelling. Microbiological examination and genetic analysis based on 16S rDNA gene sequence revealed that *Nocardia* spp. were isolated from both lesions.

**KEYWORDS:** intranasal mass, nocardiosis, radiation therapy, suppurative granulomatous inflammation


Nocardiosis is an unusual infectious disease in animals that causes suppurative to pyo-granulomatous lesions on the skin and lungs [6]. There are also systemic or disseminated types of nocardiosis [5], but intranasal lesions caused by *Nocardia* spp. are not known to have been reported previously in dogs and cats. The definitive diagnosis of nocardiosis is generally based on microbiological culture and phenotypical characterization [4]. Recently, new molecular techniques have been introduced to identify different species of the genus *Nocardia* [4]. This case describes suppurative granulomatous sinorhinitis in a cat affected with nocardiosis.

A 9-year-old spayed female cat was examined for drainage of cheek skin. Tests for feline leukemia virus antigen and feline immunodeficiency virus antibody produced negative results with a commercially available kit (SNAP FIV/FeLV Combo Test; IDEXX Laboratories, Tokyo, Japan). Upon physical examination, a slight fever (39.2°C) was recorded, and drainage from the right cheek was observed. Although the cat had been treated for these symptoms using 8 mg/kg of cefovecin sc, the cat was brought back six months after the symptomatic treatment due to the development of left-side facial deformity, intermittent drainage, epistaxis, lethargy and eye discharge. Complete blood cell count and serum chemistry revealed only mild hyperglycemia. A mass in the left nasal cavity and osteolysis of maxilla were found in a computed tomography (CT) scan of the cat’s head. The percutaneous fine needle aspiration and perinasal biopsy of the mass revealed suppurative granulomatous inflammation. Filamentous microorganisms were also found within the lesion (Fig. 1). Bacterial and mycological cultures showed no detectable filamentous bacteria or fungus. Although neoplastic lesions were not observed histopathologically, a single dose of orthovoltage x-ray radiation therapy (6.3 Gy) was administered to the intranasal mass; attempting to improve the mass and facial deformity. The cat was also treated by the subcutaneous injection and inhalation of oxytetracycline for 2 weeks. Although a discharge of pus from the cheek occurred intermittently, systemic condition and facial deformities improved. Two years (803 days) after it was first seen, the cat was examined again with a relapse of cheek drainage and facial deformity. In spite of antimicrobial therapy and the washing of the nasal cavity, the intranasal mass size increased and was exposed on the surface of the face on day 912 (Fig. 2). A subcutaneous injection of prednisolone was combined with antimicrobial therapy. However, the mass size still increased, and the cat exhibited fever (40.1°C), anorexia and progressive worsening of body condition.

The cat was referred to the Animal Medical Center at Gifu University for palliative radiation therapy on day 984. A CT scan and a biopsy revealed an intranasal mass and granulomatous inflammation (Fig. 3). Specimens from the lesion were cultured, but no microorganisms (except for *Staphylococcus intermedius* group, which was not the cause of granuloma in this case) were obtained on the agar that contain both blood agar added NAD and MacConkey agar at 35°C with 5% CO₂ for 48 hr. Also, no anaerobes were obtained on the anaerobic rabbit blood agar and BBE agar at 35°C with anaerobical condition for 72 hr. Radiation therapy of 35 Gy in 5 fractions (7 Gy/fraction) with a linear accel-
erator was conducted on the intranasal and exposed masses. The cat was treated with minocycline on the fifth radiation, because actinomycosis was suspected, and the previously cultured \textit{S. intermedius} group was susceptible to it.

The mass reduced rapidly from the first radiation and was no longer externally visible by the day of the second radiation (Fig. 2). On the day of the fifth radiation, the intranasal mass had diminished in the CT scan image (Fig. 3), however, swelling of the subcutaneous tissue on the head and left submandibular lymph node remained. Histopathologic examination of both lesions revealed the same suppurative granulomatous inflammation.

Specimen from the lesion was cultured on Chocolate agar at 37°C with 5% CO$_2$ enriched atmosphere for 96 hr. Morphological examination of isolates revealed Gram-positive, rod-shaped and filaments microorganisms. Molecular anal-

Fig. 1. Histopathological image of intranasal mass (Hematoxylin and eosin staining, \( \times 400 \)). (A) Neutrophils, macrophages and multinucleated giant cells accumulated and formed granuloma. (B) Filamentous bacteria (arrowheads) are seen in the center of the granuloma.

Fig. 2. External appearance of the cat at the day of before (A) and after (B) radiation therapy. (A) A mass originated from nasal cavity exposed on the face. (B) The mass is not seen from the outside. A part of lower eyelid is dropped due to the mass.

Fig. 3. Non contrast enhanced CT image of the cat taken in the transverse plane at the level of orbit before (A) and after (B) radiation therapy. (A) Note the attenuated soft tissue mass in the left nasal cavity. Erosion of the maxilla allowed the mass to extend into the orbit and subcutaneous tissue. (B) The mass is not seen in the nasal cavity, except for little soft tissue.
In vitro antimicrobial susceptibility test performed using broth microdilution methods (35°C with 5% CO₂ enriched atmosphere, for 48 hr and 72 hr). Isolated bacteria are susceptible to kanamycine (≤1 mg/l) and tetracycline (2 mg/l). ABPC: Ampicillin, CEZ: Cefazolin, CTX: Cefotaxime, SM: Streptomycin, GM: Gentamycin, KM: Kanamycine, TC: Tetracycline, NA: Nalidixic Acid, CPFX: Ciprofloxacin, CL: Colistin, CP: Chloramphenicol, TMP: Trimethoprim.

### Table 1. Antimicrobial susceptibility test of isolated Nocardia flavorosea

<table>
<thead>
<tr>
<th>Antibiotic</th>
<th>48 hr (mg/l)</th>
<th>72 hr (mg/l)</th>
</tr>
</thead>
<tbody>
<tr>
<td>ABPC</td>
<td>8</td>
<td>16</td>
</tr>
<tr>
<td>CEZ</td>
<td>16</td>
<td>16</td>
</tr>
<tr>
<td>CTX</td>
<td>≤0.5</td>
<td>≤0.5</td>
</tr>
<tr>
<td>SM</td>
<td>8</td>
<td>8</td>
</tr>
<tr>
<td>GM</td>
<td>≤0.5</td>
<td>≤0.5</td>
</tr>
<tr>
<td>KM</td>
<td>≤1</td>
<td>≤1</td>
</tr>
<tr>
<td>TC</td>
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<td>4</td>
</tr>
<tr>
<td>NA</td>
<td>64</td>
<td>64</td>
</tr>
<tr>
<td>CPFX</td>
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<tr>
<td>CL</td>
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<td>&gt;16</td>
</tr>
<tr>
<td>CP</td>
<td>32</td>
<td>32</td>
</tr>
<tr>
<td>TMP</td>
<td>&gt;16</td>
<td>&gt;16</td>
</tr>
</tbody>
</table>

In vitro antimicrobial susceptibility test performed using broth microdilution methods (35°C with 5% CO₂ enriched atmosphere, for 48 hr and 72 hr). Isolated bacteria are susceptible to kanamycine (≤1 mg/l) and tetracycline (2 mg/l). ABPC: Ampicillin, CEZ: Cefazolin, CTX: Cefotaxime, SM: Streptomycin, GM: Gentamycin, KM: Kanamycine, TC: Tetracycline, NA: Nalidixic Acid, CPFX: Ciprofloxacin, CL: Colistin, CP: Chloramphenicol, TMP: Trimethoprim.

The cat’s general condition after the first radiation treatment.

Many filamentous bacteria were detected in the lesion histopathologically, but bacterial cultures failed many times. Since the bacteria grew slowly, colonies were formed after 96 hr of incubation, and thus, colonies were not detected at the commercial laboratory after 48 hr of incubation in this case. There are some reports of animal nocardiosis that bacterial colonies were obtained after 72 hr to 2 weeks of incubation [4, 7]. If nocardiosis is suspected, specimens should be incubated aerobically 72 hr or more. Also, there is a case report that definite diagnosis was made by only polymerase chain reaction (PCR) to amplify the 16S ribosomal RNA gene using DNA extracted from the pus because no microbial colony was cultured from the lesion [6]. PCR analysis should be used for detection of 16S ribosomal RNA of Nocardia spp. in the case that bacterial culture failed.

Nocardia spp. are gram-positive, aerobic or partially or variably acid-fast bacteria that form branching filamentous rods [9]. Molecular analysis based on 16S rRNA gene revealed that sequence obtained from the isolate showed 99% homology with N. flavorosea. Because sequence from 16S rRNA of N. flavorosea has 99.2% similarity with that of N. carnea, further analysis like genomic DNA-DNA hybridization is needed to differentiate these two species [3].

### REFERENCES