Case Report

Zinc toxicosis in a cat associated with ingestion of a metal screw nut

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Abstract

Case summary A 3-year-old female neutered domestic shorthair cat was presented for vomiting, inappetence and weight loss. The cat developed moderately regenerative anaemia, moderately increased alanine transaminase and alkaline phosphatase activities, hyperbilirubinaemia and prolonged activated partial thromboplastin time. Abdominal ultrasound identified gastric wall thickening and changes suggestive of pancreatitis. Gastroduodenoscopy identified a metal screw nut in the pylorus, which was removed with rat tooth forceps. Metal analysis and serum zinc concentration using leftover serum collected at admission were performed after screw nut removal. Serum zinc concentration was markedly elevated, confirming a diagnosis of zinc toxicosis. Metal analysis of the screw nut showed that the major metal component was zinc. The cat recovered after screw nut removal and supportive care. Clinical signs resolved and the serum zinc concentration reduced significantly after screw nut removal.

Relevance and novel information Reports of zinc toxicosis in cats are scarce, possibly due to the more discriminating eating habits of this species. To our knowledge, this is the first report of zinc toxicosis causing haemolytic anaemia, liver enzyme activity increases, gastrointestinal signs and pancreatitis in a cat associated with ingestion of a zinc-containing metal object.

Keywords: Zinc toxicosis; haemolytic anaemia; liver; pancreatitis

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Case description

A 3-year-old female neutered domestic shorthair cat was presented to the emergency service as a referral for 2 weeks of inappetence and weight loss. One week earlier the cat was presented to the local veterinarian for inappetence, vomiting and 600 g weight loss. Haematology was unremarkable. Biochemistry revealed mildly increased alanine transaminase (ALT) (131 U/l; reference interval [RI] 12–130) and alkaline phosphatase (ALP) (256 U/l; RI 14–111). The cat was treated with intravenous (IV) fluids, amoxicillin–clavulanic acid, gabapentin, maropitant, mirtazapine and buprenorphine, and was discharged after 24 h. Five days later, inappetence continued and the cat was administered diazepam IV and dexamethasone subcutaneously to stimulate appetite. The cat then presented 1 day later for persistent inappetence and was treated with sucralfate before being transferred to the emergency service. The cat had no known history of access to toxins or foreign bodies.

Physical examination at referral revealed that the cat was quiet, alert and responsive, had pale-pink mucous membranes, tachycardia (200 beats/min), tachypnoea (60 breaths/min) and a body condition score (BCS) of 3/9. Packed cell volume (PCV) was 18% with a total protein of 66 g/l. Haematology showed a normocytic normochromic moderately regenerative anaemia with a haematocrit of 0.15 l/l (RI 0.25–0.48), an absolute reticulocyte count of 102 ×10^9/l (RI 3–50) and moderate...
neutrophilia of $20.8 \times 10^9/l$ (RI 2–13). Differential diagnoses for anaemia included haemolysis or haemorrhage. Biochemistry showed moderately increased ALT (249 U/l; RI <60) and ALP activity (277 U/l; RI <50), mild hyperbilirubinemia (10.6 µmol/l; RI 2.5–3.5) and mild hypercholesterolaemia (6 mmol/l; RI 1.9–3.9). Point-of-care feline leukaemia virus antigen and feline immunodeficiency virus antibody tests were negative. The activated partial thromboplastin time (APTT) was mildly prolonged at 132.0 s (RI 65–119) and prothrombin time (PT) was within the RI at 19.0 s (RI 15–22). Abdominal ultrasound identified diffuse concentric submucosal thickening of the gastric wall (wall thickness 0.3 cm; Figure 1) and a hypoechoic pancreas with a mild increase in peripancreatic peritoneal echogenicity (Figure 2). Ultrasonographic findings were suggestive of an intramural gastropathy and pancreatitis. Samples were sent for haemotropic mycoplasma PCR (*Mycoplasma haemofelis, Candidatus Mycoplasma haemominutum* and *Candidatus Mycoplasma turicensis*), faecal PCR (feline coronavirus, *Trichomonas foetus*, *Cryptosporidium* species, panleukopenia virus, *Clostridium perfringens*, *Giardia* species, *Salmonella* species, *Toxoplasma gondii, Campylobacter jejuni* and *Campylobacter coli*) and faecal flotation to investigate possible causes of the regenerative anaemia and gastrointestinal signs. The cat was treated with esomeprazole (1 mg/kg IV q12h) and vitamin K1 (3.8 mg/kg PO q12h). PCV decreased to 12% over 8h with a total protein of 68 g/l, and the cat received a 45 ml type A fresh whole-blood transfusion, after blood typing. The next day, PCV was 20% and the cat was eating. The cat was discharged with omeprazole (1.9 mg/kg PO q12h), sucralfate (0.1 g/kg PO q8h), marbofloxacin (4.8 mg/kg PO q24h) and vitamin K1 (3.8 mg/kg PO q12h) while test results were pending. Further diagnostic tests, including abdominal radiographs and endoscopy, were declined due to financial constraints.

Two days later, the cat presented for recurrent inappetence. Haematology revealed a non-regenerative anaemia with a haematocrit of 0.17 l/l and a mild neutrophilia (15.3 $\times 10^9/l)$. Gastroduodenoscopy was performed under general anaesthesia to investigate a possible gastrointestinal cause of inappetence, weight loss and anaemia. A non-obstructive metal object (the nut of a cam lock screw used in flat-packed household furniture) was found in the pylorus and was removed with rat tooth forceps (Figures 3 and 4). Re-inspection of the pyloric region identified a minor erosion (Figure 5). The rest of the stomach, oesophagus and duodenum were unremarkable. An oesophageal feeding tube was placed and treatment included IV fluid therapy, omeprazole, vitamin K1, sucralfate, marbofloxacin, mirtazapine (1.4 mg/kg PO q72h) and maropitant (1.5 mg/kg PO q24h).
Two days later, PCV was 20% with a total protein of 60 g/l. The cat started eating consistently and the feeding tube was removed after 6 days. Haemotropic mycoplasma PCR, faecal flotation and PCR were negative. The cat was discharged with mirtazapine only and the rest of the medications were discontinued. Body weight at discharge was 2.4 kg. After screw nut removal, metal analysis and serum zinc concentration using leftover serum collected at admission were performed. Serum zinc concentration, available 25 days later, showed a markedly elevated concentration at 448 µmol/l (RI 7.65–16.8), confirming zinc toxicosis. Metal analysis of the screw nut by inductively coupled plasma mass spectrometry (ICPMS) (Table 1) showed it was mainly composed of zinc. High levels of aluminium, copper, iron and nickel and a very low level of lead and cadmium were detected.

Seven weeks later, the cat was clinically well with a normal appetite. Examination revealed a body weight of 3.2 kg (800 g gain) and a BCS of 5/9. Haematology was unremarkable (haematocrit 0.30 l/l). Biochemistry was unremarkable: ALT, ALP and total bilirubin were all within the RIs. APTT (95 s) and PT (18 s) were within the RIs. Abdominal ultrasound was unremarkable. Repeat serum zinc concentration was 42.4 µmol/l.

### Table 1: Inductively coupled plasma mass spectrometry metal analysis of the screw nut (weight 1.340 g)

| Metal       | Concentration (µg/g screw nut) | Composition (%) |
|-------------|-------------------------------|-----------------|
| Zinc 66     | 250.835                       | 82.191          |
| Aluminium 27| 35.596                        | 11.664          |
| Copper 63   | 13.313                        | 4.362           |
| Phosphorus 31| 3.445                         | 1.129           |
| Iron 57     | 0.605                         | 0.198           |
| Sodium 23   | 0.444                         | 0.146           |
| Cadmium 111 | 0.202                         | 0.066           |
| Nickel 60   | 0.177                         | 0.058           |
| Lead 206    | 0.135                         | 0.044           |
| Lead 208    | 0.135                         | 0.044           |
| Lead 207    | 0.133                         | 0.044           |
| Potassium 39| 0.090                         | 0.030           |
| Selenium 82 | 0.031                         | 0.010           |
| Cobalt 59   | 0.012                         | 0.004           |
| Magnesium 24| 0.011                         | 0.004           |
| Chromium 52 | 0.007                         | 0.002           |
| Titanium 47 | 0.004                         | 0.001           |
| Manganese 55| 0.004                         | 0.001           |
| Zirconium 90| 0.003                         | 0.001           |
| Gold 197    | 0.002                         | 0.001           |
| Vanadium 51 | 0.001                         | 0.0003          |
| Silver 197  | 0.001                         | 0.0003          |
| Silicon 28  | ND                            | ND              |
| Mercury 202 | ND                            | ND              |
| Sulfur 32   | ND                            | ND              |
| Calcium 43  | ND                            | ND              |
| Arsenic 75  | ND                            | ND              |
| Strontium 88| ND                            | ND              |
| Barium 138  | ND                            | ND              |

ND = below the limit of detection

### Discussion

Zinc toxicosis in cats is uncommon.1 The true toxic dose for metallic zinc in cats has not been established.1 Normal serum zinc concentrations range between 0.50 and 1.10 ppm (7.65–16.8 µmol/l) in cats.1 Most cases of confirmed...
zinc toxicity have serum zinc levels exceeding 5 ppm (76.5 µmol/l). In the case reported here, the cat had a markedly increased serum zinc concentration of 448 µmol/l, accompanied by clinical signs of inappetence, vomiting and a regenerative anaemia. These findings were compatible with a diagnosis of zinc toxicosis. The serum zinc concentration had reduced to 42.4 µmol/l 7 weeks later with the resolution of clinical signs and anaemia after removal of the zinc-containing foreign object.

While the exact mechanism of zinc toxicosis is unknown, following ingestion, erosion of zinc material caused by gastric acid leads to subsequent toxicosis. The corrosive properties of zinc cause gastrointestinal signs, including vomiting, inappetence, lethargy and diarrhoea. The cat in this case presented to the referring veterinarian for vomiting prior to referral. We suspect that the zinc-containing metal screw nut was ingested at that time, 12 days prior to removal, causing gastric irritation and acute vomiting.

Following gastrointestinal absorption in the duodenum, zinc is bound to albumin and macroglobulins and is transported to the liver via the portal circulation. Approximately 30–40% of zinc is extracted by the liver and is subsequently released into the circulation. The liver, kidney, pancreas and spleen are the organs with the most rapid accumulation and turnover of zinc. In dogs, zinc toxicosis, in addition to gastrointestinal signs and haemolytic anaemia, can also cause pancreatitis, hepatopathy and acute kidney injury. In dogs with zinc toxicosis, intravascular haemolysis with normochromic or macrocytic, hypochromic anaemia and Heinz bodies is the most consistent abnormality observed. While there are no reports of anaemia secondary to zinc toxicosis in cats in the current literature, ferrets have been described to develop anaemia secondarily to gastrointestinal bleeding in an experimental study. In dogs with zinc toxicosis, intravascular haemolysis with normochromic or macrocytic, hypochromic anaemia and Heinz bodies is the most consistent abnormality observed. While there are no reports of anaemia secondary to zinc toxicosis in cats in the current literature, ferrets have been described to develop anaemia secondarily to gastrointestinal bleeding in an experimental study. In the case reported here, a moderately regenerative anaemia was identified with hyperbilirubinaemia, suggesting haemolysis. Although Heinz bodies were not observed in this case, the finding of Heinz bodies is less specific in cats and can be seen in healthy cats, cats with diabetes mellitus, hyperthyroidism and lymphoma.

The APTT was mildly prolonged and returned to within RIs after screw nut removal. Prolonged APTT has been described in dogs with zinc toxicosis. It has been speculated that zinc could cause inhibition of coagulation factors VIII, IX, XI and/or XII. Increased ALT and ALP activity, and hyperbilirubinaemia are typical findings of zinc toxicosis in dogs. These findings were also observed in the cat described here. However, it is unknown if the increase in liver enzymes was directly related to a hepatopathy or pancreatopathy. The liver is the major organ of zinc metabolism and histological lesions with centrilobular or periportal to mid-zonal hepatic necrosis have been reported in dogs with zinc toxicosis. Resolution of the anaemia and liver enzyme elevations at follow-up suggested that these clinicopathological abnormalities could be related to zinc toxicosis and/or the resolution of pancreatitis.

The ultrasonographic finding of pancreatitis in this cat was interesting. Pancreatitis and pancreatic fibrosis have been described in dogs with zinc toxicosis. In humans with zinc toxicosis, a hypoxic mechanism has been suggested as the cause of pancreatitis. Although the exact mechanism of zinc-induced pancreatitis is unknown in dogs, zinc has been shown to concentrate in the pancreas and eliminate in pancreatic secretions of dogs. Various pancreatic lesions on histopathology have been described in dogs with zinc toxicosis, including inflammation, nodular changes, fibrosis and necrosis. Pancreatic fibrosis and nodular changes were also observed in experimental cats fed with zinc. We suspect that pancreatitis in the cat reported here was associated with zinc toxicosis; however, we acknowledge that definitive diagnosis of feline pancreatitis remains challenging without histopathology and the aetiology of pancreatitis in this case remains elusive.

There are several limitations to our study. Although other metal toxicosis cannot be excluded, the clinical signs in the cat were incompatible with neurological signs expected for aluminium toxicosis. Hepatocellular injury caused by copper toxicosis is possible; however, zinc toxicosis is more likely with compatible clinical signs, and the majority of the screw nut was composed of zinc. A Coombs test or saline agglutination test was not performed as part of the diagnostic investigation for haemolytic anaemia. The justification of marbofloxacin use was based on the rapid deterioration of the cat, initial suspicion of haemotropic mycoplasmosis and hepatopathy, this was stopped when haemotropic mycoplasma PCR results returned negative. Although the use of vitamin K1 is debatable with a mildly prolonged APTT, considering the risk of coagulopathy in hepatic disease, the severe anaemia and uncommon adverse effects with oral use, vitamin K1 was given. Gastric contents and gas during the initial abdominal ultrasound precluded the identification of the metal screw nut. In the case of unexplained regenerative anaemia and gastrointestinal signs, abdominal radiographs should be performed to look for metal toxicosis. However, abdominal radiographs were declined at that time.

Conclusions

This is the first detailed clinical report of a case of zinc toxicosis in a cat. Gastrointestinal signs, regenerative anaemia, increased liver enzyme activity, prolonged APTT, gastric wall thickening and pancreatitis were observed. The clinical signs and clinicopathological abnormalities resolved after the removal of a gastric foreign object (a zinc-containing metal screw nut) and supportive treatment.
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Ethical approval The work described in this manuscript involved the use of non-experimental (owned or unowned) animals. Established internationally recognised high standards (‘best practice’) of veterinary clinical care for the individual patient were always followed and/or this work involved the use of cadavers. Ethical approval from a committee was therefore not specifically required for publication in JFMS Open Reports. Although not required, where ethical approval was still obtained, it is stated in the manuscript.

Informed consent Informed consent (verbal or written) was obtained from the owner or legal custodian of all animal(s) described in this work (experimental or non-experimental animals, including cadavers) for all procedure(s) undertaken (prospective or retrospective studies). For any animals or people individually identifiable within this publication, informed consent (verbal or written) for their use in the publication was obtained from the people involved.

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