Peters Anomaly

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Chief Complaint: Bilateral corneal clouding noted after birth

History of Present Illness

A 2-day-old, 1920 gram infant born at 36 weeks gestation via caesarean-section due to suspected intrauterine growth restriction was evaluated for bilateral corneal opacity noted at the time of delivery (Figure 1). There were no known maternal infections during the pregnancy and no problems during the perinatal period. The mother had a herpetic oral ulcer, but no genital lesions. The infant did not show evidence of tearing, blepharospasm, photophobia, drainage, or ocular redness.

Past Ocular History

- None
Past Medical History
- Intrauterine growth restriction (IUGR)

Medications
- None

Family History
- Maternal grandfather with retinal capillary hemangiomas secondary to von Hippel-Lindau syndrome

Social History
- None

Ocular Exam

Visual Acuity
- Unable to perform

<table>
<thead>
<tr>
<th></th>
<th>OD (right eye)</th>
<th>OS (left eye)</th>
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<tbody>
<tr>
<td><strong>Intraocular pressure</strong></td>
<td>16 mmHg</td>
<td>12 mmHg</td>
</tr>
<tr>
<td><strong>Corneal diameter</strong></td>
<td>8.5 mm</td>
<td>9.0 mm</td>
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<tr>
<td><strong>Corneal pachymetry</strong></td>
<td>676 microns</td>
<td>870 microns</td>
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<tr>
<td><strong>Retinoscopy</strong></td>
<td>Unable to perform</td>
<td>+2.50</td>
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*Fig. 2: Bilateral corneal opacities during examination under anesthesia. Note the more significant corneal opacification in the right eye compared to the left.*

Anterior segment examination
OD: Diffuse corneal opacification with iridocorneal adhesion temporally; minimal view of the anterior segment due to corneal opacity. 
OS: Round 5 mm corneal opacity decentered nasally, but involving central visual axis; inferonasal iridocorneal adhesion.

A-scan echography
- OD: Axial eye length, 15.99 mm; Anterior chamber depth, 1.3 mm
- OS: Axial eye length, 16.73 mm; Anterior chamber depth, 1.8 mm.

B-scan echography
- No evidence of mass or retinal detachment OU.

Dilated fundoscopic examination
- OD: No view.
- OS: Healthy-appearing optic nerve with cup-to-disc ratio 0.15; normal macula, vessels, and periphery.

High frequency ultrasound biomicroscopy
- OD: Narrow anterior chamber with multiple areas of peripheral adhesions and membranes between iris and cornea.
- OS: Deeper anterior chamber with trace iridocorneal membranes.

Laboratory studies
- TORCH titers within normal limits
- Herpes simplex virus (HSV) polymerase chain reaction (PCR) negative

Assessment
- Ocular findings were consistent with anterior segment dysgenesis, most likely Peters anomaly. There was no evidence of glaucoma in either eye.

Clinical Course
The decision was made to observe the left eye as the corneal opacification had improved at his 1 month visit and there was an excellent view to the posterior pole. However, to prevent amblyopia and provide visual rehabilitation a penetrating keratoplasty (PKP) of the right eye was recommended. The patient subsequently underwent synechiolysis of iridocorneal adhesions and PKP in the right eye. At his 4-month postoperative visit, the graft was clear and there was visual improvement OD. Genetic testing was also done for two autosomal-dominant genes for Peters anomaly, PITX2 and FOXC1 genes, but the results were negative.

Discussion
Clinical Features and Diagnosis
Peters anomaly is a rare eye malformation resulting in congenital corneal opacity and is part of a spectrum of developmental anomalies of the cornea, iris and lens termed "anterior segment dysgenesis."

Peters anomaly was first described in 1906 by Albert Peters as a central corneal leukoma with variable synechiae between the iris and cornea and a defect in Descemet's membrane and the corneal endothelium. Since then, Peters anomaly has been subdivided into three types: 1) Peters type I characterized by central corneal opacity with iridocorneal adhesions (Figure 3), 2) Peters anomaly type II with central corneal opacity and cataracts or corneolenticular adhesions, and 3) Peters-plus syndrome with Peters anomaly and short stature, developmental delay, dysmorphic facial features including cleft lip/palate along with cardiac and genital abnormalities [1, 2]. Peters anomaly has a wide spectrum of severity and corneal opacification is bilateral in approximately 80% of cases [3].

Patients with Peters anomaly are at increased risk of glaucoma, presumably due to abnormal development of the trabecular meshwork and Schlemm's canal and/or the presence of a shallow anterior chamber [1]. Approximately 50-70% of patients with Peters anomaly have associated glaucoma [3-5]. Glaucoma is more common in those with cataracts or corneolenticular adhesions. Elevated intraocular pressure (IOP) typically presents in infancy, but can also arise later in life.

The diagnosis of Peters anomaly is made clinically with the finding of corneal opacification with corneal edema and underlying loss of Descemet's membrane and endothelium. Specular microscopy may be used to assess the presence and structure of these corneal layers. Ultrasound biomicroscopy (UBM) or anterior segment optical coherence tomography (AS OCT) can be performed to aid in the identification of iridocorneal or corneolenticular adhesions [2, 6] (Figure 4).
Etiology and Pathophysiology

Most cases of Peters anomaly are sporadic without an identified genetic cause. It can occur as an isolated finding or associated with systemic disease as in Peters-plus syndrome. Abnormal formation of the anterior segment causes an incomplete separation of the cornea from the iris or lens leading to corneal opacity. In the inherited forms, mutations causing abnormal migration of neural crest cells to the posterior cornea have been linked to Peters anomaly, including PAX6, FOXC1, PITX2 and CYP1B1. Mutations in these genes can also cause other clinical conditions such as aniridia, Axenfeld-Rieger syndrome, and primary congenital glaucoma (see differential diagnosis below). Mutations in PAX6, FOXC1, and PITX2 follow an autosomal dominant inheritance pattern while mutations in CYP1B1 follow an autosomal recessive pattern [1, 2].

Management and Prognosis

Glaucoma associated with Peters anomaly is notoriously difficult to control and frequently requires surgical intervention. In a retrospective study, Yang et al. found that surgery was effective in controlling the IOP in only 32% of eyes at a median follow-up time of 11 years, even after multiple procedures [7]. There is little literature on the management of glaucoma associated with Peters anomaly and no head-to-head trials comparing the different surgical options. As a result, there is no established treatment algorithm for the condition. Based on the personal experiences of one of our senior glaucoma faculty members at the University of Iowa, patients are often initially managed medically. There are a variety of surgical options that can be attempted if medical management is insufficient. Trabeculotomy or goniotomy may be performed in eyes that have not undergone corneal surgery, though goniotomy can be difficult due to decreased visualization secondary to corneal opacification. In eyes that have already undergone corneal surgery, a glaucoma drainage device implant is often
change with surgery. However it is unclear which treatment modality is superior. Further studies are needed to compare the long-term outcomes between PKP and KPro implants in children.

**Fig. 5:** Post-operative appearance of a Boston Type I Keratoprosthesis (KPro-1) after multiple failed penetrating keratoplasties in a patient with Peters anomaly.

### Differential Diagnosis

- Peters anomaly
- Axenfeld-Rieger syndrome
- Sclerocornea
- Congenital hereditary endothelial dystrophy (CHED)
- Congenital hereditary stromal dystrophy (CHSD)
- Descemet's breaks from forceps delivery
- Congenital glaucoma
- Microbial keratitis (TORCH infections, herpes simplex virus)
- Metabolic disorders (mucopolysaccharidoses, sphingolipidoses, or mucolipidoses)

A useful mnemonic that is commonly used for the differential of a cloudy cornea at birth is **STUMPED:**

- Sclerocornea
- Tears in Descemet's membrane (forceps injury or Haab's striae)
- Ulcer
- Metabolic
- Peters anomaly
- Edema (congenital glaucoma or CHED)
References


Suggested Citation format


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the initial surgery of choice if IOP control is unsatisfactory with medical management. Trabeculectomy is usually not preferred at very young ages and cyclodestructive procedures are reserved as a last option for glaucoma control when the visual prognosis is poor.

Children with corneal opacification or cataract involving the visual axis are at risk of developing amblyopia. Penetrating keratoplasty (PKP) or cataract extraction may be indicated to reduce this risk (Video). Visual prognosis is dependent on the disease severity. Although there is no formal classification for disease severity in Peters anomaly, Chang et al. defined patients with severe disease as being characterized by dense central opacities covering more than half of the cornea, corneolenticular adhesions, or associated ophthalmic anomalies like microphthalmia, aniridia, or cataracts. These patients had a visual acuity of finger-counting or worse and had poorer visual outcomes even with PKP. Patients with mild disease characterized by corneal opacities covering less than half of the cornea or iridocorneal adhesions and visual acuities of 20/100 or better had better visual outcomes with treatment [2]. Post-operatively there is also a role for amblyopia therapy to improve visual outcomes [8].

Penetrating Keratoplasty (PKP) for Anterior Segment Dysgenesis (Pete...