X-linked retinoschisis: clinical phenotype and RS1 genotype in 86 UK patients

D Pimenides, N D L George, J R W Yates, K Bradshaw, S A Roberts, A T Moore, D Trump

Background: Inactivating mutations of the gene RS1 lead to X-linked retinoschisis, a progressive retinal dystrophy characterised by schisis within the inner layers of the neuroretina. The mutation spectrum is large and the phenotype variable.

Aim: To determine whether there is a correlation between mutation type and disease severity.

Methods: We identified the causative mutation in 86 affected patients and examined each of these patients in detail. Different categories of mutation were compared for each phenotypic characteristic.

Results: We found a reduction in visual acuity with increasing age and worsening macular pathology in patients over 30 years old (p < 0.001), but there was no correlation between mutation type and severity of disease. Furthermore, we found a wide variation in phenotype even within families.

Conclusions: Identifying the causative mutation in patients with X-linked retinoschisis is helpful in confirming diagnosis and in counselling of family members but cannot be used to predict prognosis for an individual patient.

METHODS

Patients

A total of 86 UK patients were included in the study. Each had previously been diagnosed with XLRS and all were re-examined by one of three ophthalmologists (DP, NDLG, ATM). Five had no known family history of XLRS, while the others came from 29 families with a history of XLRS. The causative mutation for each patient had either already been identified or was identified as part of the study. All patients gave consent to participate in the study.

Ophthalmological examination

The refractive error in each eye and best corrected distant and near visual acuities were recorded. Ocular motility and pupil reactions were assessed, anterior segment examination performed with a slit lamp, intraocular pressures measured, and posterior segment examination performed with indirect ophthalmoscopy and slit lamp biomicroscopy, following pupil dilation. FERG recordings were obtained in 43 of the 86 patients and were consistent with a diagnosis of XLRS.

Genetic studies

In a proportion of the patients studied the mutation had already been identified as part of a previous study. For the other patients the causative mutation was identified as previously described.

Statistical methods

The age of the patients was recorded at the time of clinical examination. Five characteristics were chosen to describe the phenotypic characteristic (i) best corrected visual acuity, (ii) best corrected near vision, (iii) presence of foveal schisis or other foveal pathology in one or both eyes, (iv) peripheral schisis in one or both eyes, and (v) complications (vitreous haemorrhage or retinal detachment).

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Abbreviations: DVA, distant visual acuity; FERG, full field electroretinogram; NVA, near visual acuity; XLRS, X-linked juvenile retinoschisis

ac.uk/uwcm/mg/hgmd0.html). The majority of these are missense mutations in exons 4–6 but also include nonsense and splice site mutations, deletions, and insertions. Many missense mutations have been shown to prevent secretion of the mutant protein leading instead to intracellular degradation.

There have been reports comparing genotype and phenotype. No correlation has been found between mutation type and disease severity, but these studies have been limited by small patient numbers. We report here the results of our study of 86 patients with XLRS in whom the causative RS1 mutation has been identified.
they affected a residue within the discoidin domain, secretory leader peptide, or the remaining part of the gene, (iii) whether they affected a cysteine residue, (iv) the exon in which they occurred, (v) the mutation type: protein truncating or missense, (vi) whether the mutations interfered with protein secretion or allowed secretion into the extracellular space, and (vii) if successfully secreted, whether the mutations interfered with oligomerisation.

Each mutation category was compared to each phenotypic characteristic, using the $\chi^2$ test of independence. Phenotypes of patients from the same family and phenotypes of unrelated patients with the same mutation and of similar age, were compared.

For comparison of age and visual acuity, data were compared with those of a previous study.\textsuperscript{2} We used linear regression using the log transformed visual acuity and compared these data sets with an analysis of covariance, where we allowed for age as a linear covariate and tested for a difference in slopes by testing the interaction between age and study.

RESULTS

A total of 86 patients fulfilled the inclusion criteria and the diagnosis was supported by the FERG findings for 43 of these. Genotype details are summarised in table 1.

Phenotype

The age range was 2–69 years with a distribution skewed towards a younger age (mean age 23.8 years, median 34.5 years). A total of 25 (29%) of our patients were younger than 10 years, 23 (27%) were between 11 and 20, 11 (13%) were between 21 and 30, and only 27 were older than 30 years (31%) (fig 1A).

Distant (DVA) and near (NVA) visual acuity were measured for the patients (fig 1B–D). DVA varied from 6/6 to 6/60 and NVA from N.5 to N.24. Visual acuity appears to worsen with age over the whole group, but there is a large variation in severity within a patient only 5 years old with a DVA of 6/60 and another 47 years old with a DVA of 6/12. We found no correlation between visual acuity and complications or degree of peripheral schisis.

All patients had evidence of maculopathy. Forty eight patients (56%) had typical bilateral foveal schisis, 18 (21%) had foveal schisis in one eye and other maculopathy in the fellow eye, and 18 patients (21%) had other maculopathy in both eyes. Two patients had dense unilateral cataracts, not allowing fundal view of one eye, but had typical foveal schisis in the fellow eye. Patients over 30 years of age had more severe macular changes ($p<0.001$) which may explain the decrease in visual acuity with age.

Bilateral peripheral retinoschisis was detected in 42 patients (49%). Fifteen patients (17%) had peripheral schisis in one eye and either no schisis or retinal detachment in the fellow eye. No peripheral schisis was found in the eyes of 27 (31%) patients. There was no difference in the proportions of patients at a younger or older age with evidence of peripheral schisis. Retinal detachment or vitreous haemorrhage had occurred in 26 patients (30%) and affected both eyes of 11 patients and only one eye of 15. The proportions of younger and older patients who developed complications were similar.

A previous study of retinoschisis had reported no change in visual acuity with age in patients without complications.\textsuperscript{4} Sixty of our 86 patients had no complications and in this group we still found an association of reduced visual acuity with increasing age suggesting that visual acuity decreases with time (linear regression on log transformed data, $p=0.0013$) perhaps due to a worsening of macular pathology.

Genotype

The causative mutation had been determined for each patient (table 1). Based on our previous molecular analysis of mutations in this protein,\textsuperscript{14,16} we were able to predict whether a particular missense mutation would lead to intracellular retention and degradation of retinoschisin or would allow secretion into the extracellular space (table 1). Only five mutations (seen in 18 patients) were predicted to lead to protein truncation.

We compared the clinical characteristics of groups of patients with similar mutations but found no evidence for a correlation between mutation type and disease severity even in patients of similar ages. We also compared visual acuity, foveal changes, peripheral schisis, and disease complications in patients with similar or different classes of mutations. We classified mutations into groups including protein truncating, missense, mutations in different exons, discoidin domain mutations and those affecting other residues, and mutations affecting cysteine residues and those affecting other residues.

### Table 1: Genotype details of patients

<table>
<thead>
<tr>
<th>Mutation</th>
<th>Effect of mutation</th>
<th>Exon</th>
<th>Discoid domain missense</th>
<th>Consequences of mutation*†</th>
<th>No. of patients (total 86)</th>
<th>No. of families with mutation</th>
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<tbody>
<tr>
<td>78G→C</td>
<td>E26*</td>
<td>2</td>
<td>No</td>
<td>Not tested</td>
<td>1</td>
<td>1</td>
</tr>
<tr>
<td>79→A/G</td>
<td>AspG splice acc</td>
<td>3</td>
<td>Intronic</td>
<td>Truncating</td>
<td>4</td>
<td>1</td>
</tr>
<tr>
<td>267T→A</td>
<td>Y92X</td>
<td>4</td>
<td>No</td>
<td>Truncating</td>
<td>8</td>
<td>1</td>
</tr>
<tr>
<td>304C→T</td>
<td>R102W</td>
<td>4</td>
<td>Yes</td>
<td>No secretion</td>
<td>16</td>
<td>6</td>
</tr>
<tr>
<td>305G→A</td>
<td>R102G</td>
<td>4</td>
<td>Yes</td>
<td>No secretion</td>
<td>4</td>
<td>1</td>
</tr>
<tr>
<td>308T→G</td>
<td>L103R</td>
<td>4</td>
<td>Yes</td>
<td>Not tested</td>
<td>4</td>
<td>1</td>
</tr>
<tr>
<td>325G→C</td>
<td>G109R</td>
<td>4</td>
<td>Yes</td>
<td>No secretion</td>
<td>7</td>
<td>2</td>
</tr>
<tr>
<td>326→TG→A</td>
<td>G63A donor sp</td>
<td>4</td>
<td>Intronic</td>
<td>Truncating</td>
<td>2</td>
<td>1</td>
</tr>
<tr>
<td>329G→A</td>
<td>C110Y</td>
<td>5</td>
<td>Yes</td>
<td>Not tested</td>
<td>1</td>
<td>1</td>
</tr>
<tr>
<td>419G→A</td>
<td>G140E</td>
<td>5</td>
<td>Yes</td>
<td>Not tested</td>
<td>4</td>
<td>1</td>
</tr>
<tr>
<td>421C→G</td>
<td>R141G</td>
<td>5</td>
<td>Yes</td>
<td>Reduced secretion</td>
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<td>1</td>
</tr>
<tr>
<td>421C→T</td>
<td>R141C</td>
<td>5</td>
<td>Yes</td>
<td>No secretion</td>
<td>14</td>
<td>5</td>
</tr>
<tr>
<td>460C→T</td>
<td>Q154X</td>
<td>5</td>
<td>Yes</td>
<td>Truncating</td>
<td>1</td>
<td>1</td>
</tr>
<tr>
<td>574C→A</td>
<td>P192A</td>
<td>6</td>
<td>Yes</td>
<td>Not tested</td>
<td>2</td>
<td>1</td>
</tr>
<tr>
<td>574C→T</td>
<td>P192S</td>
<td>6</td>
<td>Yes</td>
<td>Not tested</td>
<td>3</td>
<td>2</td>
</tr>
<tr>
<td>578C→T</td>
<td>P193L</td>
<td>6</td>
<td>Yes</td>
<td>Truncating</td>
<td>2</td>
<td>2</td>
</tr>
<tr>
<td>579→810insC</td>
<td>insC</td>
<td>6</td>
<td>Yes</td>
<td>Truncating</td>
<td>1</td>
<td>1</td>
</tr>
<tr>
<td>597C→G</td>
<td>I99F</td>
<td>6</td>
<td>Yes</td>
<td>Not tested</td>
<td>2</td>
<td>1</td>
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<tr>
<td>598C→T</td>
<td>R200C</td>
<td>6</td>
<td>Yes</td>
<td>Not tested</td>
<td>6</td>
<td>3</td>
</tr>
<tr>
<td>653G→T</td>
<td>C219G</td>
<td>6</td>
<td>Yes</td>
<td>Reduced oligomerisation</td>
<td>2</td>
<td>1</td>
</tr>
</tbody>
</table>

*Novel mutation; †mutation found as part of this study.
We found no significant correlation between mutation type and disease severity. We hypothesised that there may be a difference in disease characteristics between those patients predicted to have retinoschisin present within their retinal extracellular space (that is, mutant protein with a missense change but still allowing secretion) and those predicted to have no secreted protein (truncating mutations and missense changes shown to cause intracellular degradation of protein). However, again, we found no significant difference in any of the clinical measures of disease severity.

Intrafamilial variation and phenotypic variation in patients of similar ages with the same mutation
We reviewed the phenotypic characteristics in families where we had access to large number of affected males. For two families we were able to examine six affected relatives and found the severity of disease was still very variable. We also compared young boys with the same mutation and again found a great deal of variation, for example, in boys with the R102W missense mutation originating from different families the visual acuity varied from 6/9 (11 year old) to 6/24 (10 and 14 year old).

DISCUSSION
The availability of genetic testing for families with X-linked retinoschisis has allowed accurate diagnosis and also improved genetic counselling for patients and relatives. Women at risk of being a carrier for this disorder can now be offered molecular genetic testing to determine whether they have an RS1 mutation. Sons of women who carry the mutation can be offered mutation testing to determine whether they will develop the disease.

Molecular genetic testing allows the specific mutation to be identified and could be used to predict long term prognosis if there is a relationship between disease phenotype and type of mutation. There have been several studies of the phenotypic variation in X-linked retinoschisis and comparisons made between patients with the same mutation and in groups of patients with different mutations, but as these studies are small limited conclusions can be drawn. We have studied a large group of patients with XLRS in whom the mutation has been identified and have related the mutations to our current knowledge about the molecular mechanisms thought to contribute to disease. We found a wide variation in clinical phenotype, both between and within families, which was not
dependent on mutation type. Retinoschisin is a secreted protein thought to function within the extracellular retinal environment in cell adhesion. Many missense mutations lead to intracellular retention of the mutant protein leading to an absence of extracellular retinoschisin. However the severity of the disease in such cases is no different to those with mutations allowing secretion of retinoschisin. This, together with the variation seen within in families, suggests there are other factors influencing disease severity such as genetic modifiers (for example, genetic variation in interacting proteins) or environmental influences.

Molecular genetic testing in XLRS has facilitated diagnosis in atypical cases and has improved genetic counselling for families but unfortunately is of no value in predicting long term visual prognosis.

ELECTRONIC-DATABASE INFORMATION

The X-linked Retinoschisis sequence variation database can be found at http://www.dmd.nl/rls/, and the Human Gene Mutation Database at the Institute of Medical Genetics in Cardiff can be found at http://archive.uwm.ac.uk/uwmc/mgm/d0.html.

Authors’ affiliations
D Pimenides, J R W Yates, D Trump, Department of Medical Genetics, University of Cambridge and Addenbrooke’s NHS Hospital Trust, Cambridge, UK
D Pimenides, K Bradshaw, Department of Ophthalmology, Addenbrooke’s NHS Hospital Trust, Cambridge, UK
N D L George, Department of Ophthalmology, Ninewells Hospital, Dundee, UK
S A Roberts, Biostatistics Group, Division of Epidemiology and Health Sciences, Faculty of Medical and Health Sciences, University of Manchester, UK
A T Moore, Institute of Ophthalmology and Moorfields Eye Hospital, London, UK
D Trump, Academic Unit of Medical Genetics, St Mary’s Hospital and Centre for Molecular Medicine, Faculty of Medical and Health Sciences, University of Manchester, UK

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