Progression of Late-Onset Stargardt Disease

Stanley Lambertus,1 Moritz Lindner,2 Nathalie M. Bax,1 Matthias M. Mauschitz,2 Jennifer Nadal,3 Matthias Schmid,3 Steffen Schmitz-Valckenberg,2 Anneke I. den Hollander,1,4 Bernhard H. F. Weber,5 Frank G. Holz,2 Gert Jan van der Wilt,6 Monika Fleckenstein,2 and Carel B. Hoyng1; for the Foveal sparing Atrophy Study Team (FAST)

1Department of Ophthalmology, Donders Institute for Brain, Cognition and Behaviour, Radboud University Medical Center, Nijmegen, The Netherlands
2Department of Ophthalmology, University of Bonn, Bonn, Germany
3Institute for Medical Biometry, Informatics and Epidemiology, University of Bonn, Bonn, Germany
4Department of Human Genetics, Donders Institute for Brain, Cognition and Behaviour, Radboud University Medical Center, Nijmegen, The Netherlands
5Institute of Medicine Genetics, University of Regensburg, Regensburg, Germany
6Department for Health Evidence, Donders Institute for Brain, Cognition and Behaviour, Radboud University Medical Center, Nijmegen, The Netherlands

Correspondence: Carel B. Hoyng, Department of Ophthalmology, Radboud University Medical Center, P.O. Box 9101 (OHK400), 6500 HB Nijmegen, The Netherlands; carel.hoyng@radboudumc.nl.

SL and ML are joint first authors.

MF and CBH contributed equally to the work presented here and should therefore be regarded as equivalent authors.

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PURPOSE. Identification of sensitive biomarkers is essential to determine potential effects of emerging therapeutic trials for Stargardt disease. This study aimed to describe the natural history of late-onset Stargardt, and demonstrates the accuracy of retinal pigment epithelium (RPE) atrophy progression as an outcome measure.

METHODS. We performed a retrospective cohort study collecting multicenter data from 47 patients (91 eyes) with late-onset Stargardt, defined by clinical phenotype, at least one ABCA4 mutation, and age at disease onset ≥ 45 years. We analyzed RPE atrophy progression on fundus autofluorescence and near-infrared reflectance imaging using semiautomated software and a linear mixed model. We performed sample size calculations to assess the power in a simulated 2-year interventional study and assessed visual endpoints using time-to-event analysis.

RESULTS. Over time, progression of RPE atrophy was observed (mean: 0.22 mm/year, 95% confidence interval [CI]: 0.19–0.27). By including only patients with bilateral RPE atrophy in a future trial, 32 patients are needed to reach a power of 83.9% (95% CI: 83.1–84.6), assuming a fixed therapeutic effect size of 30%. We found a median interval between disease onset and visual acuity decline to 20/32, 20/80, and 20/200 of 2.74 (95% CI: 0.54–4.41), 10.15 (95% CI: 6.13–13.34) years, respectively.

CONCLUSIONS. We show that RPE atrophy represents a robust biomarker to monitor disease progression in future therapeutic trials. In contrast, the variability in terms of the course of visual acuity was high.

Keywords: late-onset Stargardt, fundus autofluorescence, retinal pigment epithelium atrophy, biomarker, disease progression

Stargardt disease is an autosomal recessive retinal dystrophy caused by mutations in the ABCA4 gene, and affects 1:8000 to 1:10,000 people worldwide.1 Patients generally develop central loss of vision in childhood or early adulthood.2–4 However, late-onset Stargardt can be diagnosed at age ≥ 45 years, and has been associated with carrying one or two mutant ABCA4 alleles.5,6 Patients with this late-onset variant may first present with metamorphopsia or oscillopsia without any decrease in visual acuity. Occasionally, these patients are asymptomatic and are coincidentally diagnosed during screening tests for other retinal diseases, such as glaucoma, diabetes or thyroid disease.

The natural course of late-onset Stargardt includes a typical phenotype of yellow-white flecks and retinal pigment epithelium (RPE) atrophy. Patches of atrophy initially occur in the parafoveal retina and radially expand in size over time. New atrophic areas can arise; multifocal atrophic areas coalesce. Atrophic areas can form a ring encircling the intact and still-functioning fovea. Then, it is only until late in the disease course that the fovea will be involved in the atrophic process.7–9 The patient’s fixation point eventually shifts eccentrically, which leads to a substantial loss of visual acuity.10 However, central atrophy can also develop early in the disease course, and only minor disease progression has been described in other patients.5 Indeed, substantial variations in RPE atrophy progression have been reported before in small groups of typical Stargardt patients.11 Yet analyses of the natural course of large late-onset Stargardt cohorts are missing.

In light of recently upcoming therapeutic options for Stargardt disease,12–14 accurate biomarkers to determine their potential effects are crucial. The well-defined area of RPE atrophy is a frequent feature of late-onset Stargardt, showing...
similarities to geographic atrophy in age-related macular degeneration (AMD). Fundus autofluorescence (FAF) imaging can clearly visualize such areas of RPE atrophy,\(^{15,16}\) and change in RPE atrophy over time by FAF has already been accepted as a clinical endpoint by the U.S. Food and Drug Administration in AMD.\(^{17}\) We hypothesize that areas of RPE atrophy could also serve to sensitively monitor the effect of a drug trial in late-onset Stargardt. This would make patients with late-onset Stargardt appropriate candidates for upcoming therapeutic trials. In this study, we describe the natural history in late-onset Stargardt patients, and identify cohorts based on imaging parameters that determine the visual course in these patients. We quantify atrophy progression with semiautomated software, previously validated for AMD,\(^{18}\) showing the accuracy of this outcome measure, and include sample size calculations that are valuable for the design of upcoming therapeutic trials.

**METHODS**

**Patient Selection**

We identified patients from the Stargardt database of the Department of Ophthalmology at Radboud University Medical Center (Nijmegen, The Netherlands) and from the participants in the prospective natural history study Fundus Autofluorescence in Age-related Macular Degeneration (FAM; NCT003959692). We included 47 patients with a late disease onset, defined by an age $\geq 45$ years at which symptoms were first noticed.\(^5\) If the patient did not report any symptoms, we used the age at which the patient was diagnosed by an ophthalmologist. We clinically considered patients to have late-onset Stargardt when typical yellow-white flecks or dots were seen that correlated with hyperautofluorescent flecks on 488-nm FAF imaging.

Patients were analyzed for the presence of mutations in the ATP-binding cassette, subfamily A, member 4 (ABCA4, NM_000322.4) to exclude pseudo-Stargardt pattern dystrophy. A missense mutation, ABCA4-55, was always present. Additional genetic testing of PRPH2 (NM_000322.4) to exclude pseudo-Stargardt pattern dystrophy.\(^{20}\) We included patients without evidence of $ABCA4$ mutations. In patients carrying only one $ABCA4$ mutation, we performed additional sequencing of the peripherin-2 gene (PRPH2, NM_000322.4) to exclude pseudo-Stargardt pattern dystrophy and central areolar choroidal dystrophy.\(^{19,20}\)

This cohort study was carried out with approval from the Institutional Ethics Committee at Radboud University Medical Center (Nijmegen, The Netherlands) and the University Hospital of Bonn (Bonn, Germany), and adhered to the tenets of the Declaration of Helsinki. All patients provided informed consent before giving a blood sample and receiving additional ophthalmologic examinations to complete the clinical assessment.

**Clinical Assessment**

We reviewed the patients’ records for ophthalmologic history and available technical examinations, including sex, age at disease onset, and age at baseline. Best-corrected visual acuity (BCVA) was measured using a Snellen or Early Treatment Diabetic Retinopathy Study (ETDRS) chart, then transformed into the logarithm of the minimum angle of resolution (logMAR) for subsequent analysis. Fundus characteristics were documented using fundus photography (Topcon TRC-50IX; Topcon Corporation, Tokyo, Japan; or Visucam 500; Carl Zeiss Meditec, Jena, Germany). Fundus autofluorescence (488 nm; emission 500–700 nm) and near-infrared reflectance (NIR; $\lambda = 820$ nm) imaging were performed using a confocal scanning laser ophthalmoscope (Spectralis HRA+OCT or HRA2; Heidelberg Engineering, Heidelberg, Germany) in a subset of visits. The field of view was set at $30^\circ \times 30^\circ$ or $55^\circ \times 55^\circ$ and was centered on the macula. Eyes with signs of choroidal neovascularization were excluded from further analysis.

**Image Grading and Cohorts**

For each visit, two independent graders (MF and ML), blinded to each other’s results, evaluated the status of the fovea and the presence of clearly demarcated RPE atrophy (analogous to “definitely decreased autofluorescence,” the term recently used by Kuehlewein et al.\(^{21}\)) on all available imaging modalities. Atrophy was graded as follows: (1) no RPE atrophy with an intact fovea, (2) extrafoveal (but not fovea encircling) RPE atrophy, (3) a typical “foveal sparing” phenotype in which RPE atrophy encircled the fovea by $\geq 180^\circ$, or (4) foveal involvement. Foveal involvement was indicated by a mottled or absent autofluorescent signal (equaling what was ultimately termed “well/poorly demarcated questionable decreased autofluorescence”\(^{22}\)). In cases of discrepancy, a third grader (SL) evaluated the images. His agreement with one of the independent graders was finally used. Based on this grading, eyes were exploratively analyzed in order to form cohorts that might be predictive for visual acuity loss.

**Quantitative Measurements of Retinal Pigment Epithelium Atrophy**

We reviewed data using SAS Statistical Analysis Software Version 9.2 (SAS Institute, Cary, NC, USA) and R Version 3.1.2.\(^{23}\) Supplementary Figure S1 gives an overview of the analytical process applied in this work. Changes in visual acuity over time were assessed by time-to-event curves (cumulative distribution functions), and atrophy progression was analyzed using linear mixed-effects models. We performed a simulation study for power calculation for possible future interventional trials. Unless otherwise stated, all values given in the text represent median, minimum, and maximum values. Groups were compared by Mann–Whitney $U$ tests. Details on the statistical procedures can be found in Supplementary Text S1.

**RESULTS**

**Patient Features and Initial Symptoms**

A total of 91 eyes of 47 patients (19 men, 28 women) were included in this study. Two mutations in the $ABCA4$ gene were found in 20 patients (42.6%) and one mutation in 27 (57.4%; Supplementary Table S1). The median age at disease onset was...
54 years (range, 45–84). Self-reported initial symptoms were obtained for 42 patients and included a decrease in visual acuity (n = 24; 50%), metamorphopsia (n = 12; 29%), nyctalopia (n = 5; 12%), paracentral scotomas (n = 4; 10%), or oscillopsia (n = 1; 2%). Twelve patients (29%) did not report any visual complaints. In five patients, initial symptoms were not unequivocally denoted in the patient’s file.

Course of Visual Acuity

Overall, visual acuity data were available from 632 eye visits. At the first presentation after disease onset, the median disease duration was 0.9 years (range, 0–25.6) with a median BCVA of 0.10 logMAR (range, −0.14 to 1.70; Snellen 20/25). The median follow-up time of the patients with more than a single visit (45 out of 47 patients) was 4.8 years (range, 0.04–25.0). Time-to-event analysis yielded a median and 95% confidence interval (CI) between the age at onset and a decline in BCVA to mild visual impairment (n = 62), moderate visual impairment (n = 39), and severe visual impairment (n = 35) of 2.74 (0.54–4.41), 10.15 (6.13–11.38), and 11.38 (9.34–13.34) years, respectively (Fig. 1). The median disease duration at the final visit was 6.8 years (range, 0–30.9). The median BCVA at the final visit was 0.37 logMAR (range, −0.10 to 1.80; Snellen 20/47).

Assessment of Retinal Features

For each patient, clinical imaging data were available for a subset of visits (241 eye visits of 91 eyes). At baseline (first visit with imaging data available), yellow-white flecks were observed in all but one patient, in whom small yellowish spots were noted. An apparently intact fovea (Fig. 2A) without mottled or sharply decreased autofluorescence indicating RPE atrophy was present in 58 eyes. Out of these 58 eyes without foveal involvement, 22 had no RPE atrophy (Fig. 2B), 16 had extrafoveal (but not fovea encircling) atrophy (Fig. 2C), and 20...
TABLE 1. Retinal Features in Late-Onset Stargardt Over the Entire Observational Interval

<table>
<thead>
<tr>
<th>Initial Features</th>
<th>Median Visual Acuity, logMAR (Range)</th>
<th>Change in Features</th>
<th>Medial Visual Acuity, logMAR (Range)</th>
<th>Resulting Cohort</th>
</tr>
</thead>
<tbody>
<tr>
<td>No foveal involvement</td>
<td>22 0.10 (~0.20 to 0.64)</td>
<td>20 unchanged</td>
<td>0.06 (~0.06 to 0.58)</td>
<td>I</td>
</tr>
<tr>
<td>No RPE atrophy</td>
<td>22 0.10 (~0.20 to 0.64)</td>
<td>20 unchanged</td>
<td>0.06 (~0.06 to 0.58)</td>
<td>I</td>
</tr>
<tr>
<td>Extrafoveal but not fovea encircling</td>
<td>16 0.05 (~0.10 to 0.72)</td>
<td>2 eyes developed extrafoveal atrophy</td>
<td>0.01 (~0.00 to 0.02)</td>
<td>II</td>
</tr>
<tr>
<td>I</td>
<td>16 0.05 (~0.10 to 0.72)</td>
<td>2 eyes developed extrafoveal atrophy</td>
<td>0.01 (~0.00 to 0.02)</td>
<td>II</td>
</tr>
<tr>
<td>I</td>
<td>9 unchanged</td>
<td>9 unchanged</td>
<td>0.12 (0 to 1.54)</td>
<td>II</td>
</tr>
<tr>
<td>6 progressed to foveal sparing</td>
<td>6 progressed to foveal sparing</td>
<td>0.12 (0 to 1.54)</td>
<td>II</td>
<td></td>
</tr>
<tr>
<td>1 developed foveal involvement</td>
<td>1 developed foveal involvement</td>
<td>0.36</td>
<td>IV</td>
<td></td>
</tr>
<tr>
<td>16 unchanged</td>
<td>16 unchanged</td>
<td>0.22 (~0.04 to 1.80)</td>
<td>III</td>
<td></td>
</tr>
<tr>
<td>4 eyes loss of foveal sparing</td>
<td>4 eyes loss of foveal sparing</td>
<td>1.25 (0.94 to 1.50)</td>
<td>III</td>
<td></td>
</tr>
<tr>
<td>Foveal sparing encircling ≥ 180°</td>
<td>20 0.12 (0 to 0.52)</td>
<td>20 eyes unchanged</td>
<td>0.69 (~0.10 to 1.72)</td>
<td>IV</td>
</tr>
<tr>
<td>Mottled</td>
<td>25 0.30 (~0.08 to 1.80)</td>
<td>3 eyes changed to central RPE atrophy</td>
<td>1.30 (1.10 to 1.80)</td>
<td>IV</td>
</tr>
<tr>
<td>Central RPE atrophy</td>
<td>9 0.98 (0.30 to 1.50)</td>
<td>20 eyes unchanged</td>
<td>0.69 (~0.10 to 1.72)</td>
<td>IV</td>
</tr>
</tbody>
</table>

Cohort I had only flecks without any mottled foveal alterations or RPE atrophy; cohort II had extrafoveal (but not fovea encircling) RPE atrophy until the last visit; cohort III developed foveal sparing (RPE atrophy encircling the fovea ≥ 180°); and cohort IV had foveal involvement by either a mottled fovea or central RPE atrophy without passing through a foveal sparing phenotype during the observational interval.

Based on the morphologic observation of distinct cohorts among collective late-onset Stargardt patients, we further analyzed eyes with long follow-up grouped into the “foveal sparing” (cohort III) and “foveal involvement” (cohort IV) cohorts for possible distinctive long-term visual courses among these cohorts. Eyes that formed foveal sparing cohort III took an overall favorable visual course when compared to eyes from foveal involvement cohort IV. Survival analysis for the endpoints ≥ 0.2 logMAR, ≥ 0.6 logMAR, and ≥ 1.0 logMAR showed a notably later, though not significant, occurrence of each of these events in eyes from foveal sparing cohort III (Table 2).

TABLE 2. Median Times (Years; 95% Confidence Interval) of Best-Corrected Visual Acuity Decline Since the Age at Onset Compared Between Eyes of Late-Onset Stargardt Patients Who Developed Foveal Sparing and Those Who Had Early Foveal Involvement

<table>
<thead>
<tr>
<th>Cohort</th>
<th>Mild Impairment ≤ 20/32, ≥ 0.2 logMAR</th>
<th>Moderate Impairment ≤ 20/80, ≥ 0.6 logMAR</th>
<th>Severe Impairment ≤ 20/200, ≥ 1.0 logMAR</th>
</tr>
</thead>
<tbody>
<tr>
<td>Foveal sparing, cohort III</td>
<td>0.95 (NA*–6.61) years</td>
<td>10.15 (3.09–13.34) years</td>
<td>23.5 (13.6–NA*)</td>
</tr>
<tr>
<td>Foveal involvement, cohort IV</td>
<td>0.51 (NA*–4.41) years</td>
<td>7.73 (4.30–22.89) years</td>
<td>NA* (24.0–NA*)</td>
</tr>
<tr>
<td>Log-rank test</td>
<td>0.57</td>
<td>0.07</td>
<td>0.06</td>
</tr>
</tbody>
</table>

NA*, not available.

* Values could not be calculated, as too many events occurred outside the observational interval.
to deliver robust results,24,25 as these are required for planning was well defined by clinical, genetic, and validated imaging atrophy progression rates in a large multicenter cohort that comprehensively report on visual course and specific RPE atrophic RPE lesions, in which small changes are measurable, variance2 and overall slow decline; this would result in measure in clinical trials due to its high interindividual statistical power with a cohort of as few as 35 patients (Fig. 3). Although clinical features have been previously described,5 we promising clinical biomarker to monitor disease progression. The natural history of late-onset Stargardt features expanding, well-demarcated areas of RPE atrophy, which can be a AMD,17 to which areas of RPE atrophy show close similarities. In contrast, visual acuity loss may not be a useful outcome measure in clinical trials due to its high interindividual variance2 and overall slow decline; this would result in unrealistic large cohorts and long follow-up. Surrogate biomarkers may be more convenient, ideally predicting long-term changes in functional disease progression by detecting small short-term changes.23 Indeed, both visual acuity loss and atrophic RPE lesions, in which small changes are measurable, progress over time. Although this relationship may justify using RPE atrophy as a surrogate, there is a profound disconnect, particularly in late-onset Stargardt, between the area of RPE atrophy and vision. This discordance can be explained by clinically distinct progression subtypes: progression to either a foveal sparing phenotype in which RPE atrophy encircles the fovea in a horseshoe- or donut-like fashion (cohort III),7,8,10,22 or a subtype in which no such foveal sparing occurs (cohort IV). Foveal involvement can determine the eventual vision loss, either early when the eye has an initially involved fovea, or late when it exhibits a foveal sparing phenotype. As discussed above, visual acuity can vary widely, and for now, unpredictably, depending on the eventual foveal involvement. As the present data show, a large group of 37% with foveal involvement will do poorly, the rest relatively well. To determine those patients that would benefit most from therapy in terms of future clinical trials, analysis of additional imaging modalities could be helpful. While mottled decreased areas are difficult to quantify,21 other imaging biomarkers, in particular, spectral-domain optical coherence tomography, could indicate what drives the disease process toward foveal involvement. It has been demonstrated that outer retinal involvement precedes RPE loss. Specifically, outer nuclear layer and ellipsoid zone thinning can occur in regions of normal RPE thickness, suggesting that photoreceptor thinning may precede RPE degeneration.20 Hence, outer retinal damage on spectral-domain optical coherence tomography would preclude recognition on FAF In addition, environmental and genetic factors could significantly influence the development of RPE atrophy as identified in atrophic AMD.27,28 Such data were not included in this study and need to be addressed in future work.

Further limitations include the retrospective nature of the study and the resulting heterogeneity of the patients’ data, which may have been the reason for failing to show significance between different subtypes of late-onset Stargardt. For instance, some patients did not report any symptoms, and were more likely to have no RPE atrophy or only extrafoveal RPE atrophy not encircling the fovea. These patients would need a longer follow-up to identify in which direction the disease will develop. Analogously, heterogeneity within the imaging data, for example, the fields of view in NIR and FAF imaging, might have led to the nondetection of more peripheral atrophic lesions in patients with a 30° field of view, while such lesions would have been detected in eyes imaged with a 55° objective.

In recent years, identifying biomarkers in retinal disease has become a central issue for therapeutic trials that aim to test the efficacy of a drug. A surrogate outcome measure accepted by the U.S. Food and Drug Administration is geographic atrophy in AMD,17 to which areas of RPE atrophy show close similarities. As this study now has shown that RPE atrophy can also be used as an outcome measure in late-onset Stargardt, it may even be valuable in other retinal diseases affecting the RPE. Of special interest is the precise characterization of late-onset Stargardt patients; their adult age makes them ethically more appropriate candidates to participate in clinical trials than patients who are of minor age. This study provides important knowledge on the natural history of late-onset Stargardt, quantitatively describing the course of visual loss and atrophy progression. In addition, it provides fundamental information necessary to conduct clinical trials in patients with Stargardt disease.

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