Color Vision in Aniridia

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PURPOSE. To assess color vision and its association with retinal structure in persons with congenital aniridia.

METHODS. We included 36 persons with congenital aniridia (10–66 years), and 52 healthy, normal trichromatic controls (10–74 years) in the study. Color vision was assessed with Hardy-Rand-Rittler (HRR) pseudo-isochromatic plates (4th ed., 2002); Cambridge Color Test and a low-vision version of the Color Assessment and Diagnosis test (CAD-LV). Cone-opsin genes were analyzed to confirm normal versus congenital color vision deficiencies. Visual acuity and ocular media opacities were assessed. The central 30° of both eyes were imaged with the Heidelberg Spectralis OCT2 to grade the severity of foveal hypoplasia (FH, normal to complete: 0–4).

RESULTS. Five participants with aniridia had cone opsin genes conferring deutan color vision deficiency and were excluded from further analysis. Of the 31 with aniridia and normal opsin genes, 11 made two or more red-green (RG) errors on HRR, four of whom also made yellow-blue (YB) errors; one made YB errors only. A total of 19 participants had higher CAD-LV RG thresholds, of which eight also had higher CAD-LV YB thresholds, than normal controls. In aniridia, the thresholds were higher along the RG than the YB axis, and those with a complete FH had significantly higher RG thresholds than those with mild FH (P = 0.038). Additional increase in YB threshold was associated with secondary ocular pathology.

CONCLUSIONS. Arrested foveal formation and associated alterations in retinal processing are likely to be the primary reason for impaired red-green color vision in aniridia.

Keywords: aniridia, color vision, foveal hypoplasia, retinal development

The reported prevalence of congenital aniridia in Norway and Sweden is about 1:72,000. Characteristic features in aniridia include absence or hypoplasia of the iris, foveal hypoplasia (FH) and nystagmus, while optic nerve hypoplasia (ONH) occurs but is less common. Aniridia may lead to severe visual impairment, although there is considerable phenotypic variation between and within families. Secondary progressive ocular complications, such as aniridia associated keratopathy (AAK), cataract, and glaucoma, are common from adolescence and onward.

Aniridia-like phenotypes may be caused by mutations in genes such as FOXC1 and PITX2, but typically (>85%) it is caused by loss of one functional copy of the PAX6 gene, with about one-third of cases being sporadic and two-thirds being inherited as an autosomal-dominant trait. The PAX6 gene is located on chromosome band 11p13 and regulates transcription of other genes important for ocular development. Gene dysfunction may affect multiple ocular structures. Aniridia phenotypes caused by PAX6 mutations are associated with different degrees of foveal hypoplasia, most commonly to a degree of no foveal avascular zone (FAZ) and no foveal depression. It is known that Pax6 acts on several target genes required for retinal ganglion cell development and retina neurogenesis. PAX6/Pax6 dosage expression varies during normal development, and animal studies show that abnormal PAX6/Pax6 expression affects the distribution, development, and the balance of different types of retinal cells.

Normal foveal development depends on a high ganglion cell count within the central retina and is characterized by formation of a FAZ before the foveal depression is formed, displacement of the inner retinal layers and postnatal elongation and migration of cones toward the center of the fovea. Trichromatic color vision requires a normally developed healthy functioning retina containing cone photoreceptors with three different opsins (L, M, and S cones). During human fetal retinal development, S cones appear earlier than L and M cones, with S cones present around the start of FAZ development and before the foveal depression develops. Disruptions in foveal development that occur around this time may therefore affect L and M cone development, density and its associated retinal circuitry, limiting the number of bipolar and horizontal cell contacts needed for normal red-green vision, more so than for blue-yellow vision. Thus, it is
reasonable to hypothesize that foveal hypoplasia associated with aniridia could affect red-green color discrimination.

Here, results are reported from experiments using computerized color vision tests together with retinal imaging (OCT and color fundus photography) to examine color vision and retinal layer structure in persons with congenital aniridia. These experiments provide insight into the association between the degree of arrested foveal formation in aniridia and impairment of red-green color vision and how this is accompanied by impairment of yellow-blue color vision when secondary pathology is advanced.

**METHODS**

**Participants**
We recruited 36 participants previously diagnosed with congenital aniridia (15 males), aged between 9 and 72 years, and 52 healthy, normal trichromatic controls (21 males), aged between 10 and 74 years, to the study. Participants with aniridia were recruited through the Norwegian Association of Aniridia, whereas normal controls were recruited through the National Centre for Optics, Vision and Eye Care, University College of Southeast Norway. The study was conducted in accordance with the tenets of the Declaration of Helsinki and was approved by the Regional Committee for Medical and Health Research Ethics (Southern Norway Regional Health Authority). All participants and/or their guardians signed informed consent after full explanation of the study’s purpose and procedures.

**Clinical Assessment**

The participants underwent an eye examination including subjective refraction, slit-lamp biomicroscopy, and color fundus photography (Topcon TRC-NW6S nonmydriatic fundus camera; Topcon Corp., Tokyo, Japan). Monocular visual acuity (logMAR) was measured with a digital high-contrast chart at 6 m (TestChart 2000; Thomson Software Solutions, London, UK). The test distance was reduced to 3 or 1 m if a reliable measurement could not be obtained at the longer distance. Color vision was screened binocularly with the Hardy-Rand-Rittler 4th edition (HRR; Richmond Products, Albuquerque, NM, USA) pseudochromatic plates using previously described methods. Those who made two or more errors on red-green (RG) screening plates on the second sitting were tested with the RG diagnostic plates and classified to have mild, medium, or strong RG deficiency depending on errors made on these plates. Those who made one or more errors on the yellow-blue (YB) screening plates on the second sitting were classified as mild, and if errors were made on the YB diagnostic plates, they were classified to have medium or strong YB deficiency depending on errors made on these plates. The clarity of the lens was evaluated using the Lens Opacities Classification System III (LOCS III). AAK was graded 0 to 3 where grade 0 indicated that the cornea was not affected; stage 1 indicated a partially affected corneal limbus; stage 2 indicated a totally affected limbus, without central corneal involvement; and stage 3 indicated a totally affected limbus with central corneal opacification. ONH was evaluated as present or not based on funduscopic, estimated by the ratio of disc-to-foveal distance and disc diameter (average of horizontal and vertical), and confirmed by measurements on color fundus photographs. ONH was defined as present when the ratio was >3.5, meaning that >3.5 optic discs could be apposed between the expected foveal center and the border of the optic disc. Borderline cases were also evaluated based on appearance of the optic disc, and ONH was defined as present if the optic nerve head was obviously small or if the disc showed a typical double ring sign. Glaucoma was noted as present or not based on previous ocular history and treatment.

**Foveal Hypoplasia**

Foveal hypoplasia was assessed by analyzing structural alterations on spectral domain optical coherence tomography (SD-OCT) images acquired with an OCT2 device (Spectralis; Heidelberg Engineering GmbH, Heidelberg, Germany). Volumetric scans, either 20° × 20° or 30° × 10° (consisting of 49 B-scans and 512–1536 A-scans/B-scan) were centered at the expected foveal center. Between 5 and 20 B-scans (frames) were averaged during acquisition to improve signal-to-noise ratio and compensate for eye motion (TruTrack; Heidelberg Engineering GmbH). Horizontal line scans, with a nominal scan length of 30°, were obtained in eyes where macular volumes could not be obtained. Multiple scans were acquired in the region of the expected foveal location to look for signs of foveal specialization. The lateral scale of all OCT scans was corrected for individual retinal magnification factor by multiplying the nominal scan length with the ratio between each individual’s axial length, obtained with the IOL Master (Carl Zeiss Meditec AG, Jena, Germany), and the OCT default axial length (24 mm).

Horizontal line scans were segmented at the inner limiting membrane (ILM); posterior boundary of the outer plexiform layer (OPL); center of the external limiting membrane (EJM); center of the ellipsoid zone (EZ); center of the interdigitation zone (IZ); and the retinal pigment epithelium-Bruch’s Membrane (RPE-BrM) band, using custom software implementing a method similar to that used by Park et al. The foveal center was defined as the section with the minimal foveal thickness (ILM to RPE-BrM) within the foveal depression. When no pit was present, the maximum widening of the outer nuclear layer (OPL to ELM) and/or lengthening of the photoreceptor outer segments (EZ to IZ) was used to identify the expected foveal center. Individual OCT line scans through the foveal center were used to grade foveal hypoplasia in participants with aniridia following the criteria suggested by Thomas et al:

- Absence of extrusion of plexiform layers (grade 1);
- Grade 1 plus absence of foveal depression (grade 2);
- Grade 2 plus absence of outer segment lengthening (grade 3);
- Grade 3 plus absence of outer nuclear layer widening (grade 4).

**Color Vision**

Color vision was examined with a low vision version of the Color Assessment and Diagnosis (CAD-LV) test (Color Assessment and Diagnosis Test; Cambridge Research Systems Ltd., Cambridge, UK). The CAD test has been used to examine both congenital and acquired color vision deficiencies and is validated with a reported high test-retest reliability. To ascertain the degree of noncongenital deficiencies and elucidate if it was receptorial or postreceptorial in origin, some participants were invited to do the Cambridge Color Test (CCT; Cambridge Research Systems Ltd., Cambridge, UK), and anomaloscope (HMC Oculos Anomalouscope MR, Type 47700; Oculos Optikgeräte GmbH, Wetzlar, Germany). All participants wore the appropriate refractive correction as determined during the initial clinical assessment.

**Color Assessment and Diagnosis Test**

The normal controls and 31 of the participants with aniridia were tested with the CAD. The CAD test measures red-green (RG) and yellow-blue (YB) color detection thresholds with a chromatic, moving isoluminant stimulus embedded in a
Chromatic thresholds were measured using a 4-alternative-forced-choice method along 16 hue directions in the CIE 1931 (x, y) chromaticity diagram. The test conditions were altered to compensate for low vision (LV) in aniridia—the stimulus was double its default size and moved slower to compensate for reduced visual acuity. That is, the stationary achromatic square array of 15 × 15 checkers subtended 5.7° × 5.7° in visual angle and the smaller moving chromatic square array of 5 × 5 checkers subtended 1.8° × 1.8° visual angle at a distance of 140 cm. The temporal frequency was 50% lower than the default setting, and random luminance modulation was increased by 100% to mask the detection of rod-mediated signals that might be introduced by the larger stimulus. Background luminance was 24 cd/m². The monitor (SpectraView PA241W; NED Display Solutions, Itasca, IL, USA) was calibrated daily using a photometer (Gossen Mavo Monitor USB, CO Ltd., Nürnberg, Germany).

Median RG and YB thresholds, represented as chromatic difference (ΔE) from the background chromaticity (x = 0.305, y = 0.325), were computed. The test was performed binocularly to minimize nystagmus and to indicate overall functional performance, and took 10–12 minutes to complete. As a control, RG and YB thresholds were also measured at a lower luminance level (2.4 cd/m²; a 1.0 spectrally calibrated neutral density filter was added in front of each eye) for a subset of normal controls (n = 38, age 10–67 years).

Cambridge Color Test

The normal controls and 16 of the participants with aniridia (age 11–66 years, logMAR 0.00–0.90) were tested with the CCT Trivector test following standard procedures. The test was performed binocularly and took 3 to 4 minutes to complete. Participants were tested twice, and average thresholds were used for analysis. The CCT is a computer-based pseudo-isochromatic test and the stimuli were generated via a graphics system (VSG ViSaGe; Cambridge Research Systems Ltd., Rochester, UK) and presented on a 22-inch CRT monitor (LaCie Electron 22blue; LaCie Group, Paris, France). The luminance and chromaticity of the monitor were checked daily with a colorimeter (PR650 Spectra; Photo Research, Inc., Chatsworth, MA, USA). The target was a Landolt-C with inner and outer diameters of 2.2° and 4.3°, respectively, and a gap size of 1° visual angle presented at a test distance of 305 cm. The gap position varied randomly from trial to trial (up, down, left, right), and the observer’s task was to indicate the position of the gap by pushing the correct button on a response box within 5 seconds. The test employed a staircase method (11 reversals) for measuring color-discrimination thresholds in three directions along the protan, deutan, and tritan confusion axes in the CIE 1976 (u’, v’) chromaticity diagram. The maximum color-vector length was set to 0.1600 units. The mean of the last six reversals was taken as the threshold.

Rayleigh Color Match

Ten participants with aniridia performed the Rayleigh color match (Oculus Optikgeräte GmbH) using their best eye. The Rayleigh stimulus is a 2° circular field divided in two semicircles. The upper part is the color-mixture field with a mixture of green (545 nm) and red (671 nm) light, and the lower part is the reference field with yellow light (589 nm). Using the procedure suggested by Linksz, the participant’s first task was to find the metameric match between the upper and lower fields by adjusting the luminance of the monochromatic yellow reference field and the relative amounts of red and green light in the mixture field. Theoretically, this is the match that produces the same set of quantum catches at the level of the photoreceptors. This first match was used as the starting point for finding the matching range. Several different relative amounts of red and green light in the mixture field were set by the operator using a staircase procedure, and the participant’s task was to judge whether each of them appeared uniform or not by adjusting the luminance of the reference field. The matching range was taken as the difference between the highest and the lowest red/green ratio that the participant would accept after making his/her own luminance adjustments. The mean match midpoint ± SD and matching range for color normal observers for this particular instrument is 40.3 ± 1.91 and 2.01 ± 1.09. A person with a protan deficiency requires more red to match the yellow reference field resulting in a higher than normal match midpoint, whereas someone with a deutan deficiency requires more green resulting in a lower than normal match midpoint. Larger than normal matching ranges signify poorer discrimination, with dichromats accepting all mixture ratios as metameric if the luminance is adjusted appropriately.

Genetic Analysis

All participants, with the exception of participants 5119 and 5124, gave saliva samples (Oragene-DNA, OG-500, DNA Self-Collection Kit; DNA Genotek, Inc., Ottawa, ON, Canada) for genetic analysis of cone opsins gene mutations known to be associated with congenital color vision deficiencies. DNA was also used in the PCR to amplify exons 2 through 5 of the L and M opsins separately, and exons 2, 3, and 4 were directly sequenced. To identify/confirm the genetic cause of aniridia for each participant, the PAX6 gene was amplified and sequenced using PCR primers and conditions that were described previously. Fluorescent DNA sequencing was performed on both DNA strands. For participants who were negative for PAX6 mutations, exons and intron/exon junctions of the PITX2 and FOXC1 genes were amplified and sequenced using the PCR primers described by Ansari et al.

Data Analysis

Statistical analysis were performed with R (v3.3.2), R Foundation for Statistical Computing, Vienna, Austria. The data for the aniridia group was found to be nonnormally distributed, as verified by histograms and q-q plots. The nonparametric Mann-Whitney U test was applied for independent samples and the Wilcoxon signed-rank test for paired samples. Correlations were assessed with Spearman correlation coefficients (r). Between-group differences were examined with Kruskal-Wallis 1-way ANOVA. The significance level was set to P ≤ 0.05. Bonferroni-corrected P values are reported for multiple comparisons. Bland-Altman plots were used to compare the CAD and CCT measurements with the nonparametric limits of agreement estimated as 2.5 and 97.5 quantiles of the differences and the average bias estimated as the median of the differences. The statistical normal limits for the CAD-LV test were computed based on the median color discrimination threshold and the 2.5% quantile and 97.5% quantile (giving a 95% interquantile range) for the normal controls in the study. A quantile regression was conducted to estimate and plot a curve to the median, 2.5% and 97.5% quantiles of the differences and the average bias estimated as the median of the differences. The statistical normal limits for the CAD-LV test were computed based on the median color discrimination threshold and the 2.5% quantile and 97.5% quantile (giving a 95% interquantile range) for the normal controls in the study. A quantile regression was conducted to estimate and plot a curve to the median, 2.5% and 97.5% quantile (giving a 95% interquantile range) for the normal controls in the study. A quantile regression was conducted to estimate and plot a curve to the median, 2.5% and 97.5% quantile (giving a 95% interquantile range) for the normal controls in the study. A quantile regression was conducted to estimate and plot a curve to the median, 2.5% and 97.5% quantile (giving a 95% interquantile range) for the normal controls in the study. A quantile regression was conducted to estimate and plot a curve to the median, 2.5% and 97.5% quantile (giving a 95% interquantile range) for the normal controls in the study. A quantile regression was conducted to estimate and plot a curve to the median, 2.5% and 97.5% quantile (giving a 95% interquantile range) for the normal controls in the study. A quantile regression was conducted to estimate and plot a curve to the median, 2.5% and 97.5% quantile (giving a 95% interquantile range) for the normal controls in the study.
our 52 controls), respectively. Both thresholds represented as CD \times 10^4 and thresholds expressed as CAD-LV units were used for analysis.

RESULTS

Clinical Assessment and Genetics

A total of 31 of 36 participants with aniridia (aged 10–67 years, 12 males) had cone opsin genes known to be associated with normal color vision, including four female carriers of deutan color vision deficiency. Five males with aniridia were excluded from further analysis as they were confirmed to have cone opsin genes conferring congenital deutan color vision deficiency. Table 1 shows the age distribution for all the study participants.

Those with normal opsin genes had corrected visual acuity 0.00 to 1.76 logMAR in their best eye; 11 (35.5%) of these made two or more RG errors on the HRR, four (12.9%) of which also made YB errors, while one (3.2%) made YB errors only. Summary of clinical findings and color vision test results for the participants with aniridia are presented in Tables 2 and 3. The participants who were negative for \( PAX6 \) mutations, were also negative for disease-causing mutations in both the \( PITX2 \) and \( FOXC1 \) genes.

All 52 normal controls had cone opsin genes known to be associated with normal color vision, including five female carriers of deutan color vision deficiency. None of the normal controls made more than one error on the HRR; only two made one error on plate 7 (the most difficult RG screening plate). Their corrected visual acuity was 0.16 logMAR or better in their dominant eye. The controls were healthy with no known systemic disease or ocular abnormalities. Clinical assessment, color fundus photography, and OCT imaging were performed on each participant, and all controls were found to be healthy and free of eye disease. Nuclear opalescence, nuclear color grade, cortical- and posterior subcapsular cataract were graded

#### Table 1. Age Distribution and Number of Participants in Each Age Group

| Age Group | Aniridia n | Median Age | Age Range | Normal Controls n | Median Age | Age Range |
|-----------|------------|------------|-----------|------------------|------------|-----------|
| <20       | 7          | 12         | 9–19      | 11               | 14         | 10–19     |
| 20–29     | 9          | 24         | 20–29     | 15               | 25         | 20–27     |
| 30–49     | 9          | 40         | 31–49     | 11               | 41         | 32–47     |
| 50–59     | 3          | 51         | 50–56     | 5                | 55         | 51–58     |
| >60       | 3          | 66         | 64–67     | 10               | 67.5       | 62–74     |

#### Table 2. Summary of Clinical Findings for the Participants With Aniridia

| ID   | Age | Genetic Mutation | Visual Acuity, logMAR | Nystagmus | Aniridia Associated Keratopathy, Grade | Foveal Hypoplasia, Grade | Optic Nerve Hypoplasia | Glaucoma | Lens Status |
|------|-----|------------------|-----------------------|-----------|---------------------------------------|--------------------------|------------------------|-----------|-------------|
| 5124 | 23  | N/A              | 0.40                  | N         | 0                                     | 0                        | N                      | N         | Phakic      |
| 5139 | 15  | Unknown          | 0.00                  | N         | 0                                     | 0                        | N                      | N         | Phakic      |
| 5132 | 64  | Unknown          | 0.56                  | N         | 0                                     | 1                        | N                      | Y         | Pseudophakic|
| 5134 | 49  | Unknown          | 0.18                  | N         | 1                                     | 1                        | N                      | N         | Pseudophakic|
| 5116 | 66  | PAX6             | 0.40                  | N         | 2                                     | 2                        | N                      | N         | Pseudophakic|
| 5120 | 42  | PAX6             | 0.22                  | N         | 1                                     | 2                        | N                      | N         | Phakic      |
| 5123 | 24  | PAX6             | 0.50                  | N         | 2                                     | 2                        | N                      | N         | Phakic      |
| 5154 | 24  | Unknown          | 0.72                  | Y         | 1                                     | 2                        | N                      | N         | Phakic      |
| 5114 | 56  | PAX6             | 0.86                  | Y         | 1                                     | 3                        | Y                      | Y         | Aphakic     |
| 5125 | 15  | PAX6             | 0.74                  | Y         | 1                                     | 3                        | N                      | Y         | Phakic      |
| 5126 | 12  | PAX6             | 0.80                  | Y         | 1                                     | 3                        | N                      | Y         | Phakic      |
| 5135 | 40  | PAX6             | 0.70                  | Y         | 1                                     | 3                        | Y                      | Y         | Phakic      |
| 5137 | 20  | PAX6             | 0.70                  | Y         | 1                                     | 3                        | N                      | N         | Phakic      |
| 5144 | 52  | PAX6             | 0.74                  | Y         | 2                                     | 3                        | N                      | Y         | Pseudophakic|
| 5147 | 11  | Unknown          | 0.50                  | Y         | 2                                     | 3                        | N                      | N         | Phakic      |
| 5148 | 49  | PAX6             | 0.60                  | N         | 2                                     | 3                        | N                      | Y         | Pseudophakic|
| 5113 | 32  | PAX6             | 0.80                  | Y         | 2                                     | 4                        | N                      | N         | Phakic      |
| 5117 | 20  | Unknown          | 1.00                  | Y         | 2                                     | 4                        | N                      | Y         | Pseudophakic|
| 5119 | 9   | N/A              | 1.00                  | Y         | 1                                     | 4                        | N                      | Y         | Phakic      |
| 5127 | 41  | PAX6             | 1.20                  | Y         | 2                                     | 4                        | N                      | Y         | Pseudophakic|
| 5131 | 29  | PAX6             | 1.30                  | Y         | 1                                     | 4                        | Y                      | Y         | Aphakic     |
| 5138 | 11  | PAX6             | 0.90                  | Y         | 1                                     | 4                        | N                      | N         | Phakic      |
| 5140 | 19  | PAX6             | 0.70                  | Y         | 2                                     | 4                        | N                      | N         | Phakic      |
| 5141 | 23  | PAX6             | 1.00                  | Y         | 2                                     | 4                        | N                      | N         | Phakic      |
| 5149 | 31  | PAX6             | 0.90                  | Y         | 2                                     | 4                        | N/A                    | N         | Phakic      |
| 5110 | 50  | PAX6             | 1.00                  | Y         | 3                                     | 4                        | N/A                    | Y         | Pseudophakic|
| 5118 | 25  | PAX6             | 0.74                  | Y         | 3                                     | 4                        | N/A                    | N         | Phakic      |
| 5121 | 51  | PAX6             | CF                    | Y         | 3                                     | N/A                     | N                      | N         | Phakic      |
| 5129 | 67  | PAX6             | 1.30                  | Y         | 3                                     | N/A                     | N                      | Y         | Aphakic     |
| 5145 | 36  | PAX6             | 1.76                  | Y         | 3                                     | N/A                     | N                      | N         | Pseudophakic|
| 5152 | 26  | Unknown          | 1.30                  | Y         | 3                                     | N/A                     | N                      | Y         | Phakic      |

CF, counting fingers at 0.5 m; N, no; Y, yes; N/A, not applicable.

* The participants are ordered by the grade of foveal hypoplasia.

† Measured with their best eye.
and found to be lower than NO2, NC4, C2, and P0 (LOCS III), respectively. None of the normal controls had undergone cataract surgery.

**Color Vision**

Figure 1 shows RG and YB CAD-LV thresholds for 27 participants with aniridia (circles) and 52 normal controls (squares) as a function of age. Four participants with aniridia were unable to complete the CAD-LV test, either because of too poor vision \((n = 2)\) or of other reasons \((n = 2; one child and one with cerebral palsy)\).

RG and YB CAD-LV thresholds were significantly higher for those with aniridia compared with age-matched normal controls (Mann-Whitney \(U\) test: RG: \(P < 0.001\), YB: \(P = 0.002\)). A total of 19 participants with aniridia had RG thresholds that were higher than the 97.5% quantile that was defined as the upper normal limit based our normal control data, of whom seven also had YB thresholds higher than the 97.5% quantile. None had higher YB threshold than the upper normal limit only. Figure 2 shows CAD-LV YB thresholds as a function of RG thresholds; both calculated as CAD-LV standard units. CAD-LV RG thresholds were significantly higher than the CAD-LV YB thresholds \((Z = 3.1, P = 0.001)\) in aniridia (Fig. 2).

CCT RG threshold was calculated as the mean of the protan and deutan thresholds. There was good agreement between the CAD-LV and the CCT test, but both RG and YB median CAD-LV were higher than the median CCT thresholds. The median RG difference (CAD-CCT threshold) was 12.2 and 19.7 for normal controls and participants with aniridia, respectively (Fig. 3). The median YB differences in thresholds were 69.5 and 28.9, respectively. However, for three of the participants with aniridia, the CCT tritan thresholds were more than twice as high as their CAD-LV YB threshold.

Ten of those with aniridia agreed to be measured with Rayleigh anomaloscopy. The median matching midpoint was 43.3 (range, 40.2–46.2), with a median matching range of 5.6 (range, 4.0–11.4).

| ID | Age | RG (CD ×10^4) | YB (CD ×10^4) | Protan | Deutan | Tritan | Midpoint | Range | RG | YB |
|----|-----|---------------|---------------|--------|--------|--------|----------|-------|----|----|
| 5124 | 23 | N/A | N/A | – | – | – | – | – | Normal | Normal |
| 5139 | 15 | 84 | 134 | 50 | 64 | 62 | – | – | Normal | Normal |
| 5132 | 64 | 183 | 327 | 296 | 328 | 1186 | 46.2 | 6.0 | Normal | Normal |
| 5134 | 49 | 93 | 193 | – | – | – | – | – | Normal | Normal |
| 5116 | 66 | 116 | 170 | 146 | 131 | 133 | 45.7 | 4.9 | Normal | Normal |
| 5120 | 42 | 50 | 113 | 84 | 81 | 92 | 41.0 | 5.1 | Normal | Normal |
| 5123 | 24 | 118 | 221 | 79 | 86 | 168 | 40.2 | 4.2 | Normal | Normal |
| 5154 | 24 | 110 | 167 | 41 | 54 | 72 | – | – | Normal | Normal |
| 5114 | 56 | 434 | 1198 | 320 | 407 | 1398 | – | – | Medium | Mild |
| 5125 | 15 | 196 | 317 | 73 | 74 | 103 | – | – | Normal | Normal |
| 5126 | 12 | 94 | 159 | 130 | 121 | 117 | 42.6 | 4.3 | Normal | Normal |
| 5135 | 40 | 123 | 187 | 153 | 218 | 286 | 42.2 | 6.6 | Normal | Normal |
| 5137 | 20 | 93 | 166 | 136 | 142 | 737 | 44.9 | 4.0 | Normal | Normal |
| 5144 | 32 | 104 | 144 | – | – | – | Normal | Normal |
| 5147 | 11 | 143 | 216 | 132 | 128 | 210 | 44.4 | 7.4 | Normal | Normal |
| 5148 | 49 | 192 | 497 | 96 | 149 | 1076 | 44.1 | 11.4 | Mild | Mild |
| 5113 | 32 | 154 | 177 | 128 | 120 | 115 | – | – | Medium | Normal |
| 5117 | 20 | 186 | 251 | – | – | – | – | – | Normal | Normal |
| 5119 | 9 | N/A | N/A | – | – | – | – | – | Normal | Normal |
| 5127 | 41 | 249 | 210 | – | – | – | – | – | Medium | Normal |
| 5131 | 29 | 197 | 154 | – | – | – | – | – | Medium | Normal |
| 5138 | 11 | 80 | 141 | 182 | 157 | 192 | – | – | Normal | Normal |
| 5140 | 19 | 133 | 213 | – | – | – | – | – | Normal | Normal |
| 5141 | 23 | 231 | 238 | – | – | – | – | – | Normal | Normal |
| 5149 | 31 | 181 | 277 | – | – | – | – | – | Normal | Normal |
| 5110 | 50 | 119 | 213 | – | – | – | – | – | Normal | Normal |
| 5118 | 25 | 220 | 848 | 141 | 154 | 739 | 40.5 | 7.8 | Normal | Mild |
| 5121 | 51 | N/A | N/A | – | – | – | – | – | Mild | Strong |
| 5129 | 67 | 621 | 1292 | – | – | – | – | – | Medium | Mild |
| 5145 | 36 | 124 | 196 | – | – | – | – | – | Mild | Normal |
| 5152 | 26 | N/A | N/A | – | – | – | – | – | Medium | Normal |

N/A, not applicable; RG, red-green; YB, yellow-blue.

* The participants are ordered by the grade of foveal hypoplasia (as in Table 2).
Foveal Hypoplasia and Color Vision in Aniridia

Figure 4 shows OCT scans from five participants with aniridia and foveal hypoplasia grade from 0 to 4. Foveal hypoplasia was observed in 23 of the 25 participants with aniridia imaged with OCT (six had too severe nystagmus and/or ocular media opacities to allow imaging). Two participants had grade 1, four grade 2, eight grade 3 and nine grade 4. One participant had intermediate age-related macular degeneration (which made correct foveal grading difficult) and was, therefore, excluded from the analysis which included structural changes and visual function.

There was a strong correlation between the grade of foveal hypoplasia and visual acuity ($r_s = 0.859$, $P < 0.001$). A positive correlation was found between grade of foveal hypoplasia and CAD-LV RG threshold ($r_s = 0.558$, $P = 0.007$), but not for CAD-LV YB threshold ($r_s = 0.255$, $P = 0.252$). Those with foveal hypoplasia grade 0–2 (mild) were grouped and compared with those with grade 3 (moderate) and grade 4 (complete) for between-group comparisons (Fig. 5). Kruskal-Wallis ANOVA revealed a significant difference in CAD-LV RG thresholds between the grades of FH ($\chi^2 = 6.876$, $df = 2$, $P = 0.032$). Those with complete FH (grade 4) had significantly higher CAD-LV RG thresholds than those with mild FH (grade 0–2; $P = 0.038$) (Fig. 5A).

Secondary Pathology and Color Vision in Aniridia

Thirteen of the 27 (48%) participants with aniridia who performed the CAD-LV test had previously been diagnosed with glaucoma and/or ocular hypertension. Six of these had higher than normal thresholds for both CAD-LV RG and YB. Five of these six were also tested on the CCT, and four had higher than normal thresholds for both CCT RG and YB (ID: 5132, 5114, 5148, 5118). The only person without glaucoma who had higher than normal thresholds for both CAD-LV RG and YB (5149) had severe cortical and posterior subcapsular opacification of the lens. The participants with aniridia and the highest thresholds for both CAD-LV RG and YB (5129, 5118) had severe cortical and posterior subcapsular opacification of the lens. The participants with aniridia and the highest thresholds for both CAD-LV RG and YB had either ONH in addition to glaucoma (5114) or high-grade keratopathy in addition to glaucoma (5129, 5118). The other three with ONH had only increased RG threshold.

Kruskal-Wallis ANOVA indicated a difference in CAD-LV YB thresholds between the three AAK severity groups (mild: grade 0–1; moderate: grade 2; severe: grade 3; $\chi^2 = 6.749$, $df = 2$, $P =$...
0.034), but post hoc pairwise comparisons failed to reveal any significant differences (mild/moderate: \( P = 0.167 \), mild/severe: \( P = 0.079 \)) (Fig. 5D). Note that 12 of 13 with glaucoma also had AAK, and if the 13 with glaucoma were removed from the analysis, only one of the remaining with AAK (5149) had higher than normal CAD-LV YB threshold. The person with glaucoma and no AAK (5132) had higher than normal CAD-LV YB threshold. There were no significant differences in CAD-LV RG thresholds between any of the AAK severity groups (mild/moderate and mild/severe: \( P = 0.23 \)) (Fig. 5C).

Evaluation of lens opacities based on LOCS III grading was only applicable for persons with at least one phakic eye (\( n = 18 \)). Total grading score showed no significant correlation with either RG or YB CAD-LV thresholds (\( r_s = 0.296, P = 0.305 \) and \( r_s = 0.252, P = 0.384 \), respectively).

**DISCUSSION**

The results presented here show that persons with aniridia exhibit a quantifiable loss of color vision. The greatest loss was in RG color discrimination, which was positively correlated with the grade of foveal hypoplasia. Additional loss was observed in YB color discrimination, but this was associated with secondary ocular pathology, usually glaucoma. Color vision is known to vary as a function of age with maximum sensitivity around 20 years of age, thought to reflect normal healthy development and maturation of the visual system.\(^56\)-\(^58\)

The gradual increase in color discrimination from childhood until early adulthood can be observed for the normal controls in this study, but not for those with aniridia. These findings suggest that a likely reason for loss of red-green color vision in aniridia is arrested foveal formation and associated alterations in retinal structure and processing.

Loss of color vision has been described on one occasion previously, in a family with 11 members with congenital aniridia.\(^59\) We show that loss of RG and YB color vision is frequent in aniridia but cannot always be detected with the HRR pseudoisochromatic plates. Computerized tests, that allow for more accurate measures of chromatic discrimination such as the CAD-LV or the CCT, are required to quantify the loss. General loss in red-green discrimination was also observed for the Rayleigh match, with matching midpoints more than 2 standard deviations larger than for normal trichromats.\(^50\)

The majority of persons with aniridia (\( n = 23 \)) in this study exhibited the same variation in the degree of foveal hypoplasia (grades 1–4) as previously described for aniridia caused by \( PAX6 \) mutations.\(^13\),\(^36\),\(^60\) Only two participants had grade 0, both with clearly defined FAZ and RG thresholds within the normal range. Those with complete FH had significantly higher RG thresholds than those with mild FH. There was no association between age, RG thresholds, and FH grade. The degree of FH is most likely associated with the timing of arrested foveal formation,\(^22\),\(^28\) resulting in a lower number of L and M cones in the fovea, and foveal cones that are more similar to the peripheral cones with shorter cone outer segments\(^26\) and decreased cone photopigment optical densi-
Those who have anirida and associated FH because of PAX6 mutations may also have a lower ganglion cell density\(^1\) and lesser developed cone to midget circuitry (predominance of connections from a single cone to a single midget bipolar cell and a single ganglion cell),\(^17,19,26\) normally located within the FAZ.\(^20,22\) Thus, we argue that altered spatial organization of cone photoreceptors and associated circuitry of the central retina in aniridia is the most likely cause of poorer than normal red-green color discrimination. This is supported by the reduction in cortical volume in the region where the fovea is represented, previously reported in aniridia.\(^62\) We cannot rule out that unstable fixation and

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**Figure 4.** OCT scans from the central \(10^\circ\) of five participants with aniridia (A–E) showing the grades (0–4) of foveal hypoplasia.\(^36\) The dotted rectangle delineates the \(\times2\) magnified area represented on the right. The marked retinal layers were segmented and analyzed using longitudinal reflectivity profiles averaged over a 5-pixel wide region positioned at the foveal center (vertical arrows in left column). The distance between the ELM and the posterior boundary of OPL was defined as the outer nuclear layer thickness and the cone outer segments are bounded by the hyperreflective peaks corresponding to IZ and EZ. (B) Grade 1 is defined as absence of extrusion of plexiform layers (marked with arrow in magnified section on right). (C) Grade 2 is defined as grade 1 plus absence of foveal depression (marked with arrow). (D) Grade 3 is defined as grade 2 plus absence of outer segment lengthening (marked with arrows). (E) Grade 4 is defined as grade 3 plus absence of outer nuclear layer widening (marked with arrows). Scale bars: 200 \(\mu\)m.
Secondary pathology may aggravate both spatial and color vision function early on in aniridia, because with FH there are most likely far fewer cones and possibly not the same redundancy of cells within the macular region as that observed in normally developed retinas. This is corroborated by previously reported correlations between high contrast achromatic visual acuity and the degree of retinal development as ascertained by the grades of FH. It is known from studies on other retinal degenerative diseases that visual acuity may remain within normal limits even when cone density is less than half of what is observed in normal controls.

The RG thresholds were significantly higher than YB thresholds in persons with aniridia, with loss of YB color vision in addition to loss of RG vision appearing to be most strongly associated with glaucoma. Both RG and YB color vision loss have previously been reported in glaucoma, with higher thresholds being associated with severity of the disease. The two with aniridia who had the highest RG and YB thresholds, had had glaucoma for more than 20 years (assumed to reflect severity of glaucoma). Loss of YB color vision is also commonly associated with media opacities and a reduction in retinal illumination caused by increased absorption of short wavelength light. There was no significant correlation between YB color vision loss and grade of media opacity here, perhaps because only four participants who performed the CAD test had stage 3 AAK (central corneal involvement), and because of the younger age in the phakic group. The pattern of cataract development in aniridia is also different from typical age-related changes. Combined media opacities (AAK and cataract) in aniridia may result in a general reduction of retinal illumination. But, as measured with an added ND filter (to simulate reduced retinal illumination) in normal controls, this resulted in an almost uniform increase in both RG and YB CAD-LV thresholds, as opposed to the significantly higher RG thresholds in aniridia.

Both normal and impaired color discrimination has been reported in other disorders associated with foveal hypoplasia, but it is not known if other pathophysologic mechanisms that cause FH also affect color discrimination. In aniridia, we surmise that PAX6 mutation dosage expression is likely to be directly correlated with the degree of FH and the degree of impaired red-green color discrimination. Future work including more detailed PAX6 genotyping and additional measurement techniques, such as multimodal AOSLO imaging, may enable us to test this hypothesis.

CONCLUSIONS

In conclusion, visual function loss in aniridia is not limited to loss of visual acuity. Additional loss of color vision appears to be a combined consequence of the timing of arrested foveal formation and secondary ocular pathology. It is a reminder that persons with aniridia are equally likely to inherit congenital color vision deficiencies as others.

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