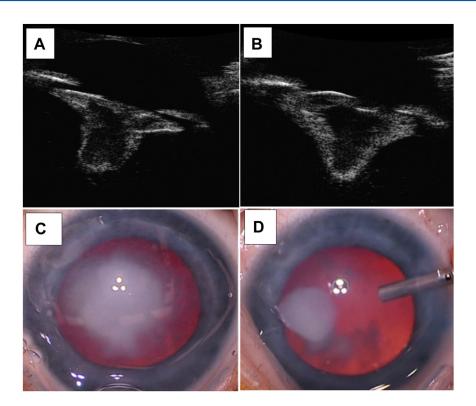
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Pictures & Perspectives



Bilateral Cataracts and Posterior Lentiglobus in USP9X Syndrome

A 3-month-old girl with mild brachycephaly and a broad nasal bridge was referred for cataract surgery. She had hearing loss in her left ear, an atrial septal defect, and a pulmonary venous anomaly. Initial examination showed blink-to-light vision without nystagmus or strabismus. Examination under anesthesia found bilateral cataracts with posterior lentiglobus (**A**, **B**, **C**). Infusion flushed the cataract out intraoperatively (**D**). Genetic testing showed a heterozygous variant (c.799_802del) in *USP9X*, on the X-chromosome. *USP9X* syndrome is a neurodevelopmental disorder predominantly found in female patients. Symptoms include intellectual disability, facial dysmorphia, language impairment, short stature, heart dysmorphia, hearing problems, abnormal skin pigmentation, and cataracts. Lentiglobus has not been previously described. (Magnified version of Figure **A-D** is available online at www.aaojournal.org).

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