INTRODUCTION

Renal oxalosis in animals is most frequently caused by ethylene glycol poisoning, ingestion of oxalate-containing plants, or ingestion of plants colonized by oxalate-producing fungi (Sanz & Reig 1992, Rahman et al. 2013). In amphibians, there are a few reports of renal oxalosis in captive-bred tadpoles and frogs consuming oxalate-rich plants or crickets that fed on oxalate-containing plants (Wright 2001). In this study, we found oxalate poisoning in free-living tadpoles of American bullfrogs *Lithobates catesbeianus* from the Kyusyu region, Japan. Severe coelomic and subcutaneous edema and nephropathy were observed in affected tadpoles, and they had features that appeared to be associated with renal nephropathy and metabolic disorder. Since the pathological features and pathogenesis of renal poisoning in tadpoles have not been characterized, the aim of the present study was to further investigate the features of this disorder, including gross renal pathological and histopathological changes. To our knowledge, this is the first report of oxalate nephropathy in free-living amphibians.
MATERIALS AND METHODS

Study area and tadpole evaluation

A total of 68 free-living American bullfrog tadpoles were examined between February and March 2014 (Table 1). These included 57 tadpoles caught in an artificial pond (pond A) in a zoological garden located in the Kyusyu region, and 11 tadpoles from another artificial pond (pond B) in the same zoological garden as a normal control group. These ponds were natural breeding grounds of American bullfrogs. Pond B was closely situated to, but located upstream of, pond A. Pond A contained large amounts of sediment containing fallen leaves and mud, whereas there was little sediment in pond B. All animal experiments in this study were carried out with permission from the Ministry of Environment (121221300) and in accordance with the Japanese Law of Animal Welfare and Care.

The tadpoles were euthanized using FA100 (Tamura-seiyaku) and necropsied. For the detection of ranavirus, DNA was extracted from the left kidney, spleen, and a part of the liver of the tadpoles using NucleoSpin® Tissue (Takara Bio), and polymerase chain reaction (PCR) was performed according to the protocol described by Marsh et al. (2002). For histopathology, visceral organs were fixed in 10% neutral buffered formalin. The tissues were embedded in paraffin, sectioned at a thickness of 3 µm and stained with hematoxylin and eosin (H&E). Sections were examined under polarized light for birefringence. In order to identify calcium oxalate, sections were stained with Alizarin Red S at pH 7.0 and 4.2, and insolubility was determined in 0.1 N hydrochloric acid (HCl) and 2 M acetic acid, as described by Proia & Brinn (1985). Coelomic fluids were aseptically collected from 4 tadpoles with severe coelomic edema and directly homogenized together with 2 ml of deionized water for 1 min. The homogenates were centrifuged at 500 \( \times g \) for 10 min, and these supernatants were centrifuged at 8000 \( \times g \) for 10 min. Mud (50 g) and fallen leaves (20 mg) were collected from pond A in March 2014 and preserved at –30°C until use. The frozen mud from pond A was directly homogenized for 15 min. Defrosted frozen leaves from pond A were homogenized together with 10 ml of deionized water for 1 min. The supernatants of homogenates were immediately filtered through filter paper. The filtrates were further filtered using a spin column Ultrafree-MC 0.45 µm PTF membrane (Millipore) and centrifuged at 4000 \( \times g \) for 30 min. HPLC analysis was performed using the Shim-pack SCR-102H column (Shimadzu) in the LC-10A system (Shimadzu) with 5 mM p-toluenesulfonic acid as the dissociate solvent, at a speed of 0.8 ml min\(^{-1}\) with the column temperature at 40°C.

RESULTS

Of the 57 tadpoles from pond A, 4 (12.3%) appeared to have obvious abdominal distension on the body with hemorrhages within the skin when captured (Fig. 1). At necropsy, severe coelomic and/or subcutaneous edema were seen in 7 of 57 (12.3%) tadpoles from pond A. Kidney enlargement and multifocal white renal calculi in the kidney (Fig. 2) were seen at gross examination in 19 of 57 (33.3%) tadpoles from pond A. The other tadpoles from pond A and 11 tadpoles from pond B were macroscopically normal. Ranavirus was not identified in the kidney, spleen, or liver from the 68 tadpoles.

On histopathological examination, crystal deposition was found in the kidneys of 35 of 57 (61.4%) tadpoles from pond A (Table 1). Minor crystal deposition, characterized by a few small crystals, detectable only under polarized light, occurred in 10 of 35 (28.6%), moderate crystal deposition was present in

<table>
<thead>
<tr>
<th>Date</th>
<th>Pond</th>
<th>Number of tadpoles</th>
<th>With coelomic edema</th>
<th>With crystal deposition</th>
</tr>
</thead>
<tbody>
<tr>
<td>20 Feb 2014</td>
<td>A</td>
<td>10</td>
<td>3</td>
<td>9</td>
</tr>
<tr>
<td>17 Mar 2014</td>
<td>A</td>
<td>18</td>
<td>nr</td>
<td>12</td>
</tr>
<tr>
<td>13 Feb 2014</td>
<td>A</td>
<td>29</td>
<td>4</td>
<td>14</td>
</tr>
<tr>
<td>13 Feb 2014</td>
<td>B</td>
<td>11</td>
<td>0</td>
<td>0</td>
</tr>
</tbody>
</table>

Table 1. Prevalence of coelomic and subcutaneous edema and crystal deposition in the kidneys of American bullfrog Lithobates catesbeianus tadpoles; nr: not recorded.
18 of 35 (51.4%), and severe crystal deposition scattered throughout histological sections was seen in 7 of 35 (20%) tadpoles. The crystals were found mostly in renal tubules that were irregularly dilated by these crystals (Fig. 3A), and the tubular epithelium showed flattening or necrosis with hyaline droplet degeneration. No crystal depositions were observed in the kidney of 11 tadpoles from pond B and within the other organs of all tadpoles examined.

The crystals were transparent, pleomorphic in shape, highly birefringent in polarized light, and arranged in a radial pattern within the renal tubular lumen (Fig. 3B). Thorough fixation and routine processing did not result in dissolution of these crystals. The crystals stained positively with Alizarin Red S at a pH 7.0 (Fig. 3C) but not at pH 4.2, and they were soluble in 0.1 N hydrochloric acid but insoluble in 2 M acetic acid.

The biochemistry of coelomic fluid from 4 tadpoles with edema and severe crystal deposition in the kidneys is shown in Table 2. The ammonia levels in the coelomic fluid were high in these tadpoles. In HPLC analyses, the chromatograms showed oxalic acid peaks in the kidneys, mud, and fallen leaves, with concentrations of 0.35, 0.02, and 0.02 mg g⁻¹, respectively.

**DISCUSSION**

The morphological characters of crystal deposition in the kidneys of the tadpoles and their birefringence...
Calcium oxalate crystals stain with Alizarin Red S at a pH of 7.0 but not at a pH of 4.2. This differentiates calcium oxalate from calcium carbonate and calcium phosphate, both of which stain at pH 7.0 as well as 4.2 (Proia & Brinn 1994, Yanai et al. 1995). Calcium oxalate deposition has been reported in koala *Phascolarctos cinereus*, white-tailed deer *Odocoileus virginianus*, green turtle *Chelonia mydas*, desert tortoise *Gopherus agassizii*, baboon *Papio ursinus*, and Japanese macaque *Macaca fuscata* (Wyand et al. 1971, McConnell et al. 1974, Yanai et al. 1995, Stacy et al. 2008, Jacobson et al. 2009, Speight et al. 2013). In these cases, most animals were asymptomatic, and these occurrences were presumed to result from the ingestion of oxalate-rich plants. Wright (2001) reported that consumption of crickets fed on oxalate-containing plants was responsible for renal oxalosis in captive-breeding waxy frog *Phyllomedusa sauvagii* and relict leopard frog *Rana onca* (Wright 2001); however, reports regarding oxalate poisoning in free-living amphibians are lacking.

In this study, we tried to find the possible source of oxalate ingestion by the tadpoles. There were no sources of ethylene glycol in pond A, nor was there any evidence of *Aspergillus* infection. American bullfrog tadpoles are herbivorous and feed on plants and algae (Altig et al. 2007). Environmental analyses revealed the presence of oxalate in the fallen leaves and mud in pond A. Even though the source of oxalate in the present case remains unknown, it was most likely that the tadpoles ingested oxalate by feeding on sediments containing oxalate-containing plants.

The American bullfrog is native to North America. It has been introduced in several other countries and islands around the world, and is considered the most important pest in these regions. Recently, American bullfrogs have been widely suggested as infection-tolerant reservoirs for emerging infectious diseases, such as those caused by viral and fungal pathogens (Une et al. 2009). In the present study, edema was one of the findings in tadpoles with renal oxalosis, and it has also been observed in emerging ranaviral infection (Pessier 2009). This suggests the importance of obtaining a differential diagnosis for both diseases.

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