




Monitoring and Management of the Patient with Stargardt Disease

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Abstract: Stargardt disease (STGD1) represents one of the major common causes of inherited irreversible visual loss. Due to its high phenotypic and genotypic heterogeneity, STGD1 is a complex disease to understand. Non-invasive imaging, biochemical, and genetic advances have led to substantial improvements in unveiling the disease processes and novel promising therapeutic landscapes have been proposed. This review recapitulates the modalities for monitoring patients with STGD1 and the therapeutic options currently under investigation for the different stages of the disease.

Keywords: Stargardt disease, inherited retinal dystrophy, multimodal imaging, optical coherence tomography, gene therapy

Introduction

Stargardt disease (STGD1) or Stargardt macular dystrophy is a recessive inherited retinal disease with an incidence of 8–10 per 100,000 persons.¹ First reported by Stargardt in 1909, it is caused by an autosomal mutation in the adenosine triphosphate binding cassette transporter 4 (*ABCA4*) gene.² Autosomal dominant pattern of inheritance of STGD1 has also been described, in association with mutations in *PRPH2*, *ELOVL4*, and *PROM1* genes.³

Yellow or white fish-shaped flecks and photoreceptors, retinal pigment epithelium (RPE) and choriocapillaris (CC) atrophy are the pathognomonic clinical features of STGD1.⁴ Flecks result from the accumulation of lipofuscin at the level of the RPE within or outside the main vascular arcades; flecks are thought to be the epiphenomenon of the RPE cell metabolic impairment in the clearance of photoreceptors debris.

STGD1 symptoms typically develop within the first two decades of life and include progressive, irreversible loss of central and color vision and delayed dark adaptation. The peripheral visual field is normal and there is no complaint of night blindness. Nevertheless, the natural course of the disease is characterized by marked clinical variability with regards to the age of onset, the pattern of fundus lesions, and the rate of progression.⁵ Unknown mechanisms as genotype-phenotype interaction or environment factors could modify the anatomical fate and the functional prognosis. To shed light on these differences, STGD1 patients have been extensively monitored by means of non-invasive imaging techniques.

Fundus autofluorescence (FAF) classically shows hyperautofluorescence corresponding to the flecks and hypoautofluorescence at the level of the RPE atrophy.⁶ Angiographic examinations, as fundus fluorescein angiography (FFA) and indocyanine

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green angiography (ICGA) show specific features but are poorly applicable in the first diagnosis of the disease. Optical coherence tomography (OCT) enlightens on the changes in the outer nuclear layer, as photoreceptor loss, RPE abnormalities, or the rare occurrence of choroidal neovascularization (CNV).⁷ Little is still known about the degree to which retinal and choroidal vascular networks are involved in STGD1, but new exciting information is coming from OCT angiography (OCTA) studies.^{8–10}

To date, no treatment is currently approved for STGD1 patients; however, stem cell therapy, gene replacement, and pharmacological strategies are the latest therapeutic promises intended to restore the RPE damage or slow down the advancement of the disease.¹¹ Recent trials are aiming to correlate clinical and functional factors with different rates of RPE atrophy enlargement, helping in stratifying clusters of patients and fixing clearer endpoints.^{5,12} The aim of this review is to recapitulate the modalities for monitoring patients with STGD1 and the therapeutic options presently under investigation for the different stages of the disease.

Molecular Basis

ABCA4 encodes for a retinal ATP-binding cassette protein located on the membrane of the outer segment discs of the cones and the rods.^{13,14} *ABCA4* works as a transporter that utilizes the energy of ATP hydrolysis to unidirectionally translocate retinoids (N-retinylidene-PE and all-trans-retinal) generated after photobleaching-induced isomerization of 11-cis-retinal, from the luminal to the cytoplasmic side of the disk membrane.¹³

Following isomerization and release from the cell, all-trans retinal travels to the RPE to be first esterified by lecithin-retinol acyltransferase (LRAT) and then converted to 11-cis-retinol by the isomerohydrolase RPE65. Finally, it is oxidized 11-cis-retinal before being transported back to the photoreceptor outer segment, where it is again conjugated to rhodopsin or cone opsin to form new, functional visual pigment.¹³ Failure of this transport results in the accumulation of lipofuscin, the main by-product of the photoreceptor visual cycle, into the RPE during the process of disk shedding. Lipofuscin and its components, especially N-retinylidene-N-retinylethanolamine (A2E), turn out to be toxic to epithelial and neuronal cells, with consequent RPE and photoreceptor degeneration.¹⁵ *ABCA4* knockout mice supported the hypothesis: accelerated deposition of lipofuscin in RPE and increased levels of N-ret-PE have been observed after light exposure in *abca4*^{−/−} mice.¹⁶

ABCA4 is a large complex gene in chromosome 1 consisting in 50 exons and has a causative role in numerous retinal diseases; mutations have been found in STGD1, cone-rod dystrophy, retinitis pigmentosa, and age-related macular degeneration (AMD).^{17,18} The extreme complexity of the gene makes it difficult to establish a thorough analysis of the genotype-phenotype interactions. Its high heterogeneity is reflected in the almost 6000 variant types reported in the literature, the majority located in the coding region of *ABCA4*,¹⁹ discriminating between normal and pathological variants is not always possible.¹⁹ To add complexity, heterogeneity was observed to vary among different populations.^{20–24} Therefore, genetic test represents a crucial inclusion criterion for access to clinical trials.

The most common genetic variants are missense and null mutations: the former are generally associated with a milder presentation, while the latter have severe visual prognosis and an early vision loss onset, with some exceptions. Some missense variants could also lead to fast-progressive disease, as p.Leu541Pro and p.Ala1038Val.²⁵ On the other hand, homozygous p.Gly1961Glu missense mutation phenotypically corresponds to adult-onset disease, with atrophy confined to limited areas of the macula, mild retinal dysfunction and variable visual acuity.²⁶

This large number of pathogenic variants in *ABCA4* only explain 60–75% of STGD1 phenotypes; other types of variants in non-coding regions, hardly identified through the most commonly-used strategies of genetic screening, may account for missing heritability in *ABCA4*-related diseases (that is, the gap between heritability estimates from genotype data and heritability estimates from twin studies).^{27,28} Some of these mutations include hypomorphic^{29,30} and deep intronic variants, which have a presumed effect on splicing.^{31–34} Non-coding deep-intronic splice variants represent potential interesting therapeutic targets for gene therapy approaches (i.e., CRISPR/Cas9 and antisense oligonucleotide-based splicing correction).³¹

So far, the use of a variety of mutation detection techniques for STGD have been reported, such as single-strand conformation polymorphism (SSCP)/heteroduplex analysis,³⁵ high-resolution melting,³⁶ microarray,³⁷ and direct Sanger sequencing.³⁸ These approaches are labor- and cost-intensive or low throughput. Next-generation sequencing (NGS) and whole exome sequencing (WES) strategies, on the contrary, allows for a comprehensive molecular diagnosis of STGD1 in a more rapid, cost-effective, and high throughput way.^{39,40} Nevertheless, a fair amount of patients with clinical diagnosis of STGD1 remains molecularly “unsolved”,^{29,41} and this

could be explained by the presence of yet unknown variants, unidentified regulatory systems, or inaccurate diagnosis (phenocopies).

Clinical Presentation

STGD1 clinical spectrum varies widely.^{4,42} Progressive bilateral loss can occur at any age; generally, two phenotypes can be distinguished on the basis of symptoms onset: a first peak around 20 years and a second peak around 40 years. Age of onset could be indicative of disease severity, with earlier onset being associated with a more aggressive disease course; however, this parameter should be interpreted cautiously, as it is mainly dependent on the degree of involvement of the fovea.

Adult-onset STGD1 is generally associated with missense mutations and better prognosis, while childhood-onset is largely linked to null variants with the poorest prognosis. Cases of very late-onset disease (i.e., after the sixth decade) are extremely rare.

Many attempts have been published to categorize the clinical spectrum of the disease; the most widely used classification belongs to Fishman, who described 4 stages based on clinical findings, and electrooculogram (EOG), electroretinogram (ERG), and psychophysical testing (Table 1).⁴³

Presenting best-corrected visual acuity (BCVA) is also heterogeneous. Moderate to high correlations between BCVA and both subjective onset ($r = 20.64$, $P = 0.001$) and duration of symptoms ($r = +0.71$, $P = 0.001$) have been

Table 1 Classifications of Stargardt Disease. The Groups Proposed Do Not Always Correspond to Different Progressive Stages of the Disease but Have Important Implications in the Clinical and Functional Assessment of the Patients

Fishman Classification ⁴³	Lois Classification ⁴⁴	FAF ⁴⁵	Early Onset STGD Fundus Grading System ⁴⁶	Genotype ⁴⁵
1. Flecks confined to the fovea, with pigmentary changes; "beaten-metal" or "snail-slime" appearance. EOG and both scotopic and photopic ERG normal (Figure 1)	1. PERG abnormality with normal full-field ERG	1. Localized low foveal FAF signal surrounded by a homogeneous background with or without perifoveal foci of high or low signal	1. Normal fundus	A. Patients with ≥ 2 severe/null variants
2. Flecks, extending anterior to the vascular arcades and/or nasally to the optic disc. Flecks might be partially or totally resorbed. Subnormal cone and rod responses on EOG and ERG, delayed dark adaptation (Figure 2)	2. PERG abnormality with cone ERG abnormality	2. Focal low FAF signal at the macula encircled by heterogeneous background and foci of high or low FAF signal anteriorly to the vascular arcades	2. Macular and/or peripheral flecks without central atrophy	B. Patients with 1 severe/null variant and ≥ 1 variant that are missense or in-frame insertion/deletion.
3. Flecks reabsorbed, atrophy of RPE and CC. EOG testing subnormal; ERG severely altered	3. Pattern ERG abnormality with generalized cone and rod dysfunction	3. Multiple areas of low FAF signal in macular area with heterogeneous background and/or foci of high or low signal	3a. Central atrophy without flecks 3b. Central atrophy with macular and/or peripheral flecks 3c. Paracentral atrophy with macular and/or peripheral flecks, without central atrophy	C. Patients with no severe/null variant, but ≥ 2 variants that are missense or in-frame insertion/deletion.
4. Further worsening of stage 3, complete reabsorption of flecks, extensive CC and RPE loss. ERG responses extinguished			4. Multiple extensive atrophic changes of the RPE, extending beyond the vascular arcades	D. Patients with 1 missense or in-frame insertion/deletion variant or with only variants predicted as less likely pathogenic or uncertain.

Abbreviations: CC, choriocapillaris; FAF, fundus autofluorescence; EOG, electrooculogram; ERG, electroretinogram; PERG, pattern electroretinogram; RPE, retinal pigment epithelium.

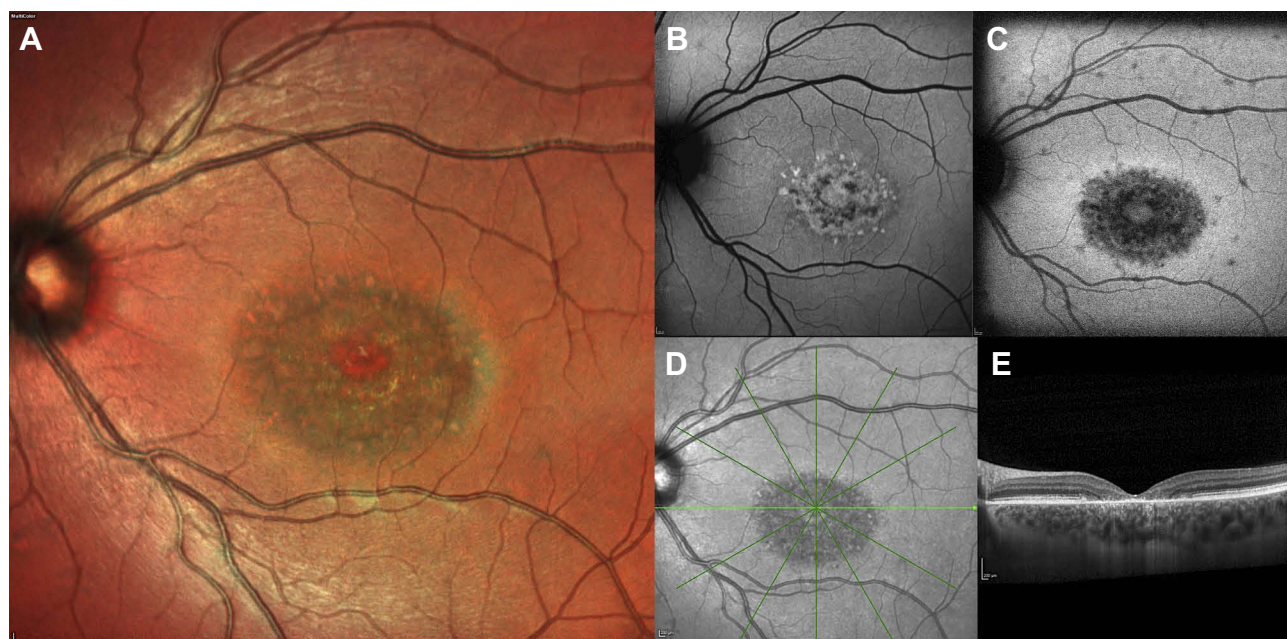


Figure 1 Multimodal imaging of a patient with STGD1 and macular involvement. Color fundus photograph (A), near-infrared-autofluorescence (B) and short wavelength fundus autofluorescence (C). Absence of RPE is visible as an area of irregular hypofluorescence on fundus autofluorescence. Several hyperautofluorescence lesions at the macula correspond to the flecks. The OCT scan (D) centered on the fovea shows atrophy of RPE and ellipsoid zone and backscattering.

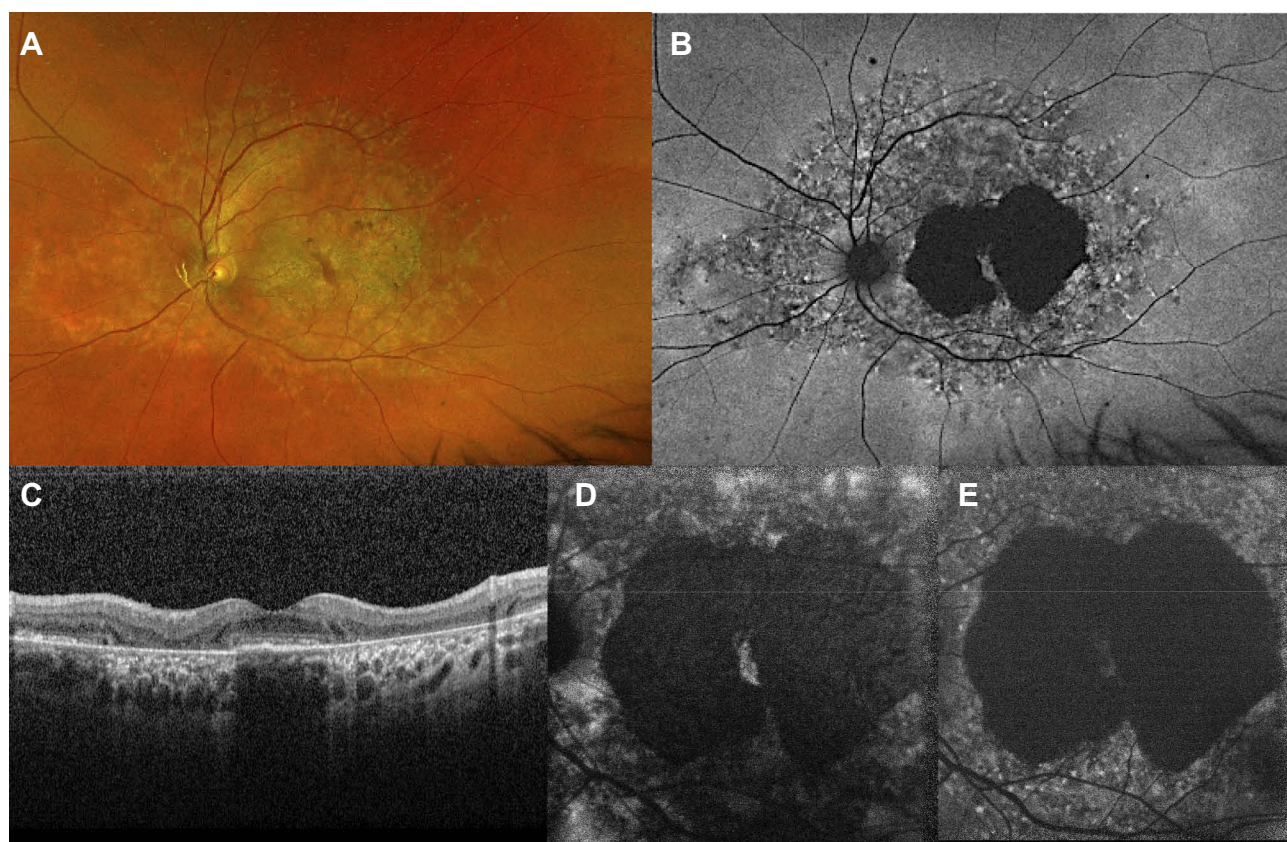


Figure 2 Ultra-widefield fundus photograph and autofluorescence of an advanced form of STGD1. The area of RPE mottling is comprised between the vascular arcades and broadens nasally to the optic disc (A, B). The macular area shows a marked hypofluorescence, with a little amount of foveal sparing (C). The OCT scan (C) demonstrates irregularity in the subfoveal RPE-photoreceptors complex and extensive atrophy in RPE and external retinal layers in the peri- and para-foveal areas. Foveal sparing is better seen on near-infrared autofluorescence (D) rather than short wavelength autofluorescence (E).

recently reported.⁴⁷ Patients with disease onset later than 20 years of age are more likely to maintain lifetime visual acuities better than 20/200 compared to patients with STGD1 presenting before 20 years of age. As predictable, the presence of foveal sparing ophthalmoscopically is associated with better visual acuity in diverse cohorts of patients.⁴⁸

The Progression of Atrophy Secondary to Stargardt Disease (ProgStar) study, a landmark project involving a large international cohort of STGD1 patients based on both prospective and retrospective analyses, reported no significant change in BCVA during a 12-month follow-up⁴⁹ and a clinically small BCVA loss over 24 months.⁵⁰ The change rate varied depending on baseline BCVA and visual involvement; eyes with no or modest acuity impairment but with a foveal lesion at baseline had the fastest loss rate. Nevertheless, due to its slow rate of change over time, BCVA might not be looked at as a reliable endpoint for putative STGD1 treatments.

Monitoring Progression of STGD1

Ophthalmoscopic signs and the visual function might not be helpful in monitoring STGD1, either in the first phase or in the later stage of the disease. The classic yellowish-white fundus flecks, the alteration of the foveal reflex, and the RPE and choroidal irregularities generally appear late. On the other hand, high BCVA might be maintained if a small island of foveal photoreceptors is preserved.

Conversely, novel non-invasive imaging techniques, as FAF and OCT, might attain valuable information on the metabolic state of the photoreceptor/RPE complex since the very early stages.

FFA reveals blockage of the choroidal fluorescence, referred to as the dark choroid sign, due to the lipofuscin accumulation into the RPE cells; however, the dark choroid is a late-onset feature and is not detectable in all cases. Moreover, in cases of advanced disease, it might be difficult to identify due to the presence of extensive window defects in regions of confluent RPE atrophy.⁵¹ ICGA provides useful information about the alterations of the choroidal vasculature, revealing hypofluorescence within the area of macular atrophy (dark atrophy).⁵² This sign has been interpreted as extended damage of the CC, and (when present) is useful to distinguish late-onset STGD1 from geographic atrophy (GA) secondary to age-related macular degeneration.

SW-FAF and NIR-FAF

Short-wavelength FAF (SW-FAF) imaging, based on the signal derived from lipofuscin excited with a 488-nm

wavelength beam, has been used as a fast and reproducible technique to calculate the progression of RPE atrophy in both monocentric and multicentric studies.¹² Reported rates of progression are slightly different across different investigations, as mainly dependent on the age of the patients, the advancement of the disease, and the method of hypo-FAF quantification.

In a large report on 68 patients, the rate of atrophy enlargement was calculated stratifying patients into 3 classes, according to the SW-FAF phenotype (Table 1).⁴⁵ The rate of progression was significantly faster in type 3 lesions (4.37 mm²/year) compared to type 1 (0.06 mm²/year) and type 2 (0.67 mm²/year) ones (Figure 3).⁴⁵

Indeed, two different types of decreased autofluorescence have been proposed based on the measurement of relative of darkness with respect to the optic nerve head (or, if not present in the image, the retinal vessels), considered as the reference for 100% blackness on the gray scale: definitely decreased autofluorescence (DDAF), defined as a signal more than 90% black in reference to the optic nerve head, and questionably decreased autofluorescence (QDAF), defined as 50% to 90% black, as compared to the reference.⁵³ Areas of QDAF tend to be unstable on consecutive examinations; they have been interpreted as a transition state between healthy retina and regions of advanced disease (ie, DDAF). Therefore, lesions of QDAF may be still amenable to rescue, and may be of interest for future therapeutic interventions. DDAF and QDAF can be summed up to calculate the total decreased autofluorescence (TDAF). Mean progression of TDAF has been estimated as 0.35 mm²/year (95% CI, 0.42–0.61) and 0.64 mm²/year (95% CI, 0.54–0.97) in the retrospective⁵⁴ and in the prospective¹² branch of the ProgStar study, respectively. The growth rates were dependent on initial lesion size, with larger lesions at baseline expanding faster than smaller ones. Unifocal lesions tended to display a milder course with respect to multifocal atrophy, regardless of lesion size at baseline.⁵⁴

Alongside the observation of hypo-FAF consequent to the RPE loss, FAF allows for the identification of other lesions pathognomonic of STGD1, as the hyperautofluorescent border surrounding the RPE atrophy in certain phenotypes,⁵⁵ which is believed to represent accelerated bisretinoid production in metabolically-deranged RPE/photoreceptor complex. This feature has been identified as a predictor of selective atrophy enlargement in STGD1, even though no association between the rate of progression and the hyper-FAF ring surrounding RPE atrophy has been proven.¹² Another distinctive feature of

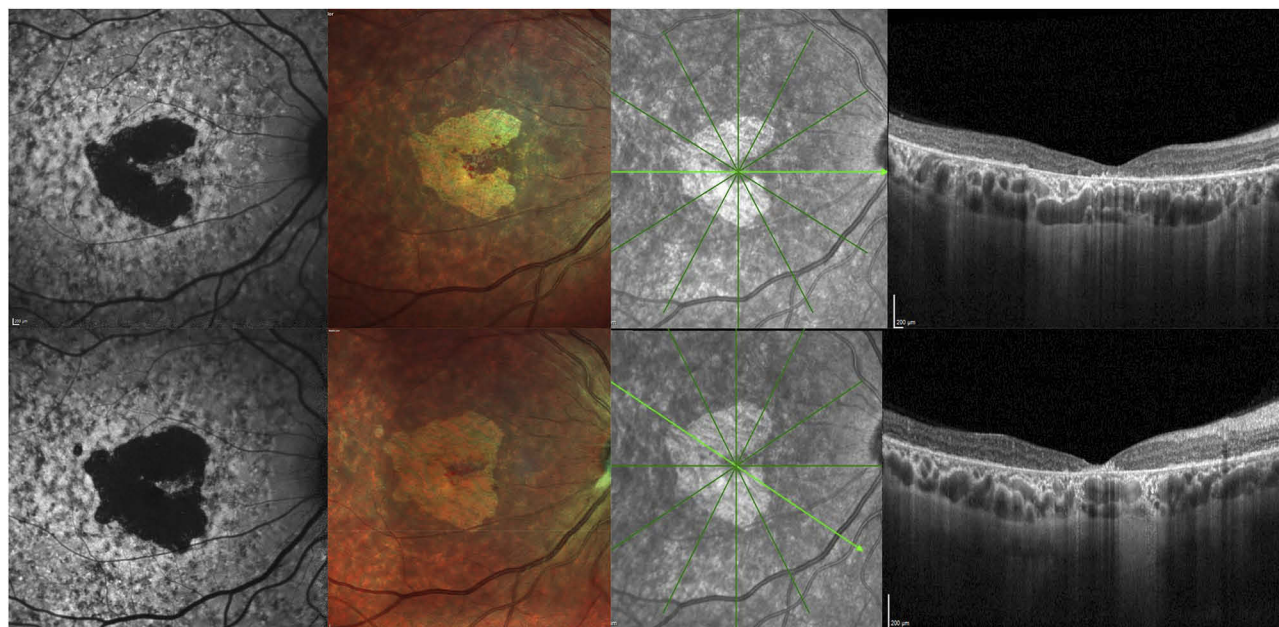


Figure 3 Progression of STGD1. First row: Short wavelength autofluorescence, Multicolor imaging, and OCT at baseline. Second row: Short wavelength autofluorescence, Multicolor imaging, and OCT after 12 months, showing enlargement of the area of RPE and photoreceptor loss on autofluorescence and central retinal thinning the foveal area.

STGD1 is the peripapillary sparing, which often helps to distinguish this disease from other phenocopies (e.g. *PRPH2*-related retinopathies).^{56,57}

Near-infrared FAF (NIR-FAF, 787 nm excitation), originating from melanin in the choroid and RPE, is another convenient method to visualize the alterations secondary to STGD1; NIR-FAF has been found to correlate consistently with SW-FAF and the loss of the photoreceptor band on OCT.^{58,59}

Theoretically, NIR-FAF has potential advantages over SW-FAF in the evaluation of STGD 1 progression. Because degenerating photoreceptor cells produce an abnormally higher amount of lipofuscin, SW-FAF signal might be falsely normal in correspondence of atrophic RPE lesions; on the other hand, NIR-FAF provides a better delineation of RPE cell loss and photoreceptor damage compared to shorter wavelength imaging techniques.⁶⁰ Also, fovea involvement may be better assessed with NIR-FAF (Figure 4). Nevertheless, NIR-FAF has not met the same enthusiasm in the international scientific community in comparison to SW-FAF. One of the drawbacks of NIR-FAF includes the technical difficulty of obtaining gradable images in the absence of adequate pupil dilatation and media transparency. Moreover, interpretation of NIR-FAF is challenging in cases of evident RPE atrophy due to the presence of melanin both within the RPE cells and the underlying choroid.

Relatively new automated or semi-automated tools to quantify FAF signal,⁶¹ as quantitative autofluorescence

(qAF),^{55,62} and new techniques, as fluorescence lifetime imaging ophthalmoscopy (FLIO),⁶³ have allowed more precise, objective and reliable methods of assessing the course of the disease. In particular, FLIO is a novel non-invasive imaging method detecting anatomic and metabolic changes of the human retina in vivo.⁶⁴ The analysis of retinal fluorescence lifetimes has been used to characterize and differentiate hyperfluorescent lesions and hypofluorescent atrophic areas in STGD1 and is potentially able to predict disease progression and assess therapeutic effects in upcoming clinical trials.⁶⁴

Optical Coherence Tomography and Optical Coherence Tomography Angiography

Spectral-domain OCT (SD-OCT) and swept-source OCT (SS-OCT) have led to consistent improvements in spatial and depth resolution in comparison to previous generation time-domain instruments. Both devices allow visualization of the individual layers of the retina and the choroid, en face reconstruction, or simultaneous imaging with infrared, color, or FAF acquisition.

A generalized thinning of the neuroepithelium and diffuse atrophic changes involving the outermost retinal layers (namely, the external limiting membrane (ELM), the ellipsoid zone (EZ) and the EPR) are the hallmarks of the disease. The EZ and the ELM are two distinct

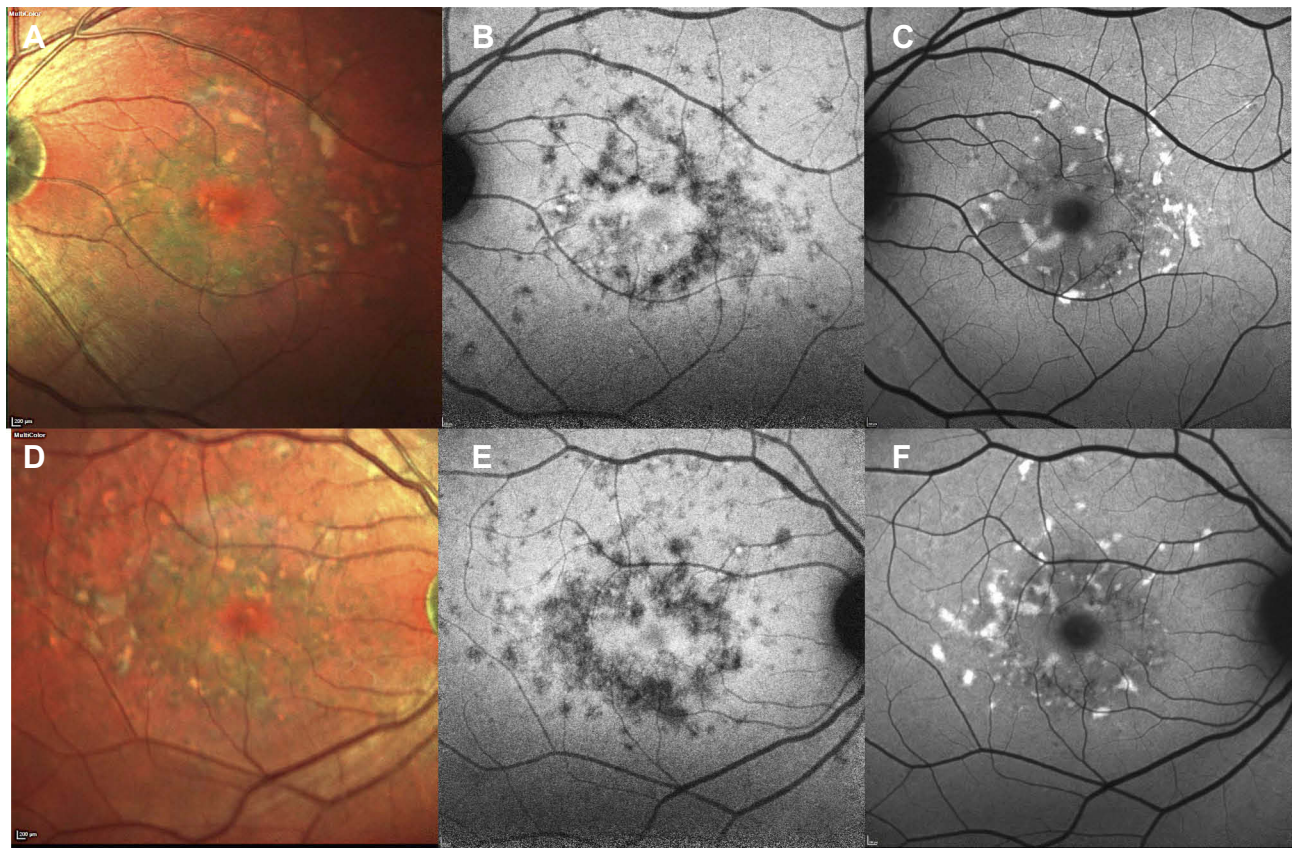


Figure 4 Comparison between short wavelength autofluorescence and near-infrared autofluorescence. Preservation of RPE in the foveal region is clearer on multicolor (**A, B**) and near-infrared autofluorescence (**C, D**) than on short wavelength autofluorescence (**E–F**), as the dark signal in the fovea in the latter due to blockage of normal choroidal fluorescence might be misinterpreted as RPE atrophy in the macular region.

hyperreflective layers in the outer retina strongly associated with photoreceptor integrity. The degree of EZ and ELM disruption correlates well with the BCVA⁶⁵ and the retinal function as measured on multifocal ERG (mfERG) or microperimetry (MP),^{60,66} consistent with the findings in other macular conditions. Using quantification of the EZ defect on serial SD-OCT imaging as the main predictor of disease progression, a rate of RPE atrophy expansion between 0.28⁶⁵ and 0.31 mm²/year⁶⁷ has been calculated. The vectors of atrophy enlargement were not uniform, and EZ changes were detected also in areas of apparently normal FAF signal. In fact, the extent of photoreceptor damage on SD-OCT appeared to exceed area of RPE loss according to different studies.^{59,68,69}

In a limited subgroup of patients, a peculiar macular phenotype has been recognized by means of SD-OCT, referred to as foveal cavitation (FC) (Figure 5).⁷ FC consists in the focal absence of the outer segments and EZ of the foveal photoreceptors in patients with general preservation of the ELM; this feature is generally stable versus

time, but is associated with a punctual profound loss of retinal sensitivity on MP and mfERG.

Finally, SD-OCT has shown reliability in the diagnosis and follow-up of patients developing macular CNV (Figure 6).⁷⁰

Enhanced depth imaging (EDI) OCT and SS-OCT allowed for deep range imaging of the CC and the choroid in STGD1 patients. Previous studies have found contradictory results with regards to subfoveal choroidal thickness (SFCT) in patients with STGD1 compared with matched normal controls.^{71–73} The analysis relying on SS-OCT revealed a generalized thinning of the choroid affecting mainly the small choroidal vessels.⁷⁴ Similarly, the choroidal vascularity index, calculated using the ratio of luminal area/total choroidal area, was significantly reduced in STGD1 patients as compared with normal eyes, with a significant negative correlation with BCVA.⁷⁵ Recently, four choroidal patterns have been described, with the following distribution: pattern 1 (normal appearing choroid) (15%), pattern 2 (reduced Sattler or Haller layer) (29%), pattern 3 (reduced Sattler and Haller layers) (26%), pattern 4 (Pattern 3 + choroidal caverns) (30%).

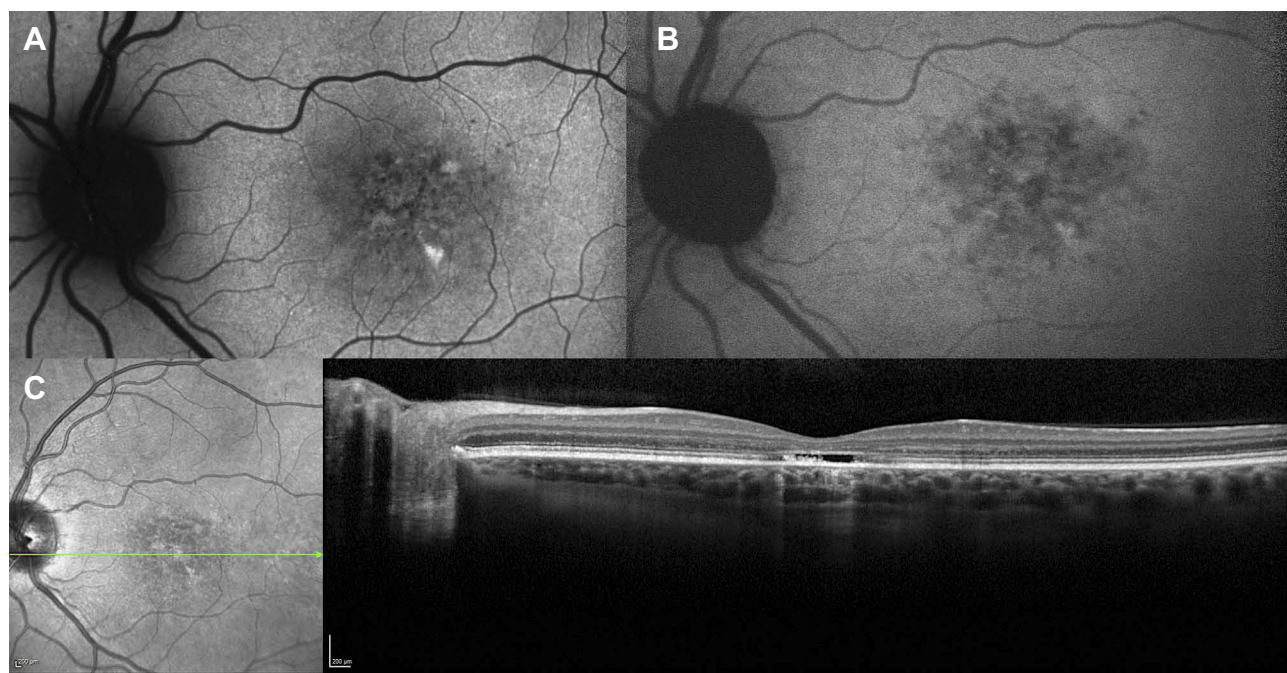


Figure 5 Foveal cavitation in STGDI. Short wavelength autofluorescence (**A**) and near-infrared autofluorescence (**B**) show macular mottling of RPE. In OCT (**C**), foveal cavitation appears as a hollow subfoveal space due to focal loss of RPE and photoreceptors with backscattering effect. External limiting membrane is preserved.

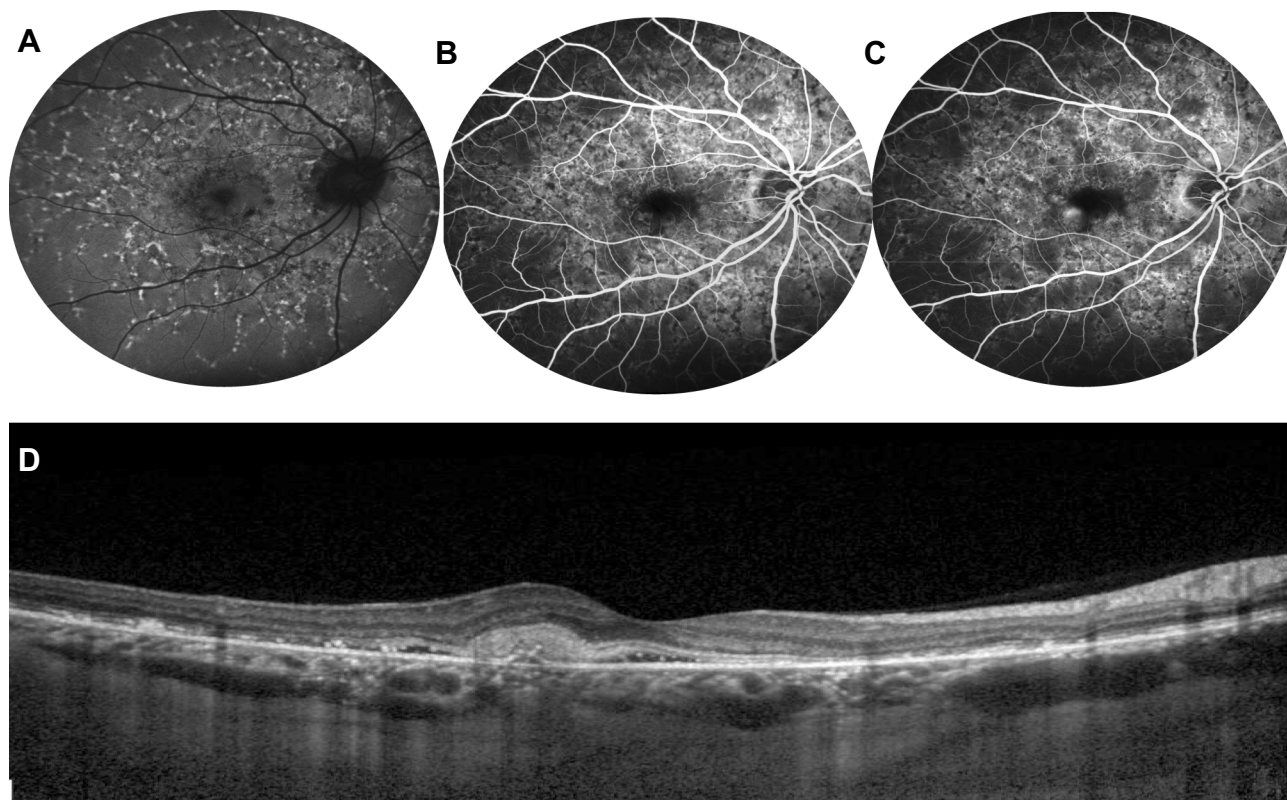


Figure 6 Choroidal neovascularization in STGDI. Fundus autofluorescence (**A**) shows hyperautofluorescent flecks and irregular macular hypoautofluorescence. Early (**B**) and late (**C**) fluorescein angiography frames demonstrate a parafoveal focal dye leakage, corresponding to a type I choroidal neovascularization (CNV). OCT scan (**D**) reveals growth of the CNV above the RPE with mild exudation, along with focal irregularity/atrophy of RPE/ellipsoid zone layers.

These patterns positively correlated with loss of retinal structural integrity on OCT. Furthermore, only advanced patterns (ie, pattern 3 and pattern 4) showed remarkable signs of progression after one-year follow-up.⁷⁶

OCTA allows to study retinal and choroidal microvasculature without dye injection and has been used to assess the superficial and the deep retinal vascular plexa as well as the CC in several retinal vascular diseases, including STGD1. There is overall agreement that patients with STGD1 show a reduction of the superficial capillary plexus (SCP), the deep capillary plexus (DCP), and the CC compared to healthy eyes (Figure 7).^{8,9,77} Comparing paired FAF and OCTA examinations, it appeared that RPE damage on FAF was significantly larger than CC loss on OCTA, which might suggest RPE damage precedes CC loss in the natural history of the disease.¹⁰ Moreover, OCTA permits an overall differentiation between STGD1 cases and GA secondary to AMD patients on the basis of CC impairment: while STGD1 eyes display a severe loss of CC with a clear demarcation between diseased CC and normal areas, GA revealed rarefied but still present CC within the area of RPE atrophy, with no detectable transition zone.⁷⁸ Longitudinal studies in GA secondary to AMD have demonstrated a significant correlation between CC impairment and disease progression,^{79,80} as longitudinal

studies involving OCTA in patients with STGD1 disease do not exist yet, the role of this technique in assessing the rate of progression of STGD1 is still vague.

Microperimetry

MP is a suitable tool to measure the residual function of the macula, the location, and the stability of fixation in STGD1 patients, thanks to the fundus tracking and fixation monitoring features embedded in most of the commercially-available instruments.^{81,82} These parameters have been used to quantify the decline of visual function in longitudinal studies of patients with STGD. As RPE atrophy progresses into the fovea, STGD1 patients begin to use an eccentric location of the retina to fixate, corresponding to a healthier retinal locus. This area is often characterized by reduced retinal sensitivity and poor fixation stability.

MP has also been used to examine the visual function changes related to the presence of retinal flecks; a significant difference in the mean retinal sensitivity has been found between flecked and “unflecked” areas, with most flecked areas having a decrease in sensitivity.⁸³

Although significant heterogeneity in retinal sensitivity and preferred retinal locus (PRL) of fixation have been recorded depending on the instrument used for the examination,⁸⁴ retinal sensitivity tends to be inversely correlated to disease duration and directly correlated with BCVA, even with some exceptions.⁸¹ Moreover, a linear relationship has been found between fixation location and BCVA (1° farther PRL eccentricity is associated with a 2.3-letter loss of BCVA) and between fixation stability and BCVA. In accordance with previous studies,⁶⁵ Schönbach et al demonstrated an association between earlier onset of disease symptoms and a longer duration of STGD1 with a more eccentric location of the PRL in a cross-sectional analysis of the ProgStar patients.⁸⁵

In conclusion, MP equipped with eye-tracking can provide a quantifiable evaluation of the remaining visual function, developing a better understanding of vision changes in STGD1 patients, and may be used as a secondary outcome measure for therapeutic trials in STGD1. These results may be important also in the rehabilitation of patients with central vision loss; biofeedback techniques coupled with MP have shown significant improvements in fixation stability, mean BCVA, mean reading speed, and contrast sensitivity for STGD1 subjects with unstable eccentric fixation.⁸⁶

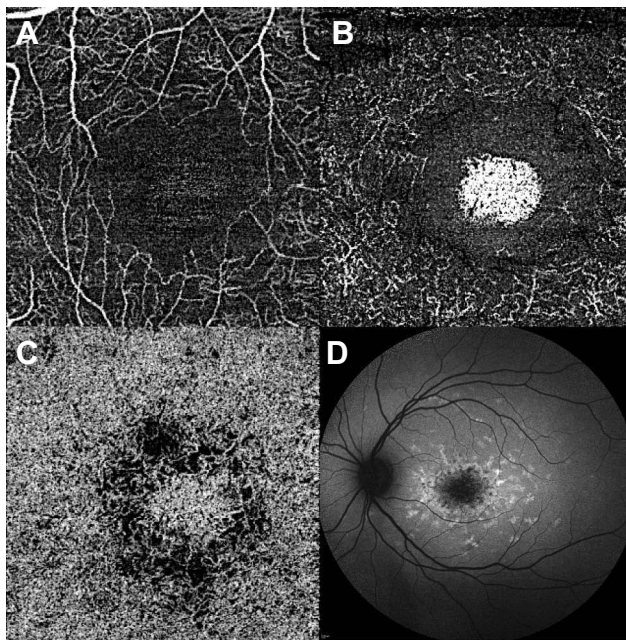


Figure 7 Optical coherence tomography angiography of STGD1. OCT angiography (left) shows foveal avascular zone enlargement and reduced vessel density on superficial capillary plexus (A) and deep capillary plexus (B), as well as damaged choriocapillaris (C). (D) Shows the corresponding autofluorescence.

Management and Treatment of STGD1

There is lack of consensus regarding the most appropriate management of patients with STGD1, and the actual therapeutic opportunities are scarce. Latest studies have focused on achieving a better understanding of the underlying genetic defects and the pathogenetic mechanisms leading to visual damage in STGD1, rather than proposing guidelines for follow-up and treatment.

Even if there is no approved treatment for STGD1, different therapies are under investigation (Table 2); the most promising are visual cycle modulators, complement inhibitors, gene therapy, and stem-cell therapy.

Visual Cycle Modulators

The aim of Visual Cycle Modulators (VCM) is to prevent the accumulation of toxic Vitamin A dimers at the level of the RPE.⁸⁷ Emixustat inhibits retinoid isomerohydrolase and decreases the conversion of all-trans-retinyl ester to 11-cis-retinol, with the aim to slow down the formation A2E. The drug has been recently tested in patients with GA secondary to AMD; the drug was overall well-tolerated,⁸⁸ but failed to exert a therapeutic effect on the rate of progression of GA in the treated arm.⁸⁹ A multicenter, randomized, double-masked,

placebo-controlled study to evaluate the efficacy and safety of emixustat compared to placebo in subjects who have macular atrophy secondary to STGD1 (SeaSTAR) is ongoing.⁹⁰

ALK-001 is a chemically modified form of vitamin A, in which three hydrogen atoms have been replaced with deuterium atoms (heavy hydrogen). In preclinical studies, ALK-001 reduced the buildup of lipofuscin and the decline in ERG amplitudes.⁹¹ It is currently being evaluated in a double-masked, placebo-controlled, clinical trial in Stargardt disease.⁹²

VM200 is a primary ammine which reacts with all-trans retinal and forms a non-toxic Schiff base, preventing the formation of A2E and cellular damage. VM200 has been tested in preclinical trials and showed positive results in murine models. Isotretinoin,⁹³ fenretinide, and other non-retinoids compounds were shown to inhibit the accumulation of lipofuscin in mouse models of recessive STGD1 by inhibiting different enzymatic steps of the visual cycle.⁹⁴

Complement Inhibition

Inflammation is thought to have a role in the progression of STGD1. A2E and other bisretinoids could activate the complement system in RPE cell, inducing inflammation

Table 2 Treatment Currently Under Investigation for Stargardt Disease

Treatment	Class	Mechanism	Company	Phase
ACU-4429 (Emixustat)	Visual Cycle Modulator	Inhibition of retinoid isomerohydrolase	Acucela Inc	Phase 3 (SeaSTAR)
ALK-001	Visual Cycle Modulator	Chemically modified Vitamin A preventing Vitamin A dimerization	Alkeus Pharmaceuticals	Phase 2 (TEASE)
VM200	Visual Cycle Modulator	Primary ammine which reacts with all-trans retinal forming a non-toxic Schiff base	Vision Medicine	Preclinical Trials
Isotretinoin	Visual Cycle Modulator	Inhibition of 11-cis-retinol dehydrogenase	Patent expired	Preclinical Trials
Fenretinide	Visual Cycle Modulator	Synthetic retinoid which binds to retinol-binding protein 4	Sirion Therapeutics	Preclinical Trials
AI120	Visual Cycle Modulator	Non retinoid RBP4 antagonist	iCura Vision	Preclinical Trials
Avacincaptad pegol (Zimura®)	Complement inhibition	Complement C5 inhibitor	Ophthotech Corporation	Phase 2b
SAR422459	Gene therapy	Recombinant lentiviral vector containing a functioning ABCA4	Sanofi, Oxford Biomedica	Phase I/II
Human embryonic stem cell-derived RPE	Stem cells therapy	Replacement of RPE cells restoring the function of overlying retina	Advanced Cell Technology	Phase I/II

and promoting RPE and photoreceptors degeneration.^{95,96} Complement inhibition has demonstrated a protective role in cultured RPE cells against apoptosis and a reduction in the accumulation of lipofuscin in a mouse model of STGD1.⁹⁷

Avacincaptad pegol (Zimura, Ophthotech Corporation) is an inhibitor of complement component 5 (C5), administered by intravitreal injection. C5 plays a pivotal role in the assemblage of membrane attack complex (MAC). Its inhibition is thought to prevent the formation of the MAC and reduce the rate of photoreceptor and RPE loss in STGD1. A phase 2b trial is ongoing.

Gene Therapy

In the majority of cases, STGD1 is caused by loss-of-function mutations of the gene *ABCA4*. Gene therapy aims to introduce a functional gene in the retina, supplementing the role of the impaired one in order to restore the normal cycling of 11-cis-retinal and preventing the accumulation of lipofuscin. There are two main ways to administer retinal gene therapy: intravitreal injection or subretinal delivery.⁹⁸ In the intravitreal injection, the vector is injected in the vitreous cavity and then diffuses to the retina. Subretinal injections are more invasive, requiring pars plana vitrectomy and a small neurosensory retinal detachment, but deliver the vectors directly between RPE cells and photoreceptors. In inherited retinal diseases, adeno-associated virus (AAV) and lentivirus have been used as viral vectors to deliver the intact target genes.⁹⁹ As the length of the *ABCA4* gene (6.8 kb) exceeds the transportation capacity of AAV (4.5–5 kb), lentiviruses have been considered the vector of choice for STGD1 trials. SAR422459 (Oxford Biomedica, Sanofi), formerly known as Stargen, is a recombinant lentiviral vector based on Equine Infectious Anemia Virus containing a functioning *ABCA4* gene. It was preclinically tested in mice, macaques, and rabbits, with encouraging results.^{100,101} A phase I/II clinical trial has started in 2011, but up to date, no preliminary results have been published.¹⁰²

Stem Cells Transplantation

Stem cells are a promising therapeutic option in STGD1, whose purpose is to replace lost RPE cell in degenerated tissue. Preclinical studies of RPE cells derived from human embryonic stem cell (hESC) showed encouraging results: the subretinal transplantation of these cells in mouse models improved visual function and photoreceptor survival in a dose-dependent way.¹⁰³ A phase I/IIa clinical trial was conducted to

test the safety profile of transplanted RPE cells originated from hESC in patients with STGD1, enrolling 13 patients. The trial met its primary endpoint, demonstrating the tolerability of stem cell therapy. A longer follow-up (5-years) of this study is still uncompleted.

Treatment of Complications

The development of CNV in patients with STGD1, although rare, can lead to rapid and severe vision loss. Laser photocoagulation¹⁰⁴ and photodynamic therapy¹⁰⁵ have been attempted for the treatment of choroidal neovascular membranes, without encouraging results.

The efficacy of vascular endothelial growth factor (VEGF) inhibition has been demonstrated for neovascularization of other rarer etiologies rather than AMD, including STGD1.^{70,106} Ranibizumab has shown slowing down CNV progression, apparently without significant visual improvement. However, due to the infrequent rate of CNV development in STGD1, it is difficult to achieve robust evidence for the long-term management of neovascular membranes in these patients.

Conclusion

STGD1 still represents one of the major common causes of inherited childhood and adulthood irreversible visual impairment. Due to its high phenotypic and genotypic heterogeneity, STGD1 is a complex disease to understand. Increasingly improved non-invasive imaging techniques, biochemical and genetic advances have led to substantial forward steps in the monitoring and management of patients with STGD1. Optimization in study design and implementation of interventional trials will be pivotal in reaching the ambitious goal to shed complete light on this disease.

Disclosure

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