Clinical Ophthalmology 2014:8 813–817
Clinical Ophthalmology downloaded from https://www.dovepress.com/ by 54.70.40.11 on 13-Jan-2019
For personal use only. Powered by TCPDF (www.tcpdf.org)

Cloudy corneas as an initial presentation of multiple myeloma

Priyanka Sharma1
Haifa A Madi1
Richard Bonshek2
Stephen J Morgan1
1Sunderland Eye Infirmary, Sunderland, UK; 2National Specialist Ophthalmic Pathology Service, Department of Histopathology, Manchester Royal Infirmary, Manchester, UK

Summary: We report a case of previously unsuspected myeloma, presenting with cornea verticillata due to intracorneal paraprotein deposition.

History: An 85-year-old female presented via her optician with a 4-month history of cloudy vision. She had undergone an uneventful bilateral phacoemulsification surgery 7 years earlier. Extensive spiraling corneal epithelial opacification was noted on slit-lamp examination. On further investigation, she was found to have a previously unsuspected low-grade multiple myeloma. We established the nature of the corneal deposits with corneal epithelial biopsy histopathology and electron microscopy. It is very rare for multiple myeloma to present in this fashion. Ophthalmologists should be aware that such a presentation may rarely be due to systemic multiple myeloma.

Keywords: corneal crystals, multiple myeloma, crystalline keratopathy, vortex keratopathy

Introduction

Corneal deposits of various types have been described in multiple myeloma, monoclonal gammapathy, and essential cryoglobulinemia.1,2 There are only a very few case reports of corneal immunoglobulin (Ig) deposition. We present one such case in which there was intraepithelial deposition of corneal IgG-kappa. In addition, it is noteworthy that the presentation to the ophthalmologist was the presenting complaint leading to the diagnosis of myeloma.

Case report

An 85-year-old lady with bilateral cloudy corneas was referred to ophthalmology as an outpatient from her optometrist. She gave a 3–4 month history of cloudy vision, mainly in her right eye. She had undergone an uneventful bilateral phacoemulsification surgery 7 years earlier.

Her visual acuity at presentation was 6/7.5 OU. On examination, she had marked grayish intraepithelial corneal opacities in a pattern of hazy spiraling lines in both eyes (Figure 1). The corneal stroma and endothelium had normal appearance; there was no evidence of corneal edema. Specular microscopy was not possible. The anterior chamber depth and contents, intraocular lenses, posterior capsule, and fundus examination were normal.

She had a past medical history of ischemic heart disease and osteopenia. Her current medications were aspirin, simvastatin, lisinopril, codeine, and paracetamol.

Systemic investigation revealed a raised serum IgG with a kappa paraprotein band (12.4 g/L) on serum protein electrophoresis (Figure 2). The erythrocyte sedimentation
rate was raised (49 mm/hour), and there was a mild kidney impairment with raised urea (8.1 mmol/L) and raised creatinine (118 µmol/L). The random blood glucose, electrolytes, liver function, lipid profile, and calcium profile were normal. Urinary Bence Jones proteins were elevated. There was no evidence of Fabry disease as the lysosomal enzymes were all found to be normal.

She was referred to the Department of Haematology, Sunderland Royal Hospital, Sunderland, UK, and a bone marrow biopsy was carried out that showed increased plasma cells (11%) with pink staining crystals in the cytoplasm. Free crystals were also seen. These findings were consistent with multiple myeloma.

A corneal epithelial biopsy was undertaken and subjected to further laboratory analysis. Immunohistochemistry of the corneal biopsy showed excessive amounts of kappa light chain staining, relative to lambda light chain staining. On transmission electron microscopy, there was evidence of intraepithelial intracellular and extracellular geometrically irregular hexagonal electron dense particles (Figure 3). These are typically found in crystalline keratopathy due to gammopathy. There was an absence of immunoactoid, a paraprotein also commonly present in these cases, but not detected in our case.

The patient was commenced on systemic chemotherapy with cyclophosphamide and dexamethasone. Six months later, there was significant improvement in corneal clarity (Figure 4).

Figure 1 Clinical photographs of the right cornea.
Notes: (A) Subepithelial deposits extending toward the corneal center by fingerlike projections; (B) at higher magnification, depicting the spiral-like pattern known as corneal verticillata.

Discussion

Corneal epithelial disturbance in a spiraling pattern is variously known as cornea verticillata, vortex keratopathy, and hurricane keratopathy. Well-recognized causes include: Fabry disease;^1^ drugs^4^ (amiodarone,^5^ chloroquine, hydroxychloroquine, indomethacin, chlorpromazine, tamoxifen, meperidine); toxicity of topical medications; and ocular surface failure secondary to chemical and thermal burns, conjunctival malignancy, and cicatrizing conjunctivitis.

In myeloma, an intracorneal accumulation of Ig is well-described. Patients may be asymptomatic or complain of decreased vision,^6^ irritation,^7^ photophobia,^8^ and erosions.^9,10^ The corneal deposition can be crystalline,^10–12^ amorphous,^6^ or a combination.

Other ocular surface manifestations of multiple myeloma include repeated subconjunctival hemorrhages,^13^ peripheral
Ig-kappa distribution can affect different corneal layers. It was reported to be deposited in: all layers of the cornea in four cases, only stromal involvement in three cases, only epithelial involvement in two cases, epithelium and stromal involvement in one case; and endothelium and Descemet’s membrane involvement in one case.

Corneal deposition of immunoglobulins in multiple myeloma is rarely in a vortex pattern. Two cases of vortex keratopathy in multiple myeloma have been previously reported. Auran et al described a case of crystalline keratopathy in a vortex pattern in a patient with multiple myeloma. However, they did not obtain any pathological material. Their patient had already undergone bilateral phacoemulsification 7 years previously and later developed an endocapsular hematoma that resolved without treatment. There were some similarities between Auran’s case and ours. Both cases had a clinical picture of vortex keratopathy and an Ig-kappa spike on serum electrophoresis. Auran’s case differed in that the patient had decreased vision and glare, while our patient only had decreased vision without glare or photophobia. Their patient had crystalline deposits in the epithelium and stroma, while our patient had deposition only in the epithelium.

Chong et al described the second case of vortex keratopathy in multiple myeloma. Their case was diagnosed with multiple myeloma 18 months after first presentation to the ophthalmologist. Again, they did not obtain a corneal biopsy. They did, however, take conjunctival specimens which stained monoclonally for IgG-kappa light chains. Conjunctival electron microscopy showed intracellular hexagonal crystalline structures – similar to our corneal biopsy picture. Their conjunctival specimen also showed some extracellular material in the form of microfibrils, which stained for kappa light chains. This was not evident in our case. Chong et al therefore assumed that the corneal crystalline vortex keratopathy was due to Ig deposition.

Buerk and Tu reported a patient with multiple myeloma crystalline keratopathy where confocal microscopy showed in vivo crystals in the corneal epithelium and the anterior stroma. They repeated confocal microscopy after 6 months of chemotherapy and demonstrated a decrease in the size and the number of the corneal crystals. Use of in vivo confocal microscopy was also used by Paladini et al in a case with bilateral crystalline corneal deposits, due to monoclonal gammopathy. Confocal microscopy has the advantage of being able to image corneal crystalline deposits in vivo and help monitor a decrease in these changes with chemotherapy more quantitatively. However, confocal microscopy is not a histological tissue test and, therefore, cannot confirm the type...
of material/Ig deposited. We did not have access to confocal microscopy at our center.

Our case is, therefore, the first published case of vortex keratopathy in multiple myeloma with corneal deposition of Ig confirmed by corneal epithelial biopsy.

It is very rare for multiple myeloma to present in this fashion. There are several causes of intracorneal crystalline deposits, and the differentiation is impossible clinically, but laboratory studies – specifically a serum protein electrophoresis – will help detect any unsuspected multiple myeloma.37 Ophthalmologists should be aware that such a presentation may rarely be due to systemic multiple myeloma.

Disclosure
The authors report no conflicts of interest in this work.

References
Cloudy corneas as initial presentation of multiple myeloma