Randomised study evaluating the pharmacodynamics of emixustat hydrochloride in subjects with macular atrophy secondary to Stargardt disease

Ryo Kubota 1, David G Birch,2 Jeff K Gregory,3 John M Koester3

ABSTRACT

Background/Aims Stargardt disease is a rare, inherited, degenerative disease of the retina that is the most common type of hereditary macular dystrophy. Currently, no approved treatments for the disease exist. The purpose of this study was to characterise the pharmacodynamics of emixustat, an orally available small molecule that targets the retinal pigment epithelium-specific 65 kDa protein (RPE65), in subjects with macular atrophy secondary to Stargardt disease.

Methods In this multicentre study conducted at six study sites in the USA, 23 subjects with macular atrophy secondary to Stargardt disease were randomised to one of three doses of daily emixustat (2.5 mg, 5 mg or 10 mg) and treated for 1 month. The primary outcome was the suppression of the rod b-wave recovery rate on electroretinography after photobleaching, which is an indirect measure of RPE65 inhibition.

Results Subjects who received 10 mg emixustat showed near-complete suppression of the rod b-wave amplitude recovery rate post-photobleaching (mean=91.86%, median=96.69%), whereas those who received 5 mg showed moderate suppression (mean=52.2%, median=68.0%). No effect was observed for subjects who received 2.5 mg emixustat (mean=−3.31%, median=−12.23%). The adverse event profile was consistent with prior studies in other patient populations and consisted primarily of ocular adverse events likely related to RPE65 inhibition.

Conclusion This study demonstrated dose-dependent suppression of rod b-wave amplitude recovery post-photobleaching, confirming emixustat’s biological activity in patients with Stargardt disease. These findings informed dose selection for a 24-month phase 3 trial (SeaSTAR Study) that is now comparing emixustat to placebo in the treatment of Stargardt disease-associated macular atrophy.

INTRODUCTION

Stargardt disease (STGD) is a rare, inherited, degenerative disease of the retina that affects 1 in 8000–10 000 individuals, making it the most common type of hereditary macular dystrophy. It typically causes vision loss during childhood or adolescence, progressing slowly over time until an individual’s vision is 20/200 or worse. Currently, a pressing need exists for novel therapies for STGD, as no approved treatments for the disease exist.

The vast majority of STGD cases are type 1 (STGD1) and caused by mutations of the ABCA4 gene, in contrast to ELOVL4 (STGD3) and PROM1-associated STGD4. ABCA4 encodes a protein that transports a precursor of toxic bis-retinoids (N-retinylidene-phosphatidylethanolamine (PE), which consists of one molecule of PE covalently bound to one molecule of cis- or all-trans-retinal) from the lumen side of photoreceptor disc membranes to the cytoplasmic side, where the retinal is hydrolysed from PE. When ABCA4 is mutated, this precursor molecule accumulates in disc membranes, is phagocytosed by retinal pigment epithelium cells, and is eventually converted into toxic bis-retinoids such as N-retinylidene-N-retinylethanolamine (A2E). Build-up of toxic bis-retinoids can lead to loss of function or death of the retinal pigment epithelium. This, in turn, leads to the dysfunction of both rod and cone photoreceptors, their eventual death and permanent loss in vision.

Evidence suggests that emixustat hydrochloride may protect against the retinal damage that occurs in STGD. Emixustat is an orally available small molecule that targets the retinal pigment epithelium-specific 65 kDa protein (RPE65), reducing the production of visual chromophore (11-cis-retinal) in a dose-dependent and reversible manner. By depleting 11-cis-retinal, and thus all-trans-retinal, the substrates necessary for the production of toxic bis-retinoids such as A2E, emixustat decreases the rate at which these harmful molecules accumulate in the retinal pigment epithelium. In an ABCA4/- mouse model of STGD, in which excessive A2E accumulates in the retinal pigment epithelium, treating young mice with emixustat for 3 months significantly reduces A2E accumulation when compared with vehicle treatment.

The primary objective of this study was to characterise the pharmacodynamic effects of emixustat on electroretinography (ERG) in subjects with macular atrophy (MA) secondary to STGD, when the drug was administered orally for 1 month. The rod b-wave amplitude of the ERG is regarded as a reliable measure of signal processing in the retina and represents the extent of rhodopsin regeneration. There is a proportional relationship between the magnitude of the rod b-wave amplitude and rhodopsin levels. Reduction of the rod b-wave amplitude indicates a reduction in rhodopsin, and thus 11-cis-retinal, levels. Following exposure to a bright bleaching light, the rod b-wave amplitude is reduced, but recovers over approximately 30 min. The rate of recovery over time reflects the regeneration rate of rhodopsin, and hence 11-cis-retinal,
which is governed by RPE65 activity. Thus, assessing the effects of emixustat on rod recovery after photobleach is an indirect measure of RPE65 inhibition.

**MATERIALS AND METHODS**

**Study design**
This multicentre, randomised, double-masked study was conducted from January to November 2017 at six study sites in the USA, at which subjects with MA secondary to STGD were assigned to one of three doses of emixustat (2.5 mg, 5 mg or 10 mg) and treated daily for 1 month. The three emixustat doses were selected based on experience in prior clinical trials. The study was conducted in accordance with the Declaration of Helsinki, the Health Insurance Portability and Accountability Act (HIPAA), and the basic principles of the International Council for Harmonisation (ICH) E6 (R1) Guideline for Good Clinical Practice. All subjects provided written informed consent before study-specific procedures began. This study was registered with ClinicalTrials.gov as NCT03033108.

**Participants**
The study planned to enrol approximately 30 subjects, with the goal of having 24 (8 per treatment group) complete the masked treatment phase.

**Inclusion criteria included**
- Ability to provide informed consent.
- ≥ 18 years of age.
- Clinical diagnosis of MA secondary to STGD in one or both eyes, with one or more pathogenic mutations of the ABCA4 gene. If only one pathogenic mutation was detected, a typical fleck phenotype was required.
- Total area of MA (definitely decreased autofluorescence (DDAF) plus questionably decreased autofluorescence (QDAF) on reduced-illuminance fundus autofluorescence (RI-FAF) in the study eye equal to 1.25–26 mm², as measured by a central image reading centre (Duke Reading Center, Durham, North Carolina, USA).
- Early treatment diabetic retinopathy study best-corrected visual acuity (BCVA) of ≥20 letters in the study eye.
- Ability to administer oral medication by self or with available assistance.

The original study protocol, which limited the total area of MA to 1.25–18 mm² of DDAF, was amended to the criterion above, as this allowed enrolment of a broader population of subjects with STGD by including larger lesions and those containing QDAF, an earlier stage of MA than DDAF. Reliable measurement of QDAF can be challenging as the borders may be poorly demarcated. All RI-FAF images were obtained with the Heidelberg SPECTRALIS; and all areas of MA were measured using Heidelberg RegionFinder software.

**Exclusion criteria included**
- MA associated with a condition other than STGD in either eye.
- Active ocular disease that compromised or confounded visual function (eg, choroidal neovascularisation, diabetic retinopathy, uveitis).
- Any intraocular or ocular surface surgery within 3 months of screening.

If both of a subject’s eyes qualified for study, the investigator selected the eye with the larger MA lesion; if both eyes had an equal MA lesion area, the right eye was selected.

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**Figure 1** Disposition of study subjects.
Subjects were treated with the assigned dose of emixustat every evening for 1 month. Scheduled visits included screening (within 30 days prior to baseline), baseline (day 1), end of treatment (EoT; day 30) and study exit (30 days after EoT).

Full-field ERG was performed for each eye in all subjects at baseline and EoT. Identical ERG systems were used at each study site (Diagnosys, Lowell, Massachusetts, USA). ERG data were sent to a central ERG reading centre (Retina Foundation of the Southwest, Dallas, Texas, USA) to be evaluated. Briefly, the ERG procedure began with the subject’s pupils being maximally dilated, followed by a period of dark adaptation that lasted for at least 30 min. At the end of this period, an electrode (Dawson Trick lens) was placed on each eye under dim red light, and a rod response and a maximal response (mixed) ERG were recorded. Following a single-flash cone response and 31 Hz flicker response at the end of a 10 min period of light adaptation, subjects underwent photobleaching for a period of 3 min. Rod responses were recorded immediately after photobleaching and at 10, 20 and 30 min following the photobleaching.

Assessments performed only at screening included ABC4A genotype typing (Molecular Vision Lab, Hillsboro, Oregon, USA) and RI-FAF (performed on a Heidelberg Spectralis). At screening, baseline, EoT, and study exit, BCVA, slit-lamp biomicroscopy, and dilated ophthalmoscopy were performed. At screening and EoT, physical examination, vital signs, electrocardiography, clinical laboratory tests and intraocular pressure were performed.

### Outcomes

The study’s primary outcome was the percent suppression of rod b-wave recovery rate after photobleaching at the end of the 1-month treatment period, compared with baseline. This was defined as the percent decrease from baseline in the rod b-wave amplitude recovery rate (slope in μV/minute) over 30 min postphotobleaching, as measured by ERG. The percent suppression prebleach was defined as the percent reduction in the pre-bleach b-wave amplitude (μV) at month 1 relative to baseline. The secondary outcomes were (1) incidence, severity, seriousness of and discontinuations due to adverse events (AEs); and (2) changes from baseline in laboratory values, vital signs, physical examination findings, ECGs and ophthalmic assessments. All AEs had their verbatim terms mapped to the corresponding thesaurus terms using the Medical Dictionary for Regulatory Activities coding dictionary, V20.1.

### Statistical methods

The study’s pharmacodynamic analysis included all randomised subjects who had (1) a baseline ERG and (2) changes from baseline in laboratory values, vital signs, physical examination findings, ECGs and ophthalmic assessments. The 21 subjects who took at least 1 dose of emixustat. All safety endpoints were summarised descriptively.

### RESULTS

The investigators screened 44 subjects (figure 1). Of these, 21 did not pass the screening process. The 23 remaining subjects...
were randomised to emixustat treatment: 7–2.5 mg, 9–5 mg and 7–10 mg. All but one subject (from the 5 mg group) completed the study. All study exit visits were completed.

Demographic characteristics were generally comparable between the three treatment groups. More than half of the subjects (15/23, 65.2%) were male, and most were white (18/23, 78.3%). Their mean age was 51.6 years, with an age range of 18–77 years, and more than half of subjects (13/23, 56.5%) were ≥45 years. Overall, study drug compliance was 95.8% and was similar between the three treatment groups; the majority of subjects (20/23, 87.0%) were >80% compliant, and only one subject (in the 5 mg group) had 25%–50% compliance.

Baseline data were also comparable between the three treatment groups, with the exception of mean total MA area in the study eye, which was largest among subjects treated with 10 mg and smallest in subjects treated with 5 mg (table 1). Of 23 subjects, 5 had one pathogenic ABCA4 mutation and the remaining 18 had two pathogenic ABCA4 mutations. The mean BCVA letter score for subjects’ right eyes was 51.3 and for left eyes was 50.7. Of the 19 subjects with evaluable baseline ERGs, 15 were assessed as Lois group 1.20

At baseline, the observed mean values for b-wave amplitudes decreased after photobleaching and recovered to near prebleach values by 30 min after photobleaching in all subjects (figure 2A). At EoT, the mean change from baseline (CFB) rod b-wave amplitude before photobleaching was similar in subjects following treatment with 2.5 mg and 10 mg, while the largest decrease in subjects treated with 10 mg was largest in subjects treated with 10 mg (−10.42 μV/minute), followed by subjects treated with 2.5 mg (−4.12 μV/minute), and 5 mg (−0.42 μV/minute). Thus, subjects treated with 10 mg showed near-complete suppression of the rod b-wave recovery rate post-photobleaching (mean=91.86%, median=96.69%), whereas subjects treated with 5 mg showed a moderate degree of suppression (mean=52.2%, median=68.0%) (table 2). No effect was observed for subjects treated with 2.5 mg (mean=−3.31%, median=−12.23%). The results were consistent between the subjects with one identified ABCA4 mutation (n=5) compared with those with two identified ABCA4 mutations (n=14).

The mean CFB for ERG rod and cone mixed response for b-wave amplitude was larger in subjects following treatment with 10 mg than in subjects treated with 5 mg or 2.5 mg. The mean changes for 5 mg and 10 mg were decreases. The mean CFB b-wave implicit times were small, and no differences could be assessed between the three treatment groups. At month 1, the mean CFB in the single-flash cone response and 31 Hz flicker cone response for b-wave or peak amplitudes, respectively, and implicit times were insignificant (data not shown).

### Table 2 Percent suppression of rod b-wave amplitude recovery rate prebleach and 30 min postbleach at month 1, relative to baseline*

<table>
<thead>
<tr>
<th>Percent suppression</th>
<th>Emixustat 2.5 mg (N=6)</th>
<th>Emixustat 5 mg (N=7)</th>
<th>Emixustat 10 mg (N=6)</th>
</tr>
</thead>
<tbody>
<tr>
<td>at month 1</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Prebleach</td>
<td>Postbleach</td>
<td>Prebleach</td>
</tr>
<tr>
<td>n</td>
<td>6</td>
<td>6</td>
<td>7</td>
</tr>
<tr>
<td>Mean (SD)</td>
<td>−18.85 (47.02)</td>
<td>−3.31 (41.43)</td>
<td>4.19 (22.12)</td>
</tr>
<tr>
<td>SE</td>
<td>19.20</td>
<td>16.91</td>
<td>8.36</td>
</tr>
<tr>
<td>Median</td>
<td>−11.03</td>
<td>−12.23</td>
<td>−3.07</td>
</tr>
<tr>
<td>Minimum, maximum</td>
<td>−80.7 to 37.5</td>
<td>−41.0 to 54.7</td>
<td>−22.8 to 35.3</td>
</tr>
</tbody>
</table>

* A positive percent suppression indicates a decrease from baseline and a negative percent suppression indicates an increase from baseline.
The majority of AEs were ocular (81.6%), with a similar proportion of ocular AEs across all treatment groups. The most common AEs were delayed dark adaptation (11/23, 47.8%), erythropsia (5/23, 21.7%), vision blurred (4/23, 17.4%) and photophobia and visual impairment (3/23, 13.0%) (table 3). Delayed dark adaptation was more frequent in the 5 mg and 10 mg groups. Headache was the only non-ocular treatment-emergent AE (TEAE) reported in >1 subject (present in two subjects). Of the 49 AEs, 39 were considered related to the study drug; the number of study drug-related AEs and proportion of subjects experiencing study drug-related AEs were comparable between subjects in the 5 mg and 10 mg groups and lower in subjects in the 2.5 mg group.

One subject, in the 5 mg group, experienced 3 TEAEs (delayed dark adaptation, night blindness, vision blurred), which led to study discontinuation; these TEAEs were of mild severity and were considered related to study drug. No subject experienced a serious AE during the study.

With regard to changes from baseline in safety assessments—including slit-lamp biomicroscopy, dilated ophthalmoscopy, intraocular pressure, clinical laboratory tests, physical examinations, vital signs or ECGs—few clinically relevant findings were observed. Across the study, mean and median changes in BCVA from baseline were small, with a range of −11 to +9 letters.

**DISCUSSION**

The primary objective of this study was to characterise the pharmacodynamics of emixustat in subjects with MA secondary to STGD. After being randomised to 1 month of daily treatment with emixustat at one of three doses (2.5 mg, 5 mg or 10 mg), subjects exhibited dose-dependent suppression of rod b-wave amplitude recovery postphotobleaching, with near-complete suppression at 10 mg but minimal to no suppression at 2.5 mg. This finding provides evidence of dose-dependent inhibition of RPE65 with emixustat in this population. Meaningful changes were not observed in single-flash cone response or 31 Hz flicker cone response. The lack of cone effect under these conditions may be attributable to the fact that cones also receive cis-related macular degeneration.13 14 As evidenced by dose-dependent suppression of rod recovery on ERG after photobleaching in these studies, the effects of emixustat on rod ERG were reversed after drug cessation; no effects on cone function were demonstrated on or off the drug.

The safety profile observed was also consistent with prior clinical trials in both healthy volunteers and patients with geographic atrophy secondary to age-related macular degeneration.14 15 23 26 The most common AEs in this study were ocular in nature, and the most common ocular AEs (delayed dark adaptation, erythropsia, vision blurred, photophobia, visual impairment) are consistent with emixustat’s mechanism of action (inhibition of RPE65, resulting in decreased availability of 11-cis-retinal to photoreceptors). Consistent with the lack of effect on ERG with 2.5 mg, there were lower incidences of delayed dark adaptation and study drug-related AEs in the 2.5 mg group.

On the basis of demographic and baseline characteristics, the population studied in this trial appears to be representative of STGD patients more generally, though the subjects in this study may be older than the average STGD patient enrolled in clinical trials. In the recent ProgStar study, the largest natural history study of STGD carried out to date, mean age at study entry was 33 years.27 The mean age of subjects in the current study was 52 years old. In terms of race and ethnicity, subjects in this study were similar to those enrolled in ProgStar. In addition, the areas of decreased autofluorescence on RI-FAF in this study fell within the range seen in ProgStar.

Limitations of this study include the small sample size and the inherent, significant intersubject and intrasubject variability observed in ERGs.22 These issues, in addition to the significant genetic and phenotypic variability across the STGD population,27 could limit the ability to apply the results of this study to the broader STGD population. However, the results of this study did demonstrate clear differences between the dose groups.

The results of this study informed the dose selection for an ongoing, 24-month clinical trial [Safety and efficacy of Emixustat in STARgardt disease (SeaSTAR) Study] comparing emixustat to placebo for the treatment of MA secondary to STGD (NCT03772665). This phase 3, randomised, double-masked, international trial is now investigating the effects of a 10 mg daily dose of emixustat on the mean rate of change in subjects’ total area of MA. If emixustat is shown to slow the growth of MA secondary to STGD disease, it could represent the first effective treatment for this condition.

**Acknowledgements**

The authors thank the Protocol 4429-204 investigators: Paul Bernstein (Moran Eye Center; University of Utah, Salt Lake City, Utah, USA), Karl Csaky (Retina Foundation of the Southwest, Dallas, Texas, USA), Kanishka Thiran Jayasundera (W.K. Kellogg Eye Center, University of Michigan, Ann Arbor, Michigan, USA), Byron Lam (Bascom Palmer Eye Institute, University of Miami-Miller School of Medicine, Miami, Florida, USA), Mandeep Singh (Wilmer Eye Institute, Johns Hopkins University, Baltimore, Maryland, USA), and Elias Traboulsi (Cole Eye Institute, Cleveland Clinic, Cleveland, Ohio, USA). The authors also thank Kristin Harper, PhD, of Harper Health & Science Communications, for providing writing and editorial support in accordance with Good Publication Practice (GPP3) guidelines.

**Contributors**

All authors contributed to the research design of the study, the analysis and interpretation of the study results, and the revision and review of the manuscript. JG contributed to study administration and supervision, and JMK contributed to the data analysis. All authors approved the final manuscript as submitted and agree to be accountable for all aspects of the work.

**Funding**

Funding for the study was provided solely by Kubota Vision. This study was registered with ClinicalTrials.gov as NCT03033108.

**Competing interests**

RK, JKG and JMK are employees of Kubota Vision. RK has pending and issued patents related to emixustat hydrochloride. DGB received grants from Kubota Vision during the conduct of the study.

**Patient consent for publication**

Not required.

**Ethics approval**

The protocol and informed consent forms were prospectively approved by either a central institutional review board (IRB; Chesapeake IRB) or site-specific IRBs (University of Utah IRB, Johns Hopkins University School of Medicine IRB, and the Cleveland Clinic IRB).

**Provenance and peer review**

Not commissioned; externally peer reviewed.

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**Table 3**

<table>
<thead>
<tr>
<th>Adverse event</th>
<th>Emixustat 2.5 mg (N=7)</th>
<th>Emixustat 5 mg (N=9)</th>
<th>Emixustat 10 mg (N=7)</th>
<th>All subjects (N=23)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Any adverse event</td>
<td>6 (85.7)</td>
<td>8 (88.9)</td>
<td>7 (100)</td>
<td>21 (91.3)</td>
</tr>
<tr>
<td>Delayed dark adaptation</td>
<td>1 (14.3)</td>
<td>6 (66.7)</td>
<td>4 (57.1)</td>
<td>11 (47.8)</td>
</tr>
<tr>
<td>Erythropsia</td>
<td>1 (14.3)</td>
<td>3 (33.3)</td>
<td>1 (14.3)</td>
<td>5 (21.7)</td>
</tr>
<tr>
<td>Vision blurred</td>
<td>2 (28.6)</td>
<td>1 (11.1)</td>
<td>1 (14.3)</td>
<td>4 (17.4)</td>
</tr>
<tr>
<td>Photophobia</td>
<td>1 (14.3)</td>
<td>2 (22.2)</td>
<td>0</td>
<td>3 (13.0)</td>
</tr>
<tr>
<td>Visual impairment</td>
<td>1 (14.3)</td>
<td>1 (11.1)</td>
<td>1 (14.3)</td>
<td>3 (13.0)</td>
</tr>
</tbody>
</table>

*Percentages were based on the total number of subjects enrolled in each treatment group for the safety analysis set (N).
REFERENCES


