Squamous Cell Carcinoma in a Capybara (Hydrochoerus hydrochaeris)

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The capybara (Hydrochoerus hydrochaeris), the world’s largest rodent, is a member of the order Rodentia, family Caviidae, and inhabits the eastern Amazon basin in South America [14]. Although many capybaras are maintained in zoos, aquariums and breeding farms worldwide, few studies have investigated capybara diseases [6, 12]. Moreover, only 1 study has reported a spontaneous tumor in a capybara [19], and previous literature had not listed reports of tumors in this species [5, 13]. Guinea pigs, another member of the family Caviidae, are closely related to capybaras [14] and are frequently used experimentally. Their diseases are the most commonly encountered in numerous species, such as dogs, cats, horses and cattle [7]. In general, cutaneous squamous cell carcinoma locally infiltrates and infrequently metastasizes to the lymph node [1, 7, 10]. In the mouse, a member of the order Rodentia, spontaneous cutaneous squamous cell tumors are rare [10]. Murine cutaneous squamous cell carcinoma can arise from a pre-existing keratoacanthoma or ulcer [1, 10]. Murine squamous cell carcinoma is histologically subdivided into well-differentiated, poorly-differentiated and anaplastic types. Well-differentiated squamous cell carcinoma is characterized by polymorphic squamous keratinized tumor cells with intercellular bridges, distinct cell boundaries, inflammatory changes and desmoplastic reaction. In poorly-differentiated squamous cell carcinoma, individual keratinized squamous tumor cells are observed. Anaplastic squamous cell carcinoma is difficult to be distinguished from mesenchymal tumor due to its spindle shape [1, 10]. In humans, cutaneous squamous cell carcinoma is histologically subdivided into numerous subtypes [3, 17].

This report describes a rare spontaneous squamous cell carcinoma in a captive capybara along with the clinical course, CT findings and pathological findings.

The 4-year and 2-month-old male capybara (Hydrochoe-
rus hydrochaeris) was born in an aquarium in Nagasaki and maintained in an aquarium in Gifu, Japan. Approximately 1 year before the gluteal mass was discovered, the capybara was diagnosed with dermatosis; the entire body was covered by abnormal keratinization characterized by adherent, dry and rough excess keratinous hyperplasia, as well as crusting and scaling. Alopecia secondary to the keratinous hyperplasia was present on the entire body, especially the limbs. Bacterial dermatitis was suspected; therefore, cefaclor [10.8 mg/kg, per os (PO), ter in die (t.i.d.)] was administered empirically. After 2 weeks of treatment, the lesion resolved. However, 1 month after remission, a similar lesion recurred. Ofloxacin (4.3 mg/kg, PO, t.i.d.) was then administered to treat a resistant bacterial infection. The lesion resolved following antibiotic therapy, but once again, the lesion recurred 1 month after remission. This cycle repeated 4 times over 10 months of antibiotic therapy including sultrimicillin [16.2 mg/kg, PO, bis in die (b.i.d.)], cefpodoxime proxetil (4.3 mg/kg, PO, b.i.d.) and doxycycline hydrochloride (4.3 mg/kg, PO, semel in die). With each recurrence, a dif-

Fig. 1. Coronal section CT of a gluteal mass (circle) in a capybara, which was highly invasive, of soft tissue density, and had an indistinct border, the femur (arrows) and calcification-like lesions (arrowheads).

Fig. 2. Macroscopic photographs of a capybara diagnosed with a squamous cell carcinoma on the buttocks. A large mass is present on the buttocks and can be correlated to the mass identified on CT.

Fig. 3. Histopathological features of a squamous cell carcinoma in a capybara. HE. a) The lesion comprised islands of invasive squamous epithelial tumor cells with a severe desmoplastic reaction. Bar=500 µm. b) The tumor cells showed various degrees of keratinization with keratin pearls. Inflammatory cells frequently infiltrated many of the keratin pearls. Bar=200 µm.
f erent antibiotic was prescribed.

One and one half months after the last remission, a 3 cm wide mass was found in haired regions of the left buttock near the anus. The mass initially appeared hemorrhagic and encrusted. Surrounding areas including subepidermal tissue relatively maintain the same width. Recurrence of the abnormal keratinization was a concern, and therefore, gentamicin sulfate was applied topically onto the lesion twice daily. However, the mass continued to rise and infiltrate the surrounding tissue over approximately 6 months, and it appeared alopecic and ulcerated. The growth was mainly exophytic, gradually endophytic, cauliflower-shaped and fibrous granulomatous. A surgical biopsy sample was obtained, fixed in 10% formalin and evaluated by the Laboratory of Veterinary Pathology at Gifu University. The tissue was then embedded in paraffin, cut into 3 to 4 μm sections and stained with hematoxylin and eosin (H&E); it was ultimately diagnosed as a squamous cell carcinoma. Since the tumor had conglutinated toxylin and eosin (H&E), it was ultimately diagnosed as a squamous cell carcinoma. The neoplastic tissue infiltrated the muscle layer, which had a particularly severe desmoplastic reaction. Metastasis was present and replaced nearly the entire inguinal lymph node. On immunohistochemistry, the tumor cytoplasm was intensely positive for cytokeratin AE1/AE3, confirming the cell origin as being epithelial.

Very few studies have investigated the diseases of capybara [6, 19]. Scabies, ticks in the genus *Amblyomma*, helminthes, cocciidia, trypanosomiasis, brucellosis and sanitary management in breeding farms are mainly reported [6], while studies of noninfectious diseases, particularly neoplasia, are scarce [6, 19]. This is the first known report of squamous cell carcinoma in a capybara. The only similar previous report described a fibrosarcoma in an 8-year-old capybara [19], far older than the capybara presently reported. Capybaras live for 8 to 10 years [4], and in guinea pigs, which are closely related to capybaras, tumors are more prevalent in older animals [9]. Therefore, an external factor other than age may have contributed to tumor development in the juvenile of the present case. There are several factors known to spur squamous cell carcinoma development. Ultraviolet light, keratoacanthoma, viruses, hyperplasia and chronic inflammation can all progress or induce cutaneous squamous cell carcinoma in mice [1, 10]. In the present case, the tumor arose mostly above the skin surface initially, and therefore, it may not have arisen from a pre-existing keratoacanthoma [1]. Notably, the patient experienced intermittent chronic episodes of abnormal keratinization remission and recurrence that predated the present tumor; this abnormal keratinization may have been related to neoplastic development. Bacterial dermatitis was initially suspected as the underlying etiology; therefore, antibiotics were administered, and the lesion temporarily responded. However, the actual cause of the dermal lesions was not diagnosed; therefore, clear causality cannot be established. Yet, the recurrent abnormal keratinization may have been an inciting factor in the tumor, based on the similar pattern of keratinization in both the dermal lesions and the mass itself.

In the present case, the tumor was diagnosed as a well-differentiated squamous cell carcinoma based on murine histological classification [1, 10] and had several remarkable characteristics. The desmoplastic reaction was severe, and tumor cells surrounded by reactive fibroblasts had infiltrated the muscle. The desmoplastic reaction of squamous cell carcinoma accompanies tumor infiltration. Human desmoplastic squamous cell carcinoma, which is a stroma-rich squamous
cell carcinoma, metastasizes and locally infiltrates more often than does the common squamous cell carcinoma [3, 17]. Similarly, in the present case, metastasis was observed at the regional lymph nodes. Based on these pathological features, the present case may represent a characteristic histological squamous cell carcinoma subtype similar to human desmoplastic squamous cell carcinoma. Unlike human lesions, cutaneous squamous cell carcinoma in animals is not differentiated into complex subtypes [1, 7, 10]. Additional investigation may reveal that highly-aggressive cutaneous squamous cell carcinoma in capybara is accompanied by a more severe desmoplastic reaction.

As reported previously [11], CT-based virtopsy was a useful guide for necropsy in the present case. The postmortem CT provided valuable information in the present case by revealing the absence of osteolysis. In addition, the histological and CT findings were correlated to a degree. A severe desmoplastic reaction found histologically may be correlated to the soft tissue density on CT [16]. However, metastasis was not detected on CT; this may be due to the unavailability of a contrast medium, which provides information about metastasis and tumor characteristics [18]. A contrast medium could not be used, because the CT was performed after euthanasia. More investigations about the filming condition are needed to use CT scans as virtopsy.

This report presents the clinical course and pathological findings of a cutaneous squamous cell carcinoma in a capybara. Postmortem CT-based virtopsy revealed that the mass was invasive and of soft tissue density with no metastasis detected. However, on histological analysis, metastasis was found in the regional lymph nodes. These differences between virtopsy and histological analysis may resolve by contrivance. Histologically, the mass showed features similar to a murine well-differentiated squamous cell carcinoma. Additional reports could further characterize tumors in capybara and the family Caviidae.

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