CASE REPORT

Non-traumatic Ocular and Periocular Hemorrhages in a Hypertensive Patient under Continuous Ambulatory Peritoneal Dialysis and Warfarin Therapy

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Abstract

We herein present the first reported case of severe proptosis caused by ocular and periocular hemorrhages in a continuous ambulatory peritoneal dialysis patient without previous history of trauma. The bleeding tendency caused by uremia and the use of warfarin during uncontrolled high blood pressure were most likely responsible for her ocular and periocular hemorrhages. Appropriate control of blood pressure and adequate self-care education are important for the prevention and treatment of any bleeding complications in uremic patients receiving both maintenance anticoagulation therapy and peritoneal dialysis.

Key words: non-traumatic ocular hemorrhage, periocular hemorrhage, sub-tenon hemorrhage, hypertension, continuous ambulatory peritoneal dialysis, warfarin

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Introduction

Ocular and periocular hemorrhages are extremely rare complications in patients undergoing regular dialysis, especially those without a history of surgery or trauma. In this report, we herein describe a case of non-traumatic ocular and periocular hemorrhages in a hypertensive patient under continuous ambulatory peritoneal dialysis and warfarin therapy.

Case Report

A 43-year-old anuric woman undergoing regular continuous ambulatory peritoneal dialysis (CAPD) for chronic glomerulonephritis-related end-stage renal disease was hospitalized due to the gradual onset of a protruding left eyeball with poor vision in her left eye for two weeks. She had chronic shortness of breath and dry cough resulting from severe anemia and fluid overload due to poor compliance to our recommendation of salt and water restriction. She denied any history of trauma to her left eye during the two weeks prior to this admission. She was taking warfarin (2.5 mg once daily) as a part of maintenance anticoagulation therapy after a mitral valve replacement for mitral valve regurgitation due to lower prothrombin time (INR) values (0.94-1.09) in the two months prior to this admission. However, her blood pressure was not adequately controlled for several weeks because of her poor compliance to anti-hypertensive agents and peritoneal dialysis. Her initial vital signs on admission were blood pressure 203/99 mmHg, heart rate 102/min, respiratory rate 20/min, and body temperature 36.8°C. Her laboratory data on admission included hemoglobin 5.9 g/dL, leukocyte 6,300/mm³, platelet 505,000/mm³, blood urea nitrogen 93 mg/dL, serum creatinine 13.48 mg/dL, calcium 9.3 mg/dL, phosphate 9.6 mg/dL, and serum β2-microglobulin 116.2 mg/L. Her indices of CAPD therapy were weekly Kt/V 2.05 and weekly creatinine clearance (WCC) 39.6 L/week/1.73 m² body surface area 2 months before this admission. The reason for inade-
Adequate dialysis was most likely her noncompliance to CAPD therapy. She was only exchanging 6-8 L dialysate per day despite of our prescription of dialysate exchange up to 10 L daily.

Contrast-enhanced computed tomography (CT) of the orbital cavity was then performed and showed several heterogeneous hematomas over retrobulbar, intraocular, sub-tenon, and subconjunctival areas of the left eyeball (Figure). Rupture of the ciliary artery was then highly suspected because it is the major source of blood supply to the sub-tenon area, thus leading to the simultaneous, very extensive hemorrhage in the ocular and periocular areas. An ophthalmologist was consulted but the visual loss of her left eye was irreversible because she had not sought medical assistance early enough. Surgical removal of the affected eyeball was suggested but she refused. Thus, only conservative treatment could be given for the treatment of her catastrophic visual loss. After admission, she exchanged 10 L dialysate daily according to our prescription, and her blood urea nitrogen was improved to 75 mg/dL, serum creatinine to 9.59 mg/dL, Kt/V to 2.22, and WCC to 49.9 L/week/1.73 m² body surface area on day 11 of admission. Her proptosis and intraocular hematoma resolved gradually in the following 4 weeks due to adequate CAPD therapy as well as better control of her blood pressure and judicious titration of the dose of anticoagulant therapy; however, her visual loss was irreversible. The patient was then discharged and her status remained unchanged for a year following routine follow-ups.

Discussion

To the best of our knowledge, this is the first reported case of severe proptosis caused by an ocular and periocular hemorrhages in a CAPD patient without previous history of trauma. Several cases of retrobulbar hemorrhage (1) or suprachoroidal hemorrhage (2) following an eye trauma or an optic surgery were reported in patients undergoing hemodialysis or peritoneal dialysis. However, it is rare for dialysis patients to suffer from ocular and periocular hemorrhages in the absence of surgery or trauma. Similarly, Tajika et al. also reported choroidal hemorrhaging without proptosis as a complication following an episode of cough in a patient with hypertension and chronic renal failure but without history of trauma or surgery (3). In our case, the bleeding tendency caused by uremia and the use of warfarin during uncontrolled high blood pressure were most likely responsible for her ocular and periocular hemorrhages. It has been reported that hemodialysis patients under warfarin treatment

Figure. (A) The transverse view of the orbital CT shows a retrobulbar hematoma (arrow); sub-tenon and subconjunctival hematoma (arrowhead); and intraocular hematoma (asterisk). (B) The transverse view of the orbital CT shows a posterior sub-tenon hematoma (arrow). (C) The coronal view of the orbital CT shows a sub-tenon and subconjunctival hematoma (arrowheads) and an intraocular hematoma (asterisk).
may have a higher risk for major bleeding episodes and inter-measurement INR variability compared with non-hemodialysis patients under warfarin treatment or hemodialysis patients without warfarin treatment (4). We infer that the gradual onset of her blurred vision and proptosis may possibly result from the slowly-developed blood leak and hematoma formation caused by her bleeding tendency, although a retrobulbar hematoma usually occurs abruptly in a traumatic patient.

Ocular hemorrhaging (this case) and intracranial hemorrhaging (ICH) in uremic patients may share some common mechanisms of major bleeding, including hypertension, higher level of uremic toxins due to inadequate dialysis, platelet dysfunction, and the use of anticoagulation. In addition to the previously mentioned common mechanisms for severe hemorrhaging in uremic patients, however, mitral valve regurgitations may increase the pulmonary venous pressure, thus leading to chronic cough associated with the simultaneously strenuous Valsalva maneuver. Elevated intraocular pressure caused by chronic kidney disease and chronic cough may also contribute to this complication (5, 6). Furthermore, pre-existent vascular risk factors, such as severe hypertension and hypercholesterolemia, may also result in endothelial dysfunction. These two specific mechanisms (i.e., a higher intraocular pressure and endothelial dysfunction) may also play important roles as predisposing risk factors in the development of the ocular and periocular hemorrhages in this case.

The spontaneous resolution of proptosis and the ocular and periocular hematoma in this patient may result from the absence of anticoagulation as well as the relatively stable hemodynamic condition during renal replacement therapy by peritoneal dialysis rather than by hemodialysis. This could be advantageous for the treatment of any bleeding complication in uremic patients under peritoneal dialysis. In summary, good control of blood pressure and adequate self-care education are important for the prevention and treatment of any bleeding complications including ocular or periocular hemorrhaging in uremic patients receiving both maintenance anticoagulation therapy and peritoneal dialysis, a choice for renal replacement therapy with a much lower bleeding risk than hemodialysis.

The authors state that they have no Conflict of Interest (COI).

References