

Rebound macular edema following oral acetazolamide therapy for juvenile X-linked retinoschisis in an Italian family

Maria Silvana Galantuomo^{1,*}
Maurizio Fossarello¹
Alberto Cuccu¹
Roberta Farci¹
Markus N Preising²
Birgit Lorenz²
Pietro Emanuele Napoli^{1,*}

¹Department of Surgical Sciences, Eye Clinic, University of Cagliari, Cagliari, Italy; ²Department of Ophthalmology, Faculty of Medicine, Justus-Liebig-University, Giessen, Germany

*These authors contributed equally to this work

Background: Juvenile X-linked retinoschisis (RS1, OMIM: 312700) is a hereditary vitreoretinal dystrophy characterized by bilateral foveal schisis and, in half of the patients, splitting through the nerve fiber layer in the peripheral retina. In the first decade of life, patients usually develop a decrease in visual acuity. Long-term visual outcomes can be poor due to the limited number of known successful treatments.

Purpose: The purposes of this study were to present, for the first time, a p.Arg197Cys missense mutation in the *RS1* gene (OMIM: 300839) in a four-generation Italian family with RS1 and to examine the clinical response to the treatment with acetazolamide tablets alone or in combination with dorzolamide eye drops as assessed by spectral-domain optical coherence tomography (SD-OCT).

Methods: Eleven individuals, including two brothers with RS1 (patients 1 and 2), underwent a full medical history examination and a comprehensive ocular assessment that involved SD-OCT, fluorescein angiography, electroretinography and DNA analysis. Each RS1 patient received oral acetazolamide (375 mg daily) during the first three months. Thereafter, patient 1 continued only with dorzolamide eyedrops three times a day for a period of three months, while patient 2 spontaneously stopped both medications.

Results: Sequence analysis of the *RS1* gene identified a hemizygous c.589C>T (p.Arg197Cys) missense mutation in exon 6, which has not been previously reported in an Italian family. A different response to the medical therapy was observed in the four eyes of the two affected brothers hemizygous for this abnormality. Of note, after acetazolamide interruption, a rebound effect on cystoid macular edema reduced the beneficial effects of the initial therapy for RS1 from p.Arg197Cys mutation. Indeed, a minimal rebound effect on cystoid macular edema, and an improvement in visual acuity, was observed in patient 1 during the six months of treatment. Conversely, in patient 2, an initial improvement in cystoid macular edema was not associated with visual acuity changes, followed by a marked rebound effect.

Conclusion: This study showed that the sequential use of acetazolamide tablets and dorzolamide eye drops should be considered and studied further as a possible treatment for macular edema and visual impairment in patients with RS1 from a hemizygous p.Arg197Cys mutation.

Keywords: juvenile X-linked retinoschisis, oral acetazolamide, topical dorzolamide, cystoid macular edema, macular schisis, foveal zone thickness

Correspondence: Pietro Emanuele Napoli
Department of Surgical Sciences,
University of Cagliari, via Ospedale 46,
Cagliari 09124, Italy
Tel +39 320 864 3697
Email pietronapoli@gmail.com

Introduction

Juvenile X-linked retinoschisis (RS1) is a hereditary vitreoretinal disorder that shows radiating cystoid changes in the macula with or without peripheral retinoschisis, early onset of visual loss, abnormal electroretinograms (ERGs) and, in some cases,

vitreous hemorrhage and/or retinal detachment. RS1 affects predominantly male subjects. It is transmitted as an X-linked recessive trait and female carriers are usually asymptomatic. The penetrance of RS1 is almost complete but clinical expression is highly variable.¹ As previously reported, the prevalence of RS1 is between 1:5,000 and 1:25,000.^{2,3} Visual impairment is usually mild until the fourth decade of life; thereafter, progressive visual deterioration often occurs.

The *RS1* gene (OMIM: 300839) maps to Xp22.13, has six exons and encodes retinoschisin, a secreted 224 amino acid protein associated with the disease, abundantly expressed in both cone and rod photoreceptors and bipolar cells of the retina and the pineal gland.

The gene contains a highly conserved region in exons 4–6 known as the discoidin domain, shared with a number of other proteins.⁴ The discoidin domain is implicated in cell–cell adhesion and phospholipid binding.⁵ Cell expression studies showed that disease-associated missense mutations result in aberrant protein synthesis and retention in the endoplasmic reticulum, producing non-functional products.

The ERGs of the most affected males demonstrate normal or near-normal *a*-waves, characteristic of photoreceptor function, but often substantially reduced *b*-waves, originating from inner retinal cell activity.⁶ For several decades, the ERG has been the major diagnostic technique for RS1.

Relatively recently, optical coherence tomography (OCT) has changed the diagnostic approach for various ocular diseases in both anterior and posterior segments,^{7–12} including RS1. Indeed, at present spectral domain OCT (SD-OCT) is the most useful diagnostic tool for this disease. Even in a non-cooperative child, a single OCT B-scan of the foveal area is sufficient to detect RS1 signs and to make a differential diagnosis with other diseases causing visual loss in young boys.

Long-term visual outcomes can be poor due to the limited number of known successful treatments. A beneficial effect from the use of a topical or, alternatively, systemic carbonic anhydrase inhibitor (CAI) has been reported in patients with cystoid macular edema in retinal dystrophies including RS1. One study described the successful treatment of a patient with a p.Arg197Cys, and other mutations, in RS1.¹³ No follow-up was reported in this study after the discontinuation of the therapy with acetazolamide. Further studies on CAI treatment showed a rebound effect in RS1 in a small subset of patients only.¹⁴ A different response to treatment with oral CAI might be possible for various genotypes in RS1 (including the side effects), as previously reported.^{15–18}

The purposes of this study were, therefore, to report for the first time a p.Arg197Cys missense mutation in the *RS1*

gene in an Italian family with RS1 and to explore the clinical response, as assessed by SD-OCT, in two affected brothers hemizygous for this mutation, to the treatment with oral acetazolamide alone or in combination with dorzolamide eye drops.

Materials and methods

Subjects and study procedure

This study was conducted at the Eye Clinic of the Department of Surgical Sciences, University of Cagliari, Italy. The study adhered to the tenets of the Declaration of Helsinki and was approved by the Office of Research Ethics, University of Cagliari. All subjects signed informed consents prior to their participation in the study.

Individuals from a four-generation family with RS1 were recruited at Budduso, a village in Olbia Province, Italy. The family consisted of 20 members with four affected subjects (Figure 1). Eleven individuals, including two brothers with RS1, 18 and 20 years old (patients 1 and 2, respectively), underwent a full medical history examination and a comprehensive ocular assessment, which involved best-corrected visual acuity, slit lamp examinations, fundus examination in mydriasis, SD-OCT scans, fluorescein angiography, electrophysiology and genetic analysis.

Patients were diagnosed as having RS1 based on the clinical findings, which included a decrease in visual acuity (VA), stellate-shaped cavities in the macular region on the retinal exam, and a decrease in the *b*-wave amplitude on ERG examination. The diagnosis was confirmed by molecular genetic analysis of *RS1* gene, which revealed a p.Arg197Cys missense mutation (Figure 1).

Genetic investigation

Peripheral blood samples with EDTA anticoagulant were collected from each participant of the study. Genomic DNA was extracted using GeneCatcher™ gDNA Automated Blood Kit. All six exons and the flanking intronic regions of the *RS1* gene were amplified by polymerase chain reaction with oligonucleotide primers published.⁴ The PCR products were analyzed on 2% agarose gel stained with ethidium bromide and purified with a DNA extraction kit (QIA-quick Gel Extraction Kit; Qiagen, Hilden, Germany). PCR products from all exons were directly sequenced with the Bigdyes termination method.

Clinical assessment

Both patients with RS1 from p.Arg197Cys mutation (patients 1 and 2) were orally treated with 375 mg acetazolamide

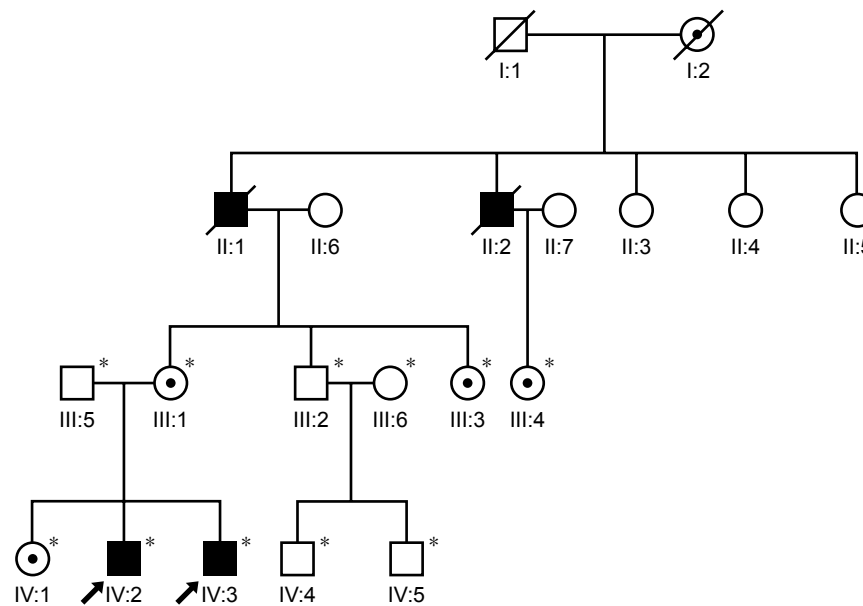


Figure 1 Family pedigree of an Italian family with juvenile X-linked retinoschisis from a p.Arg197Cys mutation (village of Budduso, Sardinia).

Notes: *Individuals genotyped for *RS1*. Unaffected individuals are represented by empty symbols. Solid squares denote affected males. Female carriers, when identified, are indicated by a shade dot inside a circle. Individuals crossed by a line represents already passed away. Each generation is identified by a Roman numeral and each individual within the same generation is numbered consecutively by an Arabic numeral. The patients are indicated by an arrow.

tablets once daily simultaneously and the treatment was maintained for three months (Diamox; Teofarma s.r.l., Via F.lli Cervi, 8 27010 Valle Salimbene, PV, Italy). At the end of the 3rd month, patient 1 continued therapy only with 2% dorzolamide collyrium (Trusopt; Santen Switzerland SA, Geneva, Switzerland) three times daily in both eyes, while the older brother (patient 2) refused any further treatment.

The response to the treatment was monitored by measuring VA and by performing SD-OCT (Cirrus HD-OCT 4000; Carl Zeiss Meditec, Inc., Dublin, CA, USA) using the macular cube 512×128 and HD 5 line raster scanning protocol, at baseline, at the end of the 3rd and 6th months. Changes in the foveal zone thickness (FZT), as detected in the central subfield of the macular thickness map (ie, in the central 1,000 μm of the foveal region in the ETDRS grid, from ILM to RPE), were calculated from the start to the end of the medical treatment. As previously reported, more than 19.6% reduction in FZT was considered to be substantial, as well as a change in logMAR score of 0.12 (based on previously reported normal variability between visits in non-treated eyes).¹⁹

Results

Sequence analysis of the *RS1* gene identified a hemizygous c.589C>T (p.Arg197Cys) missense mutation in exon 6. The patients' mother and the unaffected sister were heterozygous for the mutation c.589C>T. The nucleotide change in

the patients predicted an amino acid change from arginine (CGC) to cysteine (TGC). This mutation was previously reported in a British patient but not in patients with Italian ethnicity.^{13,20}

Fluorescein angiography showed that macular schisis cavities were not associated with late leakage. ERG examination showed that both patients had a significantly reduced rod response and decreased cone responses for the amplitudes of *a*-waves and *b*-waves.

A different response to the medical therapy was observed in the four eyes of the two brothers on examination by SD-OCT. Relative changes were calculated with respect to baseline values (Tables 1 and 2).

Patient 1 underwent complete therapy. At the start of treatment, the FZT was 563 μm in the right eye (RE) and 501 μm in the left eye (LE), and VA was 3/10 (logMAR 0.522) in both

Table 1 Changes in foveal zone thickness

	Baseline	Three months	Six months
Patient 1			
Right eye (μm)	563	430	440
Left eye (μm)	501	371	277
Patient 2			
Right eye (μm)	496	377	509
Left eye (μm)	519	455	501

Notes: Patient 1: Underwent complete therapy (sequential treatment with acetazolamide tablet and dorzolamide eye drops). Patient 2: Spontaneously interrupted any therapy after oral acetazolamide.

Table 2 Changes in visual acuity

	Baseline	3 months	6 months
Patient 1			
Right eye	3/10 (logMAR 0.522)	6/10 (logMAR 0.221)	6/10 (logMAR 0.221)
Left eye	3/10 (logMAR 0.522)	5/10 (logMAR 0.301)	5/10 (logMAR 0.301)
Patient 2			
Right eye	4/10 (logMAR 0.397)	4/10 (logMAR 0.397)	4/10 (logMAR 0.397)
Left eye	5/10 (logMAR 0.301)	5/10 (logMAR 0.301)	5/10 (logMAR 0.301)

Notes: Patient 1: Underwent complete therapy (sequential treatment with acetazolamide tablet and dorzolamide eye drops). Patient 2: Spontaneously interrupted any therapy after oral acetazolamide.

eyes (OU) on the Snellen chart. After the first three months of acetazolamide treatment (Figure 2), the FZT was 430 μ m in the RE (−23.62%) and 371 μ m in the LE (−25.94%), and VA had improved to 6/10 in the RE (logMAR 0.221) and 5/10 in the LE (logMAR 0.301). At the end of the 6th month, after treatment with 2% dorzolamide three times a day for three months, the FZT was 440 μ m in the RE (−21.84%) and 277 μ m in the LE (−44.71%), and VA remained 6/10 in the RE (logMAR 0.221) and 5/10 in the LE (logMAR 0.301) (Table 1).

In patient 2, the FZT was 496 μ m in the RE and 519 μ m in the LE, and VA was 4/10 (logMAR 0.397) in the RE and 5/10 (logMAR 0.301) in the LE before treatment. After acetazolamide treatment, the FZT was 377 μ m in RE (−23.99%) and 455 μ m in the LE (−12.33%), but VA was unchanged in OU. At the end of the 6th month, the FZT was 509 μ m in the RE (+2.62%) and 501 μ m in the LE (−3.46%), and VA was unchanged in OU (Table 2).

Discussion

RS1 is caused by mutations of the *RS1* gene that encodes for retinoschisin. Retinoschisin promotes cell adhesion and its dysfunction leads to the formation of cystoid cavities in

the inner nuclear and outer plexiform layers of the retina.¹⁵ Clinical effects of CAI are presumed to be attributed to their facilitation effect on the fluid transport across the retinal pigment epithelium, decreasing the volume of the subretinal space and increasing retinal adhesiveness.²¹ Apushkin and Fishman¹⁹ in their study demonstrated a reduction in the foveal thickness in seven of the eight patients affected by RS1, who were treated with 2% dorzolamide eye drops, and a concomitant improvement in the VA in five patients.¹⁹ In a further small case study, it was found that the response to dorzolamide eye drops was independent of the type of the underlying mutation.¹⁵

To date, 196 different mutations in the *RS1* gene are known to cause RS1 (Leiden Open Variation Database, LOVD, version 2.0, Build 35; http://grenada.lumc.nl/LOVD2/eye/home.php?select_db=RS1). More than 80% are nucleotide substitutions, with missense mutations being the predominant type, mostly found in exons 4–6 encoding the discoidin domain. However, the functional impact of most missense variants is not well known. In general, a lack of relationship between the response to dorzolamide and the type of mutation in *RS1* gene has been observed by different authors.^{13,15} In particular, structure-based analysis of

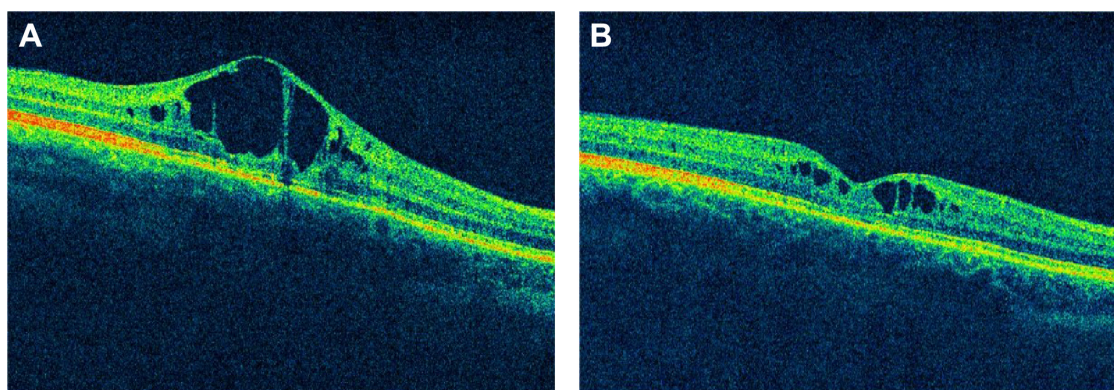


Figure 2 Spectral-domain optical coherence tomography (SD-OCT) imaging in patient 1 with juvenile X-linked retinoschisis from p.Arg197Cys mutation during the first 3 months of therapy with oral acetazolamide (Cirrus HD-OCT 4000; Carl Zeiss Meditec, Inc., Dublin, CA, USA). (A) Baseline (pretreatment) OCT scan: visual acuity was 3/10 and foveal zone thickness 563 μ m. (B) At the end of the 3rd month, OCT scan showed a significant improvement in visual acuity (6/10) and in foveal zone thickness (430 μ m).

p.Arg197Cys mutation predicts a mild change in retinoschisin protein, which should not affect the protein fold.²²

A case report on the use of oral acetazolamide in an 8-year-old child diagnosed with RS1 demonstrated a near complete resolution of macular edema, with an increased VA even when the macular edema recurred on cessation of treatment.¹⁴

Long-term treatment of oral acetazolamide may be contraindicated in some patients because of potential systemic side effects, such as electrolyte imbalance, weakness, renal dysfunction, depression, metabolic acidosis, and mental lethargy. Because of these reasons, a topical medication would be preferable for therapy over a prolonged period of time. Clearly, topical dorzolamide may also lead to side effects, which are prevalently local, such as burning sensation of eyes, eye pruritus/discomfort, and local ocular hypersensitivity reaction.

Our study showed that oral treatment of acetazolamide for 3 months substantially reduced macular thickness in both patients and increased VA in one (patient 1), without any systemic adverse drug effects. The data of our study also showed that after the cessation of acetazolamide treatment, a rebound effect on cystoid macular edema may reduce the beneficial effects of initial therapy for RS1 from p.Arg197Cys mutation. This aspect highlights the difficulties in choosing the correct treatment strategy after the cessation of oral acetazolamide.

Of note, in patient 1 (who underwent subsequent treatment with dorzolamide eyedrops) the appearance of macular thickening was limited by sustained therapy with topical CAI, avoiding the return to the pretreatment levels (baseline values). In fact, the sequential use of topical 2% dorzolamide resulted only in a slight increase in FZT without any changes in VA. Overall, during the 6 months of therapy, sequential administration of CAI, orally and topically, induced a significant improvement in both cystoid macular edema and VA, without any systemic adverse drug effects.

Conversely, in patient 2, the initial improvement in cystoid macular edema was not associated with VA changes (during therapy with oral acetazolamide), followed by a marked rebound effect after cessation of the therapy. Whether the absence of an effect on VA predicts the rebound effect is speculative since nonresponders in VA to CAI treatment have been reported previously.^{16,17,23}

Conclusion

In conclusion, the results of this study suggest that cessation of oral acetazolamide may result in a rebound effect on

cystoid macular edema in patients with RS1 from a specific gene mutation in the *RS1* (c.589C>T, p.Arg197Cys).

Also, the combined use of oral and topical CAI should be considered and studied further for a possible treatment of maculopathy and visual impairment in affected subjects with this abnormality. In this sense, CAI withdrawal should be gradual, and the dilution of the medication with eyedrops may be desirable. Consequently, until long-term efficacy and safety of a proper treatment protocol for XLRS maculopathy is established, we recommend caution in suddenly interrupting the therapeutic regimen with CAI.

Disclosure

The authors report no conflicts of interest in this work.

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