Case Report

Unilateral Subconjunctival and Retrobulbar Hemorrhage Secondary to Brodifacoum Toxicity in a Dog

Sonia E. Kuhn and Diane V. H. Hendrix

Department of Small Animal Clinical Sciences, College of Veterinary Medicine, University of Tennessee, 2407 River Drive, Knoxville, TN 37996, USA

Correspondence should be addressed to Sonia E. Kuhn; skuhn3@utk.edu

Received 31 January 2013; Accepted 28 February 2013

Abstract

An 8-year-old spayed female mixed-breed dog was presented for an acute onset of bleeding around the left eye. Mild exophthalmos and massive subconjunctival hemorrhage on the globe and nictitating membrane were present in the left eye. Retrobulbar hemorrhage was suspected, and pain was implied on opening of the mouth because the patient resisted and vocalized. No other abnormalities were found on ophthalmic or physical examination. Further questioning of the owner confirmed potential brodifacoum ingestion, and prothrombin time and partial thromboplastin time were both markedly elevated. Treatment with oral vitamin K\textsubscript{1} was implemented, and the subconjunctival hemorrhage was significantly improved within a few days of instituting treatment. All clinical signs of coagulopathy were completely resolved within 4 weeks of presentation. Coagulopathy secondary to brodifacoum ingestion can manifest as severe unilateral bulbar and nictitating membrane subconjunctival hemorrhage and exophthalmos due to retrobulbar hemorrhage without other clinical signs.

1. Introduction

Intraocular and periocular bleeding can occur with primary disease of the globe and adnexa or as manifestations of systemic disease. Clinical signs are hyphema and hemorrhage of nearly any aspect of the eye, including the uvea, vitreous, retina, subretinal space, conjunctiva, subconjunctival, and retrobulbar space. Periocular and intraocular hemorrhages are most commonly associated with uveitis or retinal detachment [1] caused by infectious diseases, including systemic fungal [2] and rickettsial diseases [3–5]; immune-mediated diseases such as uveodermatologic syndrome [6]; bleeding and vascular disorders such as hypertension [7, 8], thrombocytopenia [9, 10], anemia [11] and coagulopathy [1]; neoplasia [1, 12]; diabetes mellitus [13]; and hyperviscosity syndrome from multiple myeloma [8, 14] and polycythemia vera [8]. Additionally, persistent hyperplastic primary vitreous [15], retinal dysplasia, preiridal fibrovascular membrane formation [16], and blunt or penetrating trauma [1, 17] can also cause intraocular hemorrhage. Retrobulbar hemorrhage occurs because of trauma or coagulopathy and can cause exophthalmos [18–21].

Conjunctival or scleral hemorrhage in dogs usually occurs focally as petechiae from a primary hemostatic disorder, which is typically due to thrombocytopenia. Although subconjunctival hemorrhage is uncommon, it can occur secondary to coagulopathy, trauma, and vasculitis [21]. It has also been reported to occur with Rocky Mountain spotted fever [3] and scleral rupture [17]. The aim of this report is to describe an atypical presentation of brodifacoum rodenticide toxicity where the only clinical signs were unilateral subconjunctival and retrobulbar hemorrhages.

2. Case Presentation

An 8-year-old spayed female mixed-breed dog was presented for an acute onset of bleeding around the left eye. The bleeding began 6 hours prior to presentation and was progressively worsening. The dog had been alone in a fenced yard for most of the day. Her activity level was appropriate, and thirst and appetite were normal. No known trauma had occurred. Prior medical history was unremarkable. The dog had been sprayed by a skunk (Mephitis mephitis) 2 weeks prior to...
presentation but did not seem to suffer any ill effects from that incident. Appropriate vaccinations were current, and she was not receiving any medication aside from heartworm and flea prophylaxis.

On ophthalmic examination, direct and consensual pupillary light responses were normal in both eyes (OU). A menace response was present OU, and the dog exhibited behavior consistent with vision bilaterally. The Schirmer tear test 1 (Schirmer tear test strips, Merck Animal Health) showed normal tear production in the right eye (OD) at 19 mm/min and was not evaluated in the left eye (OS) due to the bleeding (reference range > 15 mm/min). A drop of 0.5% proparacaine hydrochloride ophthalmic solution (Alcaine, Alcon Laboratories) was applied topically OU, and applanation tonometry (Tono-Pen Vet, Reichert, Inc.) was performed. Intraocular pressure was 15 mmHg OU (reference range 6–24 mmHg) [22]. Massive hemorrhage was present beneath the bulbar conjunctiva for 360° around the left globe as well as beneath the palpebral and bulbar conjunctiva of the nictitating membrane OS (Figure 1(a)). Mild serosanguinous discharge was also present OS. The left globe was mildly exophthalmic (Figure 1(b)), and retropulsion seemed to cause moderate pain. The left eyelids were mildly swollen, and palpation of the left bony orbit was unremarkable. Slit-lamp biomicroscopy (Kowa SL-15, Kowa) and indirect ophthalmoscopy (20 D handheld lens, Volk Optical Inc.; Vantage Plus Wireless Headset, Keeler Instruments Inc.) were performed OU. Examination of the left and right anterior and posterior segments was unchanged OU from the initial presentation. The patient was reevaluated 1 month after initial presentation. Oral vitamin K₁ had been discontinued 2 days prior to examination. The PT was within normal limits at 7.4 seconds, and PTT was slightly prolonged at 26.4 seconds. The subconjunctival and periorcular hemorrhage OS was fully resolved (Figure 2(b)). Examination of the anterior and posterior segments was unchanged OU from the initial presentation. Oral vitamin K₁ therapy was not reinstituted since the coagulopathy had resolved. Communication with the owner 3 months later revealed the dog to be completely asymptomatic and free of any remaining detectable abnormalities.

The patient was admitted to the hospital for overnight monitoring and treatment. A subcutaneous injection of 93 mg (5 mg/kg) vitamin K₁ was given at admission and was repeated 12 hours later. Oral treatment with 25 mg (1.3 mg/kg) of vitamin K₁ twice daily was then instituted for 4 weeks. The left eye was lubricated with artificial tears ophthalmic ointment (15% mineral oil with 83% white petrolatum, Rugby Laboratories) four times daily to prevent exposure keratitis secondary to exophthalmos and lagophthalmos from nictitating membrane and eyelid swelling. Exercise restriction to prevent additional bleeding was also instituted.

Two days later, PT and PTT were within normal limits at 7.7 seconds and 17.8 seconds, respectively. The subconjunctival hemorrhage and eyelid swelling OS were markedly improved (Figure 2(a)). Artificial tears ointment was discontinued since the globe was no longer exophthalmic. The Schirmer tear test 1 was within normal limits OU at 15 mm/min OD and 21 mm/min OS. Applanation tonometry showed normal intraocular pressures OU of 8 mmHg OD and 10 mmHg OS. Examination of the anterior segment and fundus was unchanged OU.

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3. Discussion

Ocular lesions have been documented with anticoagulant rodenticide exposure but are rarely mentioned in texts that discuss clinical signs of this toxicity. This may be because ocular lesions are uncommon relative to other signs, or because they are usually mild in comparison to the more life-threatening hemorrhage that typically occurs, such as hemothorax. Hyphema as well as scleral and subconjunctival hemorrhage has been previously reported with anticoagulant rodenticides [1, 21, 23–25]. The previously reported scleral hemorrhage may have been referring to subconjunctival hemorrhage, though photographs of the lesions were not provided [23, 26]. Concurrent exophthalmos or suspected retrobulbar hemorrhage was not mentioned in those cases where scleral hemorrhage was noted, and one of the cases had other obvious concurrent clinical signs of coagulopathy [26]. In one retrospective study evaluating clinical signs of coagulopathy due to anticoagulant rodenticide, scleral hemorrhage was an uncommon finding that was seen in only 3 of 52 cases [23]. Whether this hemorrhage occurred with or without other clinical signs of coagulopathy was not discussed.

Massive subconjunctival hemorrhage of the nictitating membrane has not been previously documented as a clinical
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Figure 1: At initial presentation, severe subconjunctival hemorrhage of the globe, and nictitating membrane were present around the left eye. Mild exophthalmos, frank bleeding, and serosanguinous discharge were also seen. These were the only clinical signs of coagulopathy secondary to brodifacoum ingestion.

Figure 2: (a) Two days after therapy with oral vitamin $K_1$, the exophthalmos was resolved and the subconjunctival hemorrhage greatly improved. (b) The subconjunctival hemorrhage was resolved after treatment with oral vitamin $K_1$ for 4 weeks.

sign of anticoagulant rodenticide toxicity, though exophthal-
omos secondary to retrobulbar hemorrhage is known to occur [19–21]. Retrobulbar hematoma secondary to warfarin toxicity has also been reported to cause exophthalmos in humans [27]. Other causes of acquired coagulopathy that could cause subconjunctival or retrobulbar hemorrhage in dogs are severe liver disease; vasculitis; autoimmune disease directed against a coagulation factor; disseminated intravascular coagulation (DIC); anticoagulant therapy; or low levels of vitamin K secondary to obstructive hepatopathy, malabsorption, or low dietary vitamin K [21, 28, 29].

Bleeding disorders can be classified as diseases that affect fibrinolysis, primary hemostasis, and secondary hemostasis. Fibrinolysis is responsible for fibrin clot dissolution and depends on the conversion of plasminogen to plasmin [29]. Clot dissolution forms fibrin degradation products, which subsequently are removed from circulation by the liver. Accumulation of these products causes bleeding tendencies by interfering with platelet function and thrombin inhibition [29]. Primary hemostasis seals injured blood vessels with the creation of the primary hemostatic plug via the interactions of platelets and endothelium. Platelet or endothelial diseases, such as thrombocytopenia, thrombocytopenia, and vasculopathies, cause primary hemostatic disorders [29].

Secondary hemostasis has traditionally been defined using a cascade model where the intrinsic and extrinsic enzymatic pathways converge into a common pathway that results in the conversion of fibrinogen to fibrin [29]. The fibrin produced by secondary hemostasis solidifies the primary hemostatic plug. Factors VIII, IX, XI, and XII are involved in the intrinsic pathway, while tissue factor and factor VII constitute the extrinsic pathway. The common pathway involves factor X and the conversion of prothrombin to thrombin (factor II), which ultimately leads to the cleavage of fibrin from fibrinogen. While PT evaluates the extrinsic and common pathways, PTT evaluates the intrinsic and common pathways. More recently, coagulation has been described with a cell-based model, which explains in vivo deficiencies seen with the cascade model [30]. The cell-based model views coagulation as occurring in distinct yet overlapping phases rather than separate enzymatic pathways [30]. It also accounts for the role of cell surfaces in fibrin formation as well as the additional functions of coagulation proteins beyond the coagulation cascade [30]. The liver produces most of the coagulation factors necessary for secondary hemostasis [29]. Factors II, VII, IX, and X are known as vitamin K-dependent coagulation factors since they contain glutamyl residues that must be activated with carboxylation, which requires reduced vitamin $K_1$ as a cofactor [28, 31]. These glutamyl residues allow for binding of the coagulation protein to a cell membrane surface via calcium binding, and calcium binding cannot occur unless carboxylation occurs [30].
Carboxylation results in the oxidation of vitamin K₁, and the enzyme vitamin K₁ epoxide reductase is necessary to reduce vitamin K₁ back to its active form so that it can be recycled to activate additional coagulation factors [28]. Disorders of secondary hemostasis are due to decreased concentrations of or ineffective coagulation factors and result in coagulopathies.

Anticoagulant rodenticides, as described in this case, cause coagulopathy by depletion of vitamin K-dependent coagulation factors via inhibition of vitamin K₁ epoxide reductase [20, 31]. Warfarin and pindone are first-generation anticoagulants, while brodifacoum, bromadiolone, and diphenacine are second-generation anticoagulants [31]. The second-generation anticoagulants have longer half-lives, less drug-acquired resistance, and increased potency [19, 20, 31, 32]. Clinical signs of coagulopathy secondary to brodifacoum manifest 2 to 5 days after ingestion and vary based on the location and severity of the hemorrhage [20, 29, 31]. The most common are dyspnea, lethargy, coughing, hemoptysis, pale mucous membranes, and tachycardia [23, 24]. Bleeding typically occurs into body cavities causing hemothorax, hemoadenopathy, and retroperitoneal hemorrhage [20, 23]. Less frequent signs are melena, hematochezia, prolonged bleeding at injection sites, epistaxis, gingival bleeding, and neurologic signs [23, 24]. Case reports of atypical presentations of coagulopathy due to anticoagulant rodenticide include lameness from hematrhiosis [33], pericardial effusion [34], hematemesis [35], hydrenephrosis secondary to blood clots in the urinary bladder [36], tracheal obstruction [37], and submucosal gastric hemorrhage [38].

The earliest laboratory abnormality detected after second-generation anticoagulant rodenticide toxicity is elevation of the proteins induced by vitamin K absence or antagonism [31]. Prolongation of PT occurs within 36–72 hours of ingestion and precedes prolongation of PTT because of the short half-life of factor VII [20, 29]. Neither PTT nor activated clotting time is prolonged until greater than 72 hours after ingestion [20]. Decontamination of the patient is not indicated after signs of coagulopathy have developed since the toxin was consumed several days before examination. Patients with severe hemorrhage and subsequent anemia may require transfusions of whole blood or fresh frozen plasma to provide red blood cells and coagulation factors to halt bleeding. Oral vitamin K₁ at 1.25–2.5 mg/kg twice daily for 4 weeks is needed to resolve coagulopathy caused by second-generation anticoagulants [31]. Therapy is needed for only 2 weeks for first-generation anticoagulants and for 3 weeks with bromadiolone since these agents are shorter acting [20]. Absorption of oral vitamin K₁ is improved if given with a fatty meal [20]. A subcutaneous loading dose of 2.5–3.3 mg/kg of vitamin K₁ is recommended by some clinicians and discouraged by others [19, 20, 31, 39, 40]. Injections can cause additional bleeding and hemotoma formation and are not more bioavailable than the oral formulation [19]. Intravenous administration is not recommended due to the risk of anaphylaxis [39]. Treatment should be discontinued for 2 days before repeated testing of PT to ensure that the patient is able to produce active coagulation factors without supplementation. If PT is still prolonged after discontinuation of vitamin K₁, then supplementation is continued for another week and the patient is then retested [20].

This case is an unusual presentation of coagulopathy secondary to brodifacoum ingestion in that unilateral subconjunctival and retrobulbar hemorrhages were the only apparent clinical signs. Subconjunctival hemorrhage can occur because of scleral or conjunctival bleeding, or anterior migration of retrobulbar blood [21, 25]. Retrobulbar hemorrhage was presumed because of the exophthalmos and severe subconjunctival hemorrhage OS, though this was not confirmed with imaging or sample collection. While fresh frozen plasma is usually indicated for patients with active bleeding, it was not administered in this case since no other significant detectable body cavity effusion had occurred at the time of diagnosis and the patient was not anemic. Anemia is present in the majority of coagulopathic dogs after rodenticide ingestion and can be seen in 83% of cases [24]. Monitoring was continued for the first 24 hours of vitamin K₁ therapy so that fresh frozen plasma could have been administered if necessary. This patient had complete resolution of all clinical signs with oral vitamin K₁ supplementation, and no permanent adverse effects occurred to the globe or periocular structures OS. The mild prolongation of PTT seen in this patient at the conclusion of vitamin K₁ therapy was not indicative of an unresolved coagulopathy since the PT was normal. Mild elevations of PTT are often clinically insignificant, and PT is a more sensitive indicator of coagulopathy secondary to vitamin K deficiency [29].

As with other reports of atypical presentations of coagulopathy secondary to anticoagulant rodenticide ingestion, this case further reinforces that patients with this syndrome can present with bleeding in nearly any location. Prognosis is excellent as long as a prompt diagnosis is made and proper treatment instituted. An incorrect initial diagnosis occurs in up to 25% of cases of coagulopathy due to anticoagulant rodenticide toxicity, which can be life threatening because treatment is delayed and hemorrhage continues [24]. Patient history can also be misleading, as owners denied pet exposure to anticoagulant rodenticide in over 50% of confirmed cases in one study [41]. The massive unilateral subconjunctival hemorrhage in this case allowed for rapid diagnosis of the patient’s coagulopathy before more serious bleeding and anemia occurred. In summary, anticoagulant rodenticides should be strongly considered in cases of unilateral subconjunctival hemorrhage of the globe and nictitating membrane or when retrobulbar hemorrhage is suspected.

References


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