

## Case Report

# Salivary Mucocele in a Laboratory Beagle

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**Abstract:** The histologic characteristics of a salivary mucocele in a beagle used in a toxicity study are described in this report. A pale yellowish cyst under the mandibular skin containing frothy mucus was observed at necropsy. Microscopically, numerous villous projections arose from the internal surface of the cyst and were lined by stratified epithelial-like macrophages, which were immunopositive for macrophage scavenger receptor A. A ruptured sublingual interlobar duct connected to the lumen was observed near the cyst. Luminal amorphous material showed a positive reaction with Alcian blue and periodic acid-Schiff staining as did mucin in the sublingual gland. Ultrastructurally, the epithelial-like macrophages had numerous vacuoles containing electron-lucent material, which was presumed to be lysosomal in origin, and had pseudopods on their cell surfaces interdigitating with those on the adjacent cells. This case report helps to understand the diversity of the background findings in beagles used in toxicity studies. (DOI: 10.1293/tox.24.131; *J Toxicol Pathol* 2011; 24: 131-135)

**Key words:** salivary mucocele, sialocele, macrophage, beagle, toxicity study

A salivary mucocele is defined as an accumulation of leaking salivary secretion in single or multiloculated cavities in the connective tissue of the mouth or neck contiguous to a salivary gland or duct<sup>1,2</sup>. In general, it can be observed in the form of a cyst lined with granulation tissue and the absence of any epithelial lining. A salivary mucocele can occur in several kinds of animals including dogs of any breed<sup>1</sup>. The cause is generally not identified; however, blunt trauma, foreign body and sialolith have been suspected as major causes of salivary mucocele. It arises most commonly from the sublingual salivary gland, either from individual units of the polystomatic portion or from the duct of the monostomatic portion in dogs<sup>1-3</sup>. Saliva usually leaks from the torn portion, and accumulates in the adjacent tissue. Consequently, the accumulated saliva induces an inflammatory response. A wall of granulation tissue is gradually developed in response to the inflammation. The diagnosis is usually made based on the history, physical examination by palpation and cytological examination after aspiration of the cyst and can be confirmed by histopathological examination<sup>4</sup>.

In a toxicological study, we encountered a laboratory beagle with a salivary mucocele characterized by lining

epithelial-like cells. It was considered to be a spontaneous lesion because there were no similar lesions in the other animals. Unexpectedly, there are only a few reports focused on the histological features of salivary mucoceles in animals. Therefore, we describe the histopathological characteristics of this lesion.

The animal was a male beagle (Kitayama Labes Co., Ltd., Yamaguchi, Japan) treated with a compound in a 2-week repeated-dose toxicity study with a 4-week recovery period. The experiment procedures were approved by the Animal Ethics Committee of Takeda Pharmaceutical Company Limited. No abnormality was observed in the appearance of the neck and mandible or in the clinical pathology. The animal was sacrificed at 14 months old by exsanguination under thiopental sodium anesthesia and subjected to a complete necropsy. At necropsy, a pale yellowish cyst, 10 × 40 × 12 mm in size, containing frothy mucus was observed under the mandibular skin. The proximal edge of the cyst was not clearly identified because it was buried between the digastric and mylohyoid muscles. The other organs had no abnormal gross findings. The cyst was fixed in 10% neutral buffered formalin solution. Routine paraffin embedded sections were stained with hematoxylin and eosin (HE). Alcian blue (pH 2.5)-periodic acid-Schiff (AB-PAS) double staining and immunohistochemical staining for human cytokeratin (hCK), which identifies cytokeratin 5, 6, 8, 17 and 19, and macrophage scavenger receptor A (MSR-A) were performed. The antibodies used for this study were as follows: anti-human cytokeratin mouse monoclonal antibody

(diluted 1:500, clone: MNF116, Dako Cytomation, Tokyo, Japan) and anti-human macrophage scavenger receptor A mouse monoclonal antibody (diluted 1:100, clone: SRA-E5, Transgenic Inc., Kobe, Hyogo, Japan). For electron microscopy, small pieces of the cyst wall, originally fixed in 10% neutral buffered formalin solution, were postfixed in osmium tetroxide, dehydrated and embedded in epoxy resin. Ultrathin sections were cut and stained with lead citrate and uranyl acetate and examined using a transmission electron microscope.

Light microscopically, the cyst was encapsulated by dense connective tissue and numerous villous projections arose from the internal surface of the cyst (Fig. 1a). Villous projections had fibrovascular stalks with lymphoplasmacytic cells and pigmented macrophages and were lined by stratified epithelial-like cells (Fig. 1b). The epithelial-like lining cells had round to oval nuclei and slightly eosinophilic and foamy cytoplasm. The lumen of the cyst was filled with eosinophilic amorphous material with a few desquamated cells (Fig. 1b). Some multinucleated giant cells were also observed on the surface of the lining cells (Fig. 1c). The sublingual gland (Fig. 1d) and a ruptured sublingual interlobar duct (Fig. 1e) connected to the cyst were observed in the surrounding connective tissue. The amorphous material within the cyst showed a positive reaction for AB-PAS with a bluish or reddish violet coloration (Fig. 1f). The epithelial-like cells also had AB-PAS-positive reddish violet colored granules in their cytoplasm (Fig. 1g). As expected, secretory granules of the sublingual gland and saliva in the ducts showed a similar positive reaction for AB-PAS (Fig. 1h).

Immunohistochemically, the epithelial-like cells were positive for MSR-A, a macrophage marker in dogs<sup>6</sup> (Fig. 2a-b), which strongly suggests that these cells were macrophages. On the other hand, the multinucleated giant cells were negative for MSR-A (Fig. 2b: Inset). As described in previous papers, the expression of MSR-A increases during differentiation and maturation of macrophages<sup>7</sup>. Therefore, expression of MSR-A might be changed during the process of multinucleated giant cell transformation. Predictably, the epithelial-like cells were negative for hCK (Fig. 2c), while the epithelial cells of the interlobar duct were strongly positive for hCK (Fig. 2d).

Electron microscopically, the epithelial-like cells had numerous vacuoles containing electron-lucent material, which was presumed to be lysosomal in origin (Fig. 3a). This finding probably corresponded to the reddish violet colored granules stained by AB-PAS. In addition, the epithelial-like cells had pseudopods on their cell surfaces interdigitating with those on the adjacent cells (Fig. 3b). Structures of the basal lamina or the cell junctions were not observed.

Based on the results described above, we concluded that the epithelial-like cells lining the cyst were macrophages because they had typical macrophage characteristics, such as positive staining for MSR-A, and contained many lysosomes without a basement membrane. As far as we know, cystic structures characterized by epithelial-like macrophages have not been reported except for salivary

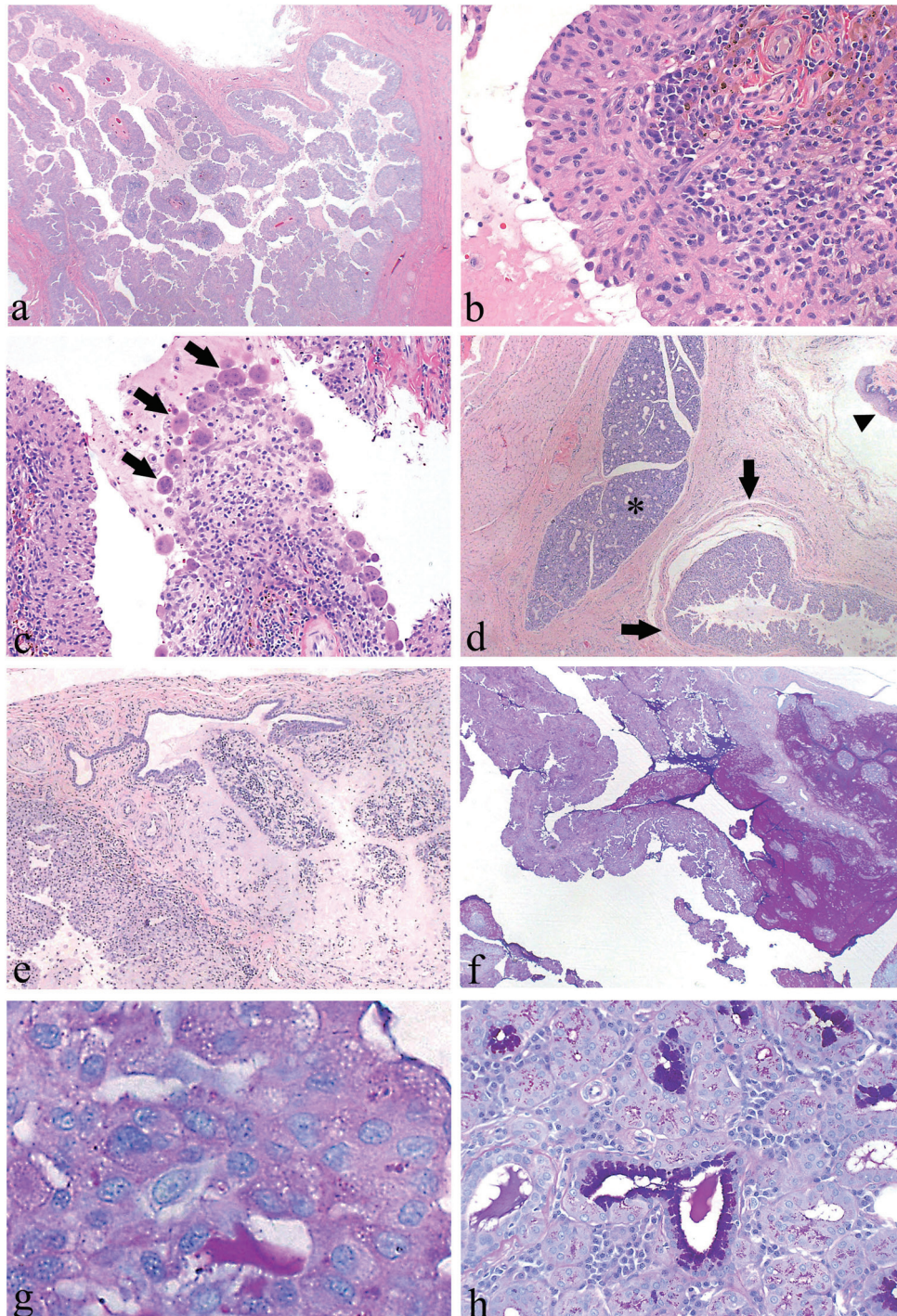
mucocele of the soft palate with fibrocytes giving the appearance of an epithelial lining<sup>8</sup>. In addition, the macrophages had no nuclear atypia, few mitoses and no monotonous proliferation. Therefore, this lesion was not considered to be a neoplastic lesion but an inflammatory response to sublingual mucoid material. In animals, cystic structures in the salivary gland are classified into two subtype, the extravasation type, a salivary mucocele, also called a sialocele or extravasated pseudocyst, and the retention type, a true salivary cyst, also called a dilatation of the duct<sup>1,3</sup>. Salivary mucoceles lack an inner luminal epithelial lining. Conversely, true salivary cysts have an epithelial lining, such as a nonkeratinized stratified squamous or interlobular duct epithelium<sup>1-3</sup>. In the present case, the cyst was lined by epithelial-like macrophages instead of epithelial tissues. Therefore, we diagnosed it as a salivary mucocele. In humans, a salivary cyst on the floor of the mouth is clinically called a ranula with or without an epithelial lining<sup>9-11</sup>. On the other hand, a textbook for animals<sup>1</sup> states that the term "ranula" should be only applied to a cystic distension of the duct, although some of the literature for animals has also applied the term "ranula" to a salivary gland cyst occurring from the sublingual gland with or without an epithelial lining<sup>5,12,13</sup>. It seems that the nomenclature of these lesions is controversial in animals. Therefore, we did not apply the term "ranula" in the diagnosis.

In the present case, fortunately, we found a ruptured sublingual interlobar duct that could be a cause of this lesion. However, it is not adequate to allege that the ruptured duct was the primary site of leakage. It is reported that salivary mucoceles sometimes have more than one communication between the sublingual gland and the cyst.<sup>14,15</sup> In view of the orientation of the cyst, it was speculated that the primary leakage probably occurred in the individual units of the polystomatic portion or the duct of the monostomatic portion.

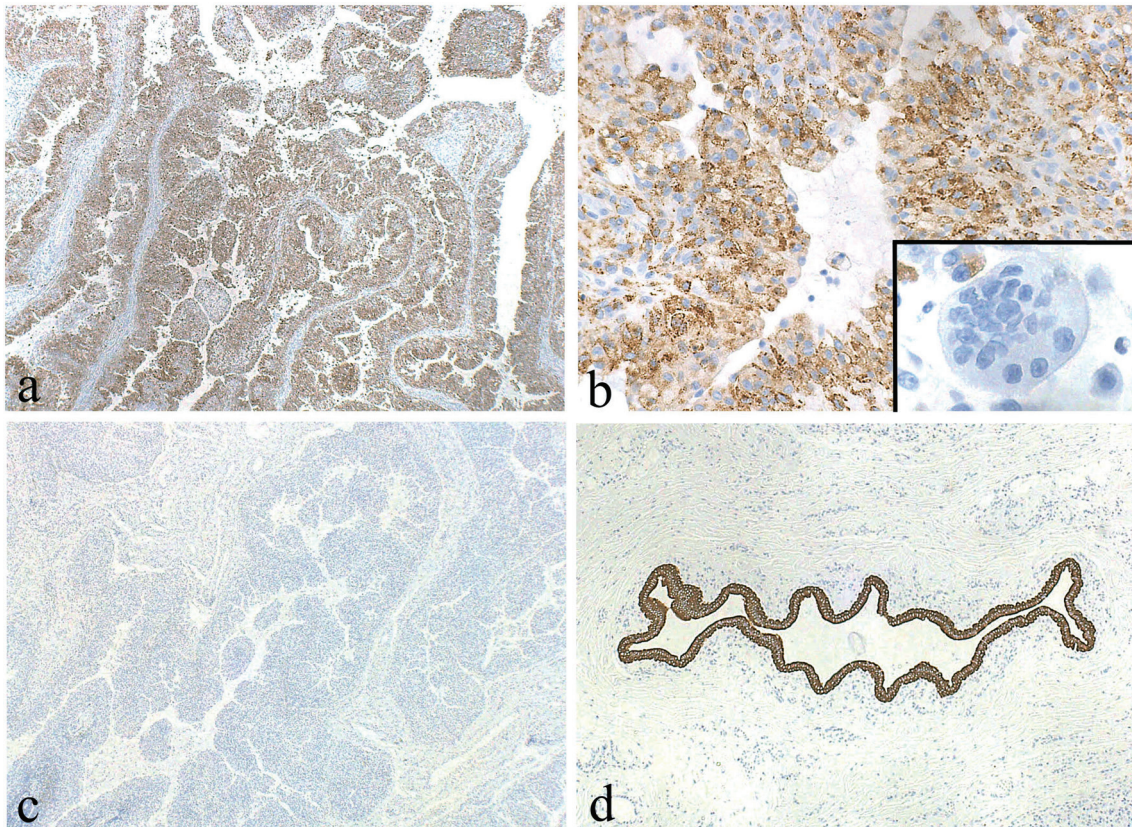
The histological appearance of salivary mucoceles varies greatly depending on the stage of development<sup>1</sup>. This case was recognized as a relatively long-standing lesion because the cyst was circumscribed by mature dense connective tissue with abundant vessels. On the other hand, the lumen of the cyst was filled with eosinophilic amorphous material and desquamated cells, suggesting that sublingual secretion could still be leaking out consistently and that this lesion would be in the process of formation. These prolonged irritations might induce accumulation of a large number of macrophages and influence its morphological changes, leading to an epithelial-like appearance.

We believe this case report helps to understand a diversity of the background findings in beagles used in toxicity studies.

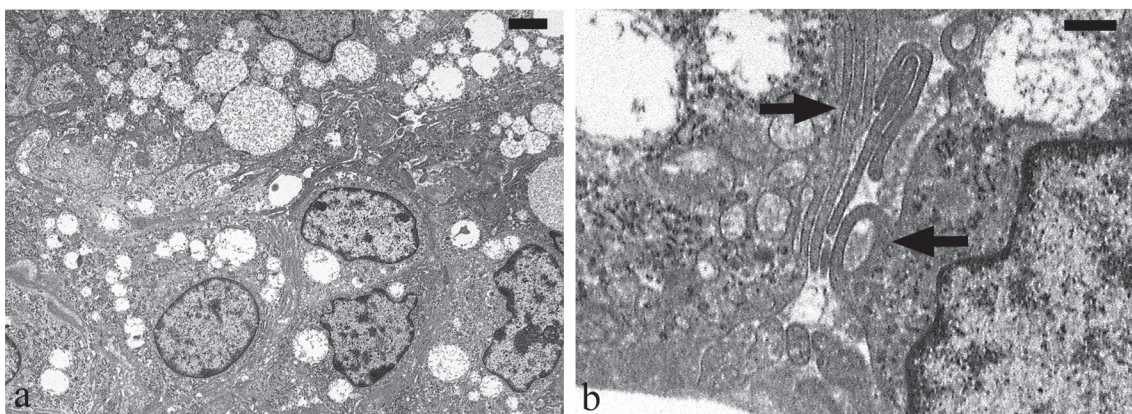
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**Fig. 1.** Histopathology of the cyst in the laboratory beagle. (a) The cyst was circumscribed by mature dense connective tissue, and the wall frequently projected into the lumen with fibrovascular connective tissue stalks. (b) The lining cells were morphologically similar to epithelial cells and projected up toward the lumen. The epithelial-like cells, lining cells, had round to oval nuclei and slightly eosinophilic and foamy cytoplasm. Granulation tissue, abundant vessels with fibroblasts, lymphocytes, pigmented macrophages and plasma cell infiltrations were observed inside of the villous projection. The lumen of the cyst was filled with eosinophilic amorphous material with a few desquamated cells. (c) Some multinucleated giant cells (arrows) were also observed in the lumen side. (d) Normal sublingual gland tissue (asterisk) was observed in the connective tissue near the cyst (arrows). In the upper right side, the normal oral mucosa was observed (arrowhead). (e) A ruptured sublingual interlobar duct connected to the lumen was observed in the peripheral connective tissue. (f) The amorphous material showed a positive reaction with a bluish or reddish violet coloration. (g) The epithelial-like cells had reddish violet colored granular staining in their cytoplasm. (h) In the nearby normal sublingual gland, the cytoplasm of the mucous cells and the luminal side of the serous cells showed violet colored granular staining. The secretion in the lumen showed a positive reaction with a bluish or reddish violet coloration. HE:  $\times 11$  (a),  $\times 206$  (b),  $\times 110$  (c),  $\times 17$  (d),  $\times 55$  (e). AB-PAS double stain:  $\times 17$  (f),  $\times 432$  (g),  $\times 103$  (h).



**Fig. 2.** Immunohistochemical staining for MSR-A and hCK. (a-b) The epithelial-like cells were positive for MSR-A. Multinucleated giant cells were negative for MSR-A (b: Inset). (c) The epithelial-like cells were negative for hCK. (d) The epithelial cells of the interlobar duct were strongly positive for hCK. Immunohistochemistry, counterstained with hematoxylin,  $\times 17$  (a, c),  $\times 41$  (d),  $\times 103$  (b) (Inset:  $\times 450$ ).



**Fig. 3.** Electron microscopy of the epithelial-like cells. (a) The epithelial-like cells had numerous vacuoles containing electron-lucent material, which was presumed to be lysosomal in origin. (b) The epithelial-like cells had pseudopods (arrows) on their cell surfaces interdigitating with those on the adjacent cells. Scale bars: 2  $\mu\text{m}$  (a), 0.5  $\mu\text{m}$  (b).

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