Multimodal Imaging in Best Vitelliform Macular Dystrophy

Jose Ronaldo Lima de Carvalho Jr,1–5 Maarjaliis Paavo,1 Lijuan Chen,1,4 John Chiang,5 Stephen H. Tsang,1,6 and Janet R. Sparrow1,6

1Department of Ophthalmology, Harkness Eye Institute, Columbia University, New York, New York, United States
2Department of Ophthalmology, Empresa Brasileira de Servicos Hospitalares (EBSERH) - Hospital das Clinicas de Pernambuco (HCPE), Federal University of Pernambuco (UFPE), Recife, Brazil
3Department of Ophthalmology, Federal University of São Paulo (UNIFESP), São Paulo, Brazil
4Department of Ophthalmology, People’s Hospital of PuTuo District, Shanghai, China
5Department of Ophthalmology, Oregon Health and Science University, Portland, Oregon, United States
6Department of Pathology and Cell Biology, Columbia University, New York, New York, United States

Correspondence: Janet R. Sparrow, Department of Ophthalmology, Columbia, University, New York, NY 10032, USA; jrs88@columbia.edu.
Submitted: January 10, 2019
Accepted: April 8, 2019

The integral membrane protein bestrophin-1 (BEST1) is encoded by the gene BEST1 (11q13) and is expressed on the basolateral membrane of RPE cells.1,2 Mutations in BEST1 are associated with early and adult-onset disease.3–5; the most common of these is juvenile-onset Best vitelliform macular dystrophy (BVMD) (OMIM 607854; BEST1).3–4 More than 200 disease-causing mutations have been described, most of which are missense mutations5 and located in the intracellular N-terminal portion of the protein. BEST1 also has been associated with defects in ocular development, specifically nanophthalmos.6

The BEST1 protein functions as an anion channel that is permeable to chloride and that is activated by changes in cytosolic calcium concentration.2–6,7–17 Under physiological conditions there may be additional activators.18 The crystal structures of bacterial and chicken homologs of BEST1 have confirmed that bestrophin-1 oligomerizes to form a pentamer with each protomer containing four transmembrane helices with cytosolic N- and C-termini.7–15 Mutations in BEST1 may be associated with disease due to the production of defective channels composed of mutant and wild-type subunits or because of disruption of the putative calcium-binding domain around the carboxylate loop.7,19

BVMD is typically inherited in an autosomal dominant manner, although autosomal recessive bestrophinopathy also is recognized and is modeled in a naturally occurring canine BEST1 knockout model.21 Although overt disease in BVMD manifests as bilateral disease that is often restricted to the macula,3 atypical BVMD can present as multifocal and extramacular involvement22,23 and unilateral disease can occur.24 The central oval lesion appears egg yolk–like (vitelliform lesion) in color fundus photographs, and exhibits intense near-infrared fundus autofluorescence (NIR-AF) and spectral-domain optical coherence tomography (SD-OCT) signal. At all stages of BVMD, nonlesion qAF was within the 95% confidence intervals for healthy eyes. Similarly, the NIR-AF intensity measurements outside the vitelliform lesion were comparable to the healthy control eye. SD-OCT scans revealed a fluid-filled detachment between the ellipsoid zone and the hyperreflectivity band attributable to RPE/Bruch’s membrane.

CONCLUSIONS. NIR-AF imaging can identify the pre-vitelliform stage of BVMD. Mutations in BEST1 are not associated with increased levels of SW-AF outside the vitelliform lesion. Elevated SW-AF within the fluid-filled lesion likely reflects the inability of RPE to phagocytose outer segments due to separation of RPE from photoreceptor cells, together with progressive photoreceptor cell impairment.

Keywords: best vitelliform macular dystrophy, bestrophin, bisretinoid lipofuscin, near-infrared fundus autofluorescence, optical coherence tomography, quantitative fundus autofluorescence, retina, retinal pigment epithelium
fluid-filled separation between photoreceptor cells and RPE.26–28

Electrophysiological testing of BVMD patients usually reveals a normal full-field electroretinogram, although there can be exceptions.29 On the other hand, an electrooculogram (EOG), a test that measures changes in the transepithelial potential across the RPE,30 can be diagnostic for BVMD31 if the recorded light peak/dark trough ratio (Arden ratio) is less than 1.5. The light peak is considered to reflect a depolarization of the basolateral membrane of RPE due to activation of a chloride conductance in response to changes in intracellular calcium concentration.30,32,33

This association led to the suggestion that BEST1 mediates the conductance involved in the light peak of the EOG.34 A recent study of microperimetry in BVMD reported that although sensitivity is particularly decreased in the affected central maculopathy, a reduction in retinal sensitivity also can occur in nonlesion areas.35

Emission spectra recorded spectrofluorometrically within the vitelliform lesions of BVMD patients exhibit maxima (580–620 nm) that are consistent with emission spectra of RPE lipofuscin recorded in healthy eyes.26 Because the vitelliform lesion is also notably hyperautofluorescent in SW-AF images, it has been assumed that this intensity represents a generalized increase in lipofuscin autofluorescence throughout the retina of BVMD patients.36 Nevertheless, we have previously observed that SW-AF (488 nm) autofluorescence, measured as quantitative fundus autofluorescence (qAF) is not elevated outside the vitelliform lesion in BVMD.26 Here we report qAF analysis of SW-AF images acquired from a second cohort of 14

<table>
<thead>
<tr>
<th>Patient</th>
<th>Family</th>
<th>Sex</th>
<th>Age</th>
<th>Ethnicity</th>
<th>Disease Stage OD</th>
<th>Disease Stage OS</th>
<th>BCVA* OD</th>
<th>BCVA* OS</th>
<th>Refraction OD</th>
<th>Refraction OS</th>
<th>Mutation</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>I</td>
<td>M</td>
<td>44</td>
<td>Caucasian</td>
<td>Vitelliform</td>
<td>Preclinic</td>
<td>0.10</td>
<td>0.00</td>
<td>+0.25</td>
<td>plan</td>
<td>c.884T&gt;C:p.Ile295Thr</td>
</tr>
<tr>
<td>2</td>
<td>II</td>
<td>F</td>
<td>34</td>
<td>Caucasian</td>
<td>Vitelliruptive</td>
<td>Vitelliruptive</td>
<td>0.00</td>
<td>0.00</td>
<td>plan</td>
<td>plan</td>
<td>c.727G&gt;A:p.Ala243Thr</td>
</tr>
<tr>
<td>3</td>
<td>III</td>
<td>F</td>
<td>12</td>
<td>African American</td>
<td>Vitelliruptive</td>
<td>Vitelliruptive</td>
<td>0.62</td>
<td>0.10</td>
<td>+6.0</td>
<td>+4.0</td>
<td>c.28G&gt;A:p.Ala10Thr</td>
</tr>
<tr>
<td>4</td>
<td>IV</td>
<td>M</td>
<td>59</td>
<td>Hispanic</td>
<td>Vitelliruptive</td>
<td>Vitelliruptive</td>
<td>0.88</td>
<td>0.50</td>
<td>+2.5</td>
<td>+2.75</td>
<td>c.653G&gt;A:p.Arg218His</td>
</tr>
<tr>
<td>5</td>
<td>V</td>
<td>M</td>
<td>49</td>
<td>Caucasian</td>
<td>Vitelliruptive</td>
<td>Atrophic</td>
<td>0.40</td>
<td>0.50</td>
<td>-1.5</td>
<td>-1.25</td>
<td>c.900G&gt;C:p.Glu300Asp</td>
</tr>
<tr>
<td>6</td>
<td>VI</td>
<td>M</td>
<td>55</td>
<td>Caucasian</td>
<td>Vitelliruptive</td>
<td>Vitelliruptive†</td>
<td>0.70</td>
<td>0.68</td>
<td>+2.0</td>
<td>+2.0</td>
<td>c.89A&gt;G:p.Lys30Arg</td>
</tr>
<tr>
<td>7</td>
<td>VII</td>
<td>M</td>
<td>23</td>
<td>Caucasian</td>
<td>Vitelliruptive</td>
<td>Vitelliruptive†</td>
<td>0.02</td>
<td>0.00</td>
<td>+1.25</td>
<td>+4.0</td>
<td>c.887A&gt;G:p.Asn296Ser</td>
</tr>
<tr>
<td>8</td>
<td>VII</td>
<td>M</td>
<td>61</td>
<td>Caucasian</td>
<td>Vitelliruptive</td>
<td>Vitelliruptive</td>
<td>0.68</td>
<td>0.62</td>
<td>+4.0</td>
<td>+4.0</td>
<td>c.887A&gt;G:p.Asn296Ser</td>
</tr>
<tr>
<td>9</td>
<td>VIII</td>
<td>F</td>
<td>32</td>
<td>Caucasian</td>
<td>Pseudohypopyon</td>
<td>Pseudohypopyon</td>
<td>0.10</td>
<td>1.2</td>
<td>+0.25</td>
<td>plan</td>
<td>c.727G&gt;A:p.Ala243Thr</td>
</tr>
<tr>
<td>10</td>
<td>IX</td>
<td>M</td>
<td>60</td>
<td>Caucasian</td>
<td>Vitelliruptive</td>
<td>Vitelliruptive</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>c.274C&gt;T:p.Arg92Cys</td>
</tr>
<tr>
<td>11</td>
<td>X</td>
<td>M</td>
<td>43</td>
<td>Caucasian</td>
<td>Vitelliruptive†</td>
<td>Vitelliruptive†</td>
<td>1.30</td>
<td>0.30</td>
<td>+0.75</td>
<td>+1.00</td>
<td>c.218T&gt;A:p.Ile73Asn</td>
</tr>
<tr>
<td>12</td>
<td>XI</td>
<td>F</td>
<td>43</td>
<td>Asian</td>
<td>Vitelliruptive</td>
<td>Vitelliruptive</td>
<td>0.20</td>
<td>0.00</td>
<td>plan</td>
<td>plan</td>
<td>c.652C&gt;T:p.Arg218Cys</td>
</tr>
<tr>
<td>13</td>
<td>XII</td>
<td>M</td>
<td>57</td>
<td>Caucasian</td>
<td>Atrophic</td>
<td>Vitelliruptive</td>
<td>1.30</td>
<td>0.60</td>
<td>-2.75</td>
<td>-2.50</td>
<td>c.652C&gt;T:p.Arg218Cys</td>
</tr>
<tr>
<td>14</td>
<td>XIII</td>
<td>M</td>
<td>45</td>
<td>Caucasian</td>
<td>Pseudohypopyon</td>
<td>Vitelliform</td>
<td>0.22</td>
<td>0.10</td>
<td>+0.50</td>
<td>+0.50</td>
<td>c.653G&gt;A:p.Arg218His</td>
</tr>
</tbody>
</table>

* BCVA: Best-corrected visual acuity logMAR equivalent.
† Eyes excluded from the qAF analysis.

**TABLE.** Clinical, Demographic, and Genetic Data

**FIGURE 1.** Quantitative fundus autofluorescence image analysis. Mean GLs recorded from eight circularly arranged segments (outlined in green) centered on the fovea were used to calculate qAF. (A) Vitelliruptive stage (P7). In this patient, three segments were excluded because they overlapped the lesion area. (B) Vitelliform stage (P14). All eight segments were analyzed in this patient.
By way of extending our previous work, we present not only qAF values reflecting SW-AF intensities outside the fovea, but also qAF color-coded images that illustrate the topographic distribution of SW-AF intensities. In addition, the signal derived primarily from RPE melanin was studied qualitatively and semiquantitatively in NIR-AF images acquired from the BVMD patients.

METHODS

Subjects

Fourteen patients heterozygous for BEST1 disease-causing variants were recruited for the present study. All patients presented to the Edward S. Harkness Eye Institute, Columbia University. Demographic, clinical, and genetic data are presented in the Table. The clinical diagnosis of BVMD was based on fundus appearance, family history, and low Arden ratio on EOG. The International Society for Clinical Electrophysiology of Vision (ISCEV) standards were accorded when performing the EOG. The anterior segment evaluation was unremarkable; all the patients were phakic and had clear media. BVMD patients were staged according to SD-OCT findings as published previously.

All procedures adhered to the tenets of the Declaration of Helsinki, and written informed consent was obtained from all patients after a full explanation of the procedures was provided. The protocol was approved by the Institutional Review Board of Columbia University.

Image Acquisition and Analysis: Short-Wavelength Autofluorescence (SW-AF)

qAF imaging was performed as described previously. SW-AF images (488-nm excitation, barrier filter transmitted light from 500 to 680 nm, 30° × 30° field) were acquired using a confocal scanning laser ophthalmoscope (cSLO; Spectralis HRA+OCT, Heidelberg Engineering, Heidelberg, Germany) equipped with an internal fluorescent reference for correction of variable laser power and differences in detector sensitivity. Pupils were dilated to at least 7 mm in diameter using 1% tropicamide and 2.5% phenylephrine before image acquisition. Room lights were turned off. A near-infrared reflectance image (NIR-R) was taken first and after switching to qAF mode, photoreceptor cells were bleached (20 to 30 seconds) during focusing and alignment to produce a uniform and maximum signal. The images were acquired in high-speed mode (8.9 frames/s), as a minimum of 12 frames (video format). The image quality was evaluated after recording and at least 7 of 12 frame images were required per video for further analysis. Misaligned frames (due to eye movements) or frames having diminished AF signal (due to eyelid interference or iris obstruction) were excluded.
FIGURE 3. Pre-vitelliform stage. P1. (A) SW-AF. The fundus appearance is normal. (B) NIR-AF imaging reveals foveal hypofluorescence. (C) qAF color-coded image of P1 presents intensities comparable to that in an age-similar healthy eye (E). (D) SD-OCT scan. The green line in A indicates the position of the SD-OCT scan; no abnormalities are noted.
An averaged non-normalized image was generated from each video and two images from each session were analyzed (four averaged images per eye).

For analysis, images were exported to IGOR (WaveMetrics, Lake Oswego, OR, USA). Mean gray levels (GLs) were determined in eight circular segments at an eccentricity of approximately 7º to 9º from the fovea (middle ring, Figs. 1A, 1B). Only segments that did not overlap the lesion were included in the measurement. The outer contour of the high SW-AF signal defined the outer limits of the lesion. The presence of vessels in the sampling area, which would decrease the qAF level, was accounted for by the software algorithm. qAF values were calculated after GLs were calibrated to GLs in the reference; and after accounting for the zero-GL of the laser, refractive error, image magnification, and age-adjusted lens transmission. For each eye, a qAF value was computed as the mean of the qAF values of the segments (qAFs).

Comparison was made to 374 eyes of 277 controls of similar age range (5 to 60 years old), without eye disease and having the following ethnic composition: 87 white, 79 Hispanic, 47 African American, 43 Asian, 6 Indian, and 15 mixed ethnicity (i.e., descending from more than one distinct race). Color-coded qAF maps were computed based on pixel-wise transformation of qAF values (WaveMetrics).

Near-Infrared Autofluorescence (NIR-AF)

The HRA2-SLO (Heidelberg Engineering) was used with the indocyanine-green angiography mode (787 nm excitation, 830 nm emission) to capture NIR-AF images (30 × 30º field). With a sensitivity of 96 (within a range of 51%–100% to adjust brightness) and the eye-tracking function, 100 single frames were averaged to obtain high-quality images saved in non-normalized mode. ImageJ (http://imagej.nih.gov/ij/; provided in the public domain by the National Institutes of Health, Bethesda, MD, USA) (Microsoft Java 1.1.4) was used to analyze and plot the NIR-AF signal. All images were aligned using i2kRetina software (DualAlign LLC, Clifton Park, NY, USA). Nineteen subjects (mean age 35.96 years) without a history of

Figure 4. Vitelliform stage. P1. (A) SW-AF Hyperautofluorescent signal within the lesion is considerably brighter than in surrounding fundus. (B) NIR-AF. Lesion has a dark central area and hyperfluorescent puncta. (C) qAF color-coded image. Hyperautofluorescence is confined to the lesion. qAF is not elevated outside the lesion. (D) Horizontal SD-OCT scans reveal a dome-shaped lesion. At the edge of the lesion the hyperreflective bands attributable to the ELM, EZ, and IZ are displaced anteriorly, separating these bands from the RPE/BM. Within the lesion, the IZ is disorganized. The positions of the scans are indicated by green lines in (A). The AF puncta in (A) and (B) (blue and yellow arrows) correspond to hyperreflective foci in the SD-OCT scans. (E) qAF color map of a healthy eye.

Figure 5. Pseudohypopyon stage. P9. (A) SW-AF discloses a lesion with hyperautofluorescence that is more pronounced inferiorly. (B) NIR-AF presents a dark central lesion with AF foci. The lesion is surrounded by hyperautofluorescence that is typical of the macula. (C) qAF color-coded image reveals an intensely AF punctum within the lesion. (D) The SD-OCT scan. The ELM and EZ bands follow the anterior contours of the lesion. The hyperreflective IZ is disorganized within the lesion. The optical clarity of the lesion is consistent with fluid in the line 1, while photoreceptors outer segment debris are likely the source of the dense deposits in the inferior region (line 2). The positions of the scans are indicated by green lines in (A). (E) qAF color-coded image acquired from an age-similar healthy eye demonstrates that qAF outside the lesion in (C) is within the normal range.
eye disease served as the healthy-eye group. These individuals self-identified as Caucasian (11), African American (3), Asian (3), and Hispanic (2).

Spectral-Domain Optic Coherence Tomography (SD-OCT)

SD-OCT images were acquired with the Spectralis HRA+OCT (Heidelberg Engineering) as horizontal 9 × 9-mm scans (870 nm; 7 microM axial resolution) through the macula acquired in high-resolution mode with averaging of 100 single scans. The scans were registered automatically to a simultaneously acquired IR-R (820 nm) fundus image, which was later used for point-to-point correlation with other fundus images. Nomenclature used to identify reflectivity bands in SD-OCT images was as published.38

RESULTS

We analyzed 28 eyes from 14 patients having a clinical diagnosis of BVMD that was confirmed by genetic testing. For all patients, heterozygous mutations were found in the BEST1 gene (Table). The age of the patients ranged from 12 to 60 years (mean 43.3 years; median 45 years). Patients were grouped as to stage of disease based on SD-OCT findings. 28 Of the 14 patients, 10 were male. Of the 26 eyes (13 patients) for which refraction was recorded, 16 were hyperopic, 4 were mildly myopic, and 6 were emmetropic. The visual acuity measured in logMAR varied from 0.00 to 1.30 (mean 0.43, median 0.35) in 26 eyes (for one patient visual acuity was not recorded). Most of the eyes included in this study were in the vitelliruptive stage (20), whereas one was classified in the pre-vitelliform stage, two were in the vitelliform, three in the pseudohypopyon stage, and two were atrophic.

The appearance of the fundus in SW-AF and NIR-AF images in BVMD patients varied with the stage of the disease. In an eye staged as pre-vitelliform, the central elliptical-shaped NIR-AF signal characteristic of a healthy eye (Fig. 2B) was altered by a central zone of reduced NIR-AF (Fig. 3B). Conversely, no abnormalities were noted in the SW-AF and SD-OCT images in the subclinical stage (Figs. 3A, 3D) when compared with a healthy eye (Figs. 2A, 2C).

In SW-AF images, vitelliform lesions were hyperautofluorescent and exhibited a small central decreased SW-AF (Fig. 4A). On the other hand, in NIR-AF images, the vitelliform lesions exhibited reduced signal (Fig. 4B). In SD-OCT scans, these lesions were located in central macula and presented as dome-shaped separations between the ellipsoid zone and RPE/Bruch’s membrane reflectivity layers (Fig. 4D). Hyperreflective material extended posteriorly into the fluid-filled space; based on continuity with the interdigitation zone, this material is assumed to be outer segments. Some hyperreflective foci associated with the lesion and visible in the SD-OCT (Fig. 4D)
corresponded to points of hyperautofluorescence in the NIR-AF image (Fig. 4B).

For the three eyes in the pseudohypopyon stage, the SW-AF signal in the lesion was more intense in the gravitationally dependent inferior zone of the lesion (Fig. 5A). Thus, in the corresponding SD-OCT images (Fig. 5D), the scan through the superior aspect of the lesion (Fig. 5A, line 1) revealed hyperreflectivity, whereas in the more inferior scan (Fig. 5A, line 2) the lesion was hyperreflective (Fig. 5D, lines 1 and 2). In the NIR-AF image, the lesions presented a generalized hypofluorescent signal with foci of normal or slightly increased fluorescence (Fig. 5B).

In the eyes in the vitelliruptive stage, zones of both hypo- and hyperautofluorescence were visible in SW-AF and NIR-AF images (Figs. 6A, 6B). Regions of increased fluorescent signal in SW-AF images (Fig. 6A, red asterisk) colocalized in SD-OCT scans with hyperreflective material that projected anteriorly; the RPE/Bruch’s membrane band was displaced and followed the contours of this material (Fig. 6C1, red arrow). Other anterior projections were visible in the SD-OCT scans and NIR-AF images (Figs. 6B, 6C2, 6C3; blue asterisks) but were hypautofluorescent in the SW-AF image. The apparent continuity of RPE over the projection is consistent with their detection in NIR-AF. The detection in NIR-AF images versus SW-AF images could also constitute an optical effect; absorption of NIR wavelengths by tissue (through which the light beam passes) is less than the blue SW light. Eleven of 20 vitelliruptive eyes presented with this hyperreflective anterior projection; this was not observed at other stages. This hyperreflective material has been previously described as choroidal neovascularization and fibrosis. Transmission of SD-OCT signal into the choroid was reduced posterior to the projection in SD-OCT scans (Fig. 6C1–3), whereas at other lesion-positions, transmission of SD-OCT signal was increased, indicating that the RPE was not intact.

For eyes in the atrophic stage, the lesions in SW-AF and NIR-AF images were largely absent of signal with only limited spots of brightness (Figs. 7A, 7B; red, yellow, and blue arrows), whereas in the SD-OCT there was a loss of outer retinal reflectivity layers (Fig. 7C1–3). Transmission into the choroid was increased in some areas (Fig. 7C3).

An additional disease feature that was observed was a halo of decreased NIR-AF and SW-AF signal just outside the border of the vitelliform lesion (Figs. 6A, 6B; 7A, 7B; 8A, 8B). The halo was more visible superiorly and laterally than inferiorly and was more prominent in the SW-AF images (Figs. 8B, 8C). Twenty-three of 26 eyes exhibited this halo; these eyes represented all stages of disease except the pre-vitelliform stage. Two eyes had a large lesion that extended beyond the vascular arcades, thus precluding analysis of the presence or absence of halo at the border of the lesion. The NIR-AF signal was also reduced in this region, except in two eyes. In the SD-OCT scans the halo corresponded to thinning of the outer nuclear layer (Fig. 8D).

qAF and semiquantitative NIR-AF measurements were acquired from 28 eyes of 14 BVMD patients. After image analysis, one patient (P11, two eyes) was excluded from the qAF analysis because the lesions extended beyond the arcades, precluding measurements outside the lesion. Another eye was excluded (P6, left eye) due to eye movement artifacts. In Figure...
qAF values measured outside the vitelliform lesion are plotted as a function of age for all remaining 25 eyes. Plotting the qAF values acquired from the single Hispanic, African American, and Asian patients versus ethnicity-matched healthy eyes revealed that in all three cases the qAF value of the BVMD patient was well within the 95% confidence intervals of the ethnicity-matched healthy eyes plotted as a function of age (data not shown). Therefore, for presentation (Fig. 9A) we compared all BVMD eyes with our database of 374 healthy eyes (277 control subjects) with the ethnic composition as described in the Methods section. Stage of disease is also indicated (Fig. 9B). In all cases, nonlesion qAF in the BVMD eyes was within the 95% confidence intervals for healthy eyes, independent of the stage of the disease (Figs. 9A, 9B).

Color-coded qAF images scaled to qAF units (1–1200) demonstrated foci of hyperautofluorescence inside the lesion. For instance, in the pseudohypopyon stage, the SW-AF signal of the vitelliform lesion was more intense inferiorly than superiorly and this difference was more noticeable in the qAF color-coded map (Fig. 5C). In the pre-vitelliform stage, SW-AF levels were comparable to a healthy eye (Figs. 3C, 3E). At other stages, the vitelliform lesions presented with variable but high SW-AF levels (Figs. 4C, 6D, 7D, 8C). Areas of atrophy and/or fibrosis showed low or absent SW-AF signal (Figs. 6D, 7D, 8C). In all stages of BVMD, qAF levels outside the lesion were normal (Figs. 3–7).

We also generated NIR-AF intensity profiles along a horizontal axis through the fovea and compared the profiles acquired from BVMD patients with profiles from healthy subjects (19 subjects). As shown in Figure 10, the NIR-AF signal in the healthy eyes increased to a peak in the fovea, whereas in the BVMD eyes the NIR-AF signal was depressed. The NIR-AF intensity was also reduced within the halo (Fig. 10, area between black and yellow lines) ($P = 0.03$).

**DISCUSSION**

We used the qAF approach to noninvasively measure SW-AF in BVMD patients and conclude that in fundus areas outside the central lesion, RPE lipofuscin levels are not increased. Thus, a generalized retina-wide increase in RPE lipofuscin is unlikely to be one of the primary features underlying the pathogenesis of BVMD. That said, the fluorescence within the vitelliform lesions exceeded the levels typically measured in the macula.26
Given that bisretinoid lipofuscin forms in photoreceptor outer segments before transfer to RPE, one need not assume that the autofluorescence within the vitelliform lesion is emitted from RPE. The most parsimonious explanation for the intense autofluorescence within the dome-shaped vitelliform lesion is that the emission originates from accumulating outer segment debris within the lesion, not because of an intrinsic inability of the RPE to phagocytose but because of separation of photoreceptor cells from the RPE monolayer.

The intense autofluorescence within the lesion probably also indicates that the rate of bisretinoid production in the photoreceptor cell outer segments is accelerated within the lesion. This could be due to the impairment of photoreceptor cells located in association with the fluid-filled lesion. As observed in a model of Merk deficiency, defective phagocytosis of outer segments by RPE leads not only to deposition of outer segment debris at the RPE-photoreceptor cell interface but also to elevated production of bisretinoid fluorophores within the accumulating outer segments and eventually photoreceptor cell degeneration.

Thus, the hyperautofluorescence of the vitelliform lesion is unlikely to be a primary disease feature but rather a secondary consequence of the fluid-filled lesion.

The focal reduction in the NIR-AF signal we observed in the pre-vitelliform stage (Fig. 3), has been noted previously. This variance in the NIR-AF signal was present without evidence of a lesion in SW-AF images or by SD-OCT. The anomaly may be indicative of an aberration in melanin. Alternatively, it could be due to an absorbance change. Water does not absorb light in the visible range (e.g., at the 488-nm wavelength used for SW-AF detection) but it does absorb in the near-infrared range (650–1350 nm). Thus, in the early stage of the disease, changes in the ionic and fluid composition of the cellular environment in the subretinal space may be the cause of the attenuated NIR-AF signal in the pre-vitelliform (subclinical) stage. We note, however, that fluid is not visible in the SD-OCT image at this stage (Fig. 3D). Mean retinal sensitivity measured by microperimetry was reported to be suppressed within 2 to 4 degrees eccentricity. This zone may correspond to the hypautofluorescent area detected by NIR-AF.

The hyper- and hypautofluorescence associated with the lesion at other stages of BVMD in NIR-AF images could reflect various processes. For instance, the depressed signal associated with the vitelliform lesion (Figs. 4–7, 10), particularly when viewed in conjunction with RPE/Bruch’s membrane thinning, likely reflects a loss of RPE cells. Alternatively, hyperautofluorescent foci within the lesion (Figs. 4–7) could be a sign of displacement and overlapping of RPE cells and/or elevated bisretinoid lipofuscin that contributes to the NIR-AF signal.

The presence of the fluid-filled lesion as observed in SD-OCT scans is likely the key to understanding disease processes in BEST1 disease. Because water follows the movement of anions, the outwardly directed movement of chloride is accompanied by fluid transport across the RPE cell from the subretinal space to the choroid. Accordingly, the formation of the fluid-filled detachment between photoreceptor cells and RPE in BVMD is attributable to impaired fluid transport secondary to the loss of anion channel activity. Reduction of anion currents in the presence of mutations in BEST1 has been demonstrated by whole-cell patch clamp recording.

**Acknowledgments**

Supported by grants from the National Eye Institute/NIH EY024091 (JRS); the Global Ophthalmology Awards Program, a Bayer-sponsored initiative committed to supporting ophthalmic research across the world (JRLC); Edward N. & Della L. Thome Memorial Foundation (SHT); Jonas Children’s Vision Care (SHT, JRS); and a grant from Research to Prevent Blindness to the Department of Ophthalmology, Columbia University.
References


12. Neussert R, Muller C, Milenkovic VM, Strauss O. The presence of bestrophin-1 modulates the Ca2+ recruitment from Ca2+-stores in the ER. Pflugers Arch. 2010;460:163–175.


