

# Aniridia : A window to wilm's tumour

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## Abstract

We are reporting a case with aniridia which lead to the diagnosis of Wilm's tumour in a twelve month male child.

Keywords : aniridia, Wilm's tumour, vitA deficiency.

## Introduction

Aniridia in majority of the cases occurs isolated without any systemic involvement due to dominantly inherited mutations or deletions of the paired box gene-6 (*PAX6*) where as in minority of cases it occurs as part of the WAGR (Wilms tumor-aniridia-genital anomalies-retardation) syndrome caused by deletion of 11p13, which includes the Wilms' tumor (*WT1*) and aniridia gene (*PAX6*) loci<sup>1</sup>. The major diagnostic feature is congenital partial or complete hypoplasia of the iris; foveal hypoplasia with reduced visual acuity and early onset nystagmus. Other associated ocular abnormalities include cataract, glaucoma and corneal opacification and vascularization.<sup>2</sup> Our case was diagnosed with aniridia leading to diagnosis of Wilm's tumour along with vitamin A deficiency.

## Case Report

Twelve months old male child presented to us with history of photophobia and whitish lesion in right eye. On ophthalmological examination visual acuity was 20/30 at 38cms with Teller's acuity chart. Conjunctiva of right eye showed bitot's spot temporally and left eye showed

Subsequently child was referred to nephrologist and was diagnosed with Wilm's tumour. He underwent partial nephrectomy surgery and is on regular cycle of chemotherapy (adrimycin). Child was started immediately on vitamin A 1,00,000 IU I.M, subsequent next day and 4 weeks. Follow up after one month showed resolution of bitot's spot and conjunctival xerosis.

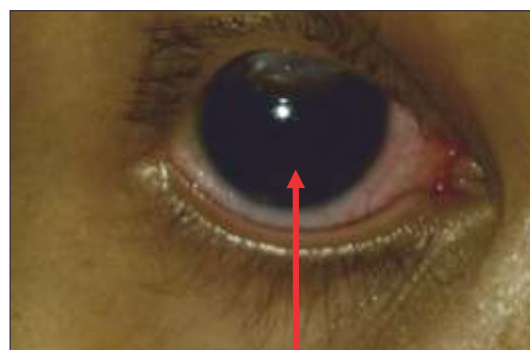


Fig 1: Shows the right eye with aniridia.

## Discussion

Aniridia is seen in approximately 1.8/100 000 live births.<sup>3</sup> The incidence ranges from 1:40000 to 1:100000. No significant racial or gender predilection has been described.<sup>4</sup> The frequency of Wilms' tumour in the general population is one in 10,000 to 50,000 individuals. Fraumeni and Glass found seven Wilms' tumour patients among 28 cases of aniridia which indicates that Wilms' tumour is more frequent in patients with aniridia than in general population.<sup>5</sup> WAGR syndrome, characterized by the clinical association of Wilms' tumors with aniridia, ambiguous genitalia, genitourinary anomalies, and mental

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conjunctival xerosis. Aniridia in both eyes with clear lens (Figure 1). Bilateral fundus was normal. There was no family history of aniridia or renal tumors. No delay in language and gross-motor development.

retardation, which was first described by Miller et al.<sup>6</sup>The underlying defect of Aniridia-Wilms' tumour was found to be a small interstitial deletion of chromosome band 11 p 13 . In patients with WAGR syndrome, the risk for developing a Wilms' tumor has been estimated to be up to 45%.<sup>7</sup> When associated with aniridia, a Wilms' tumor is diagnosed before age 5 years in 80% of cases.<sup>8</sup> In a study of 28 patients with non-familial bilateral aniridia Wilms' tumour syndrome, mental retardation along with microcephaly and craniofacial dysmorphism was observed in 71%, and cataract and glaucoma in 78% of cases .<sup>9</sup> optic nerve hypoplasia resulting from inactivation of the aniridia gene Pax6, which lies telomeric of *WT1* on chromosome 11p13.<sup>10</sup> In our case aniridia with wilm's tumour was associated with

ocular signs of vitamin A deficiency such as bitot's spot and conjunctival xerosis.

Therefore, renal ultrasound is recommended every 3-6 months until approximately 5 years of age .<sup>5</sup> Boys are often born with genital abnormalities, such as cryptorchidism or hypospadias, but more rarely ambiguous genitalia. However, genitourinary anomalies are not always present as in our case where no genital malformation was seen. Mental retardation in WAGR ranges from borderline-to-severe. However, individuals with normal intelligence have been reported <sup>11</sup>which was similarly seen our case.All the patients presenting with aniridia must be screened for wilm'stumour.

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