Two Novel Mutations in the EYS Gene Are Possible Major Causes of Autosomal Recessive Retinitis Pigmentosa in the Japanese Population

Katsuhiro Hosono1*, Chie Ishigami2, Masayo Takahashi2, Dong Ho Park3, Yasuhiko Hirami4, Hiroshi Nakanishi5, Shinji Ueno6, Tadashi Yokoi7, Akiko Hikoya1, Taichi Fujita1, Yang Zhao1,8, Sachiko Nishina7, Jae Pil Shin3, In Taek Kim3, Shuchi Yamamoto6, Noriyuki Azuma3, Hiroko Terasaki6, Miho Sato1, Mineo Kondo6, Shinsei Minoshima8, Yoshihiro Hotta1

1 Department of Ophthalmology, Hamamatsu University School of Medicine, Hamamatsu, Japan, 2 Laboratory for Retinal Regeneration, RIKEN Center for Developmental Biology, Kobe, Japan, 3 Department of Ophthalmology, Kyungpook National University Hospital, Daegu, Korea, 4 Department of Ophthalmology, Institute of Biomedical Research and Innovation Hospital, Kobe, Japan, 5 Department of Otolaryngology, Hamamatsu University School of Medicine, Hamamatsu, Japan, 6 Department of Ophthalmology, Nagoya University Graduate School of Medicine, Nagoya, Japan, 7 Department of Ophthalmology and Laboratory of Cell Biology, National Center for Child Health and Development, Tokyo, Japan, 8 Department of Photomedical Genomics, Basic Medical Photonics Laboratory, Medical Photonics Research Center, Hamamatsu University School of Medicine, Hamamatsu, Japan, 9 Department of Ophthalmology and Visual Science, Chiba University Graduate School of Medicine, Chiba, Japan

Abstract

Retinitis pigmentosa (RP) is a highly heterogeneous genetic disease including autosomal recessive (ar), autosomal dominant (ad), and X-linked inheritance. Recently, arRP has been associated with mutations in EYS (Eyes shut homolog), which is a major causative gene for this disease. This study was conducted to determine the spectrum and frequency of EYS mutations in 100 Japanese arRP patients. To determine the prevalence of EYS mutations, all EYS exons were screened for mutations by polymerase chain reaction amplification, and sequence analysis was performed. We detected 67 sequence alterations in EYS, of which 21 were novel. Of these, 7 were very likely pathogenic mutations, 6 were possible pathogenic mutations, and 54 were predicted non-pathogenic sequence alterations. The minimum observed prevalence of distinct EYS mutations in our study was 18% (18/100, comprising 9 patients with 2 very likely pathogenic mutations and the remaining 9 with only one such mutation). Among these mutations, 2 novel truncating mutations, c.4957_4958insA (p.S1653fsX2) and c.8868C>A (p.Y2956X), were identified in 16 patients and accounted for 57.1% (20/35 alleles) of the mutated alleles. Although these 2 truncating mutations were not detected in Japanese patients with adRP or Leber’s congenital amaurosis, we detected them in Korean adRP patients. Similar to Japanese arRP results, the c.4957_4958insA mutation was more frequently detected than the c.8868C>A mutation. The 18% estimated prevalence of very likely pathogenic mutations in our study suggests a major involvement of EYS in the pathogenesis of arRP in the Japanese population. Mutation spectrum of EYS in 100 Japanese patients, including 13 distinct very likely and possible pathogenic mutations, was largely different from the previously reported spectrum in patients from non-Asian populations. Screening for c.4957_4958insA and c.8868C>A mutations in the EYS gene may therefore be very effective for the genetic testing and counseling of RP patients in Japan.

Introduction

Retinitis pigmentosa (RP [MIM 268000]) is a highly heterogeneous genetic disease characterized by night blindness and visual field constriction leading to severe visual impairment. The disease appears with different modes of inheritance including autosomal recessive (ar), autosomal dominant (ad), and X-linked, and currently over half of cases are isolated in Japan.

To date, 53 causative genes and 7 loci of RP have been identified (http://www.sph.uth.tmc.edu/Retnet/), including the eyes shut homolog (EYS) gene encoding an ortholog of Drosophila spacemaker (spam), a protein essential for photoreceptor morphology. EYS spans over 2 Mb, making it one of the largest known genes expressed in the human eye [1,2]. EYS gene mutations, primarily truncating and some missense mutations, have been detected in arRP families of different ancestral origin and have reported to account for 5–16% of arRP [3–6]. Most gene mutations (e.g., RH0, PRPH2, PRPF31, RP1, and IMPDH1) have been found in Japanese patients with adRP, with few reports describing mutations in arRP [7,8]. Therefore, the genes causing arRP in most Japanese families have yet to be identified.

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* E-mail: hosono@hama-med.ac.jp
In this study, we screened all \emph{EYS} gene exons in 100 unrelated Japanese \emph{RP} patients. We found 2 novel truncating \emph{EYS} gene mutations that were surprisingly related to 16\% of Japanese \emph{arRP} patients, but were not detected in Japanese patients with either \emph{adRP} or Leber’s congenital amaurosis (LCA [MIM:204000]), the earliest onset and most severe form of hereditary retinal dystrophy with several clinical features overlapping with those of \emph{RP}. Additionally, these mutations were also detected in 9\% of Korean \emph{arRP} patients.

**Methods**

**Patients and clinical evaluation**

We screened all \emph{EYS} gene exons in 100 unrelated Japanese \emph{RP} patients with no systemic manifestations, excluding families with obvious autosomal dominant inheritance. Some pedigrees showed a pattern compatible with the recessive mode of inheritance; the other patients were considered isolated cases. In addition, 200 unrelated and non-RP Japanese individuals were screened as controls to evaluate the frequency of the mutations found in the patient samples. We also screened a part of \emph{EYS} gene exons 26 and 44 in 19 unrelated Japanese \emph{adRP} patients, 28 unrelated Japanese LCA patients, and 32 unrelated Korean \emph{arRP} patients. The 19 Japanese \emph{adRP} patients had already been screened for some principal \emph{adRP}-causing genes, but the pathogenic mutations have not yet been detected. Examples of the screening list for \emph{adRP}-causing genes and targeted exons include exon 3 and 4 in \emph{RP1}; exon 1, 2, 3, 4, and 5 in \emph{RHO}; exon 1, 2, and 3 in \emph{PRPH2}; exon 2, 3, and 4 in \emph{CRX}; exon 11 in \emph{PRP3}; exon 10, 11, and 12 in \emph{IMPDH1}; exon 2 in \emph{NRL}; exon 43 in \emph{PRPF8}; exon 1 and 2 in \emph{ROM1}; exon 5 and 6 in \emph{RP9}; exon 2, 3, 5, 6, 7, 8, 11, and 12 in \emph{PRPF31}; exon 11 and 15 in \emph{SEMA4A}; exon 1 in \emph{C4f}; exon 3 in \emph{GUCA1B}; exon 3 in \emph{SP4}; and exon 3 in \emph{TOPORS}.

Japanese \emph{RP} patients were examined either at the Department of Ophthalmology, Hamamatsu University Hospital in Hamamatsu (by YH), Department of Ophthalmology, Kobe City Medical Center General Hospital in Kobe (by MT), or Department of Ophthalmology, Nagoya University Hospital in Nagoya (by MK). Patients’ origin varied widely, from the Tokyo to Osaka areas in Japan. Japanese LCA patients were examined at the Department of Ophthalmology and Laboratory of Cell Biology, National Center for Child Health and Development in Tokyo (by NA). LCA patients’ origin varied widely, from all over Japan except the Okinawa islands. Meanwhile, Korean \emph{RP} patients were examined at the Department of Ophthalmology, Kyungpook National University Hospital in Daegu (by ITK). The Korean patients’ origin varied widely, from Daegu to Jeongju and Pohang areas in Gyeongsangbuk-do, Korea. A full ophthalmic examination was performed. Clinical diagnosis for \emph{RP} was based on visual field, fundus examination, and electroretinogram findings, and clinical diagnosis for \emph{LCA} was based on fundus examination and electroretinogram findings.

**Ethics statements**

This study was approved by the Institutional Review Board for Human Genetic and Genome Research at the 6 participating institutions (Hamamatsu University School of Medicine, RIKEN Center for Developmental Biology, Nagoya University Graduate School of Medicine, National Center for Child Health and Development, Chiba University Graduate School of Medicine, and Kyungpook National University Hospital), and its procedures conformed to the tenets of the Declaration of Helsinki. Written informed consent was obtained from all participants before molecular genetic studies.

**Mutation analysis**

Genomic DNA in Japanese samples was extracted from the peripheral lymphocytes using standard procedures. In Korean samples, whole blood samples were collected on FTA cards (GE Healthcare). Blood samples were spotted onto the cards and air-dried for 1 h at room temperature. For polymerase chain reaction (PCR) amplification, a 1:2-mm disk was punched from a dried blood spot using a Harris micro-punch tool (GE Healthcare) and processed according to the manufacturer’s instructions. PCR was performed using the KOD -Plus- ver, 2 PCR kit (Toyobo) with the primer sets described in Table S1 for 35 cycles of 98\C for 30 s, 60\C for 30 s, and 68\C for 1 min in an automated thermal cycler (GeneAmp PCR System 9700; Applied Biosystems). PCR products were purified with Wizard SV Gel and PCR Clean-up System (Promega) or treated with Exonuclease I and Antarctic Phosphatase (New England Biolabs). Direct sequencing was performed using the BigDye Terminator v3.1 Cycle Sequencing Kit on an ABI3100 autosequencer (Applied Biosystems). For Japanese \emph{arRP} patients, all 44 exons, including 3 non-coding exons (exons 1–3) that cover the 5’ untranslated region and 41 coding exons (exons 4–44), were analyzed in both sense and antisense directions. For Japanese \emph{adRP} and LCA patients, and Korean \emph{arRP} patients, parts of exons 26 and 44 were analyzed (Table S1).

**Assessment of pathogenicity**

A sequence variant was considered pathogenic if it represented a truncating mutation (nonsense or frameshift), large-scale deletion mutation, or missense mutation affecting a conserved amino acid residue and did not appear in control samples (number of alleles studied ≤400) and/or in a public SNP database (http://www.ncbi.nlm.nih.gov/projects/SNP/). Particularly, missense mutations were considered pathogenic if found together with a second variant, especially if it was truncating. As reference data, we employed 4 computational algorithms to evaluate the pathogenicity of missense mutations: SIFT (http://sift.jcvi.org/www/SIFT_seq_submit2.html), PolyPhen2 (http://genetics.bwh.harvard.edu/pph2/), PMut (http://mbc.pcb.ub.es/PMut/), and SNAP (http://roslab.org/services/snap/).

**Results**

**Mutation analysis**

Mutation analysis of \emph{EYS} in 100 unrelated Japanese patients revealed 7 very likely pathogenic mutations in 18 patients (18\%). Of these 18 patients, a second mutant allele could not be detected in 9 patients. The very likely pathogenic mutations consisted of 3 truncating mutations, 1 deletion mutation, 2 missense mutations, and 1 previously described mutation (Fig. 1, Table 1, and Table 2). In addition, we also identified 6 possible pathogenic mutations in 8 separate patients (Table 1 and Table 2).

A novel truncating insertion, c.4957_4958insA, was detected in 12 patients and accounted for 15 of the 35 mutated alleles detected (42.9\%) (Table 1 and Table 2). Three patients were homozygous for the c.4957_4958insA mutation, and the other 9 patients were heterozygous. Of the latter, 3 patients showed the second mutation while 6 did not. This insertion creates a frameshift mutation that predicts a premature stop at codon 1654 (p.S1653KfsX2). A novel truncating nonsense mutation c.8868C>G (p.Y2956X) was identified in 4 patients and accounted for 5 of the 35 mutated alleles detected (14.3\%). Thus, these 2 novel truncating mutations were identified in 16 separate patients, resulting in a very high frequency of the 2 mutations in Japanese \emph{arRP} patients.
Families with very likely pathogenic mutations and both alleles affected

Nine of the 18 patients bearing very likely pathogenic mutations appeared to have both alleles affected, suggesting that they received one mutated allele from each unaffected parent (Table 1 and Table 2). In 4 patients (RP3H, RP48K, RP56K, and RP81K), segregation analysis was performed, and the 2 pathogenic alleles were considered to be on different chromosomes (Fig. 2).

1. In RP3H, proband (II-6) was homozygous for c.4957_4958insA. The mutation co-segregated with the phenotype: the unaffected brother (II-4) demonstrated wild-type alleles, while the affected brother (II-5) was homozygous for the mutation.
2. In RP48K, proband (II-1) was homozygous for c.4957_4958insA. The unaffected brother (II-2) was heterozygous for the mutation.
3. In RP56K, proband (II-1) was compound heterozygous for c.4957_4958insA and missense mutation c.8351T>G (p.L2784R). The mutation co-segregated with the phenotype: the affected brother (II-2) also showed both mutations, while the unaffected brother (II-3) was heterozygous for c.4957_4958insA.
4. In RP81K, proband (II-5) was compound heterozygous for truncating insertion c.2522_2523insA (p.Y841X) and missense mutation c.6557G>G (p.G2186E). This insertion results in premature termination of the encoded protein at codon 841 (p.Y841X). Missense mutation c.6557G>G has been previously reported as disease causing in one Korean/American and one Chinese patient [3,6]. The unaffected mother (I-2) was heterozygous for c.2522_2523insA, while the unaffected sister (II-6) was heterozygous for c.6557G>G.

For the other patients, segregation analysis could not be performed due to difficulties in collecting samples from the families of patients (Table 1). RP54K and RP53K were homozygous for truncating mutation c.4957_4958insA and c.8868C>A, respectively. RP21H was homozygous for deletion in exon 32, an in-frame deletion that results in the replacement of amino acids from D2142 to S2191 with G2142 (p.D2142_S2191delinsG) and disrupts the second laminin G domain (Fig. 3). RP44K and RP87N were heterozygous for truncating and missense mutations, c.4957_4958insA/c.6657G>G (p.G2186E) and

| Family ID | Nucleotide change | Predicted effect | Domain* | Location in gene | Type of change | Reference |
|-----------|-------------------|------------------|---------|------------------|----------------|-----------|
| RP3Hb     | c.4957_4958insA/  | p.S1653KfsX2     | Close to coiled-coil | Exon 26/Exon 26 | Homozygous     | This study |
|           | c.4957_4958insA   |                  | Close to coiled-coil |                      |                |           |
| RP48Kb    | c.4957_4958insA/  | p.S1653KfsX2     | Close to coiled-coil | Exon 26/Exon 26 | Homozygous     | This study |
|           | c.4957_4958insA   |                  | Close to coiled-coil |                      |                |           |
| RP54K     | c.4957_4958insA/  | p.S1653KfsX2     | Close to coiled-coil | Exon 26/Exon 26 | Homozygous     | This study |
|           | c.6557G>G         | p.G2186E         | Laminin G           |                      |                |           |
| RP44Kb    | c.4957_4958insA/  | p.S1653KfsX2     | Close to coiled-coil | Exon 26/Exon 26 | Homozygous     | This study |
|           | c.8868C>A         | p.Y2956X         | EGF                 | Exon 44/Exon 44    | Homozygous     | This study |
| RP81Kb    | c.2522_2523insA/  | p.Y841X/p.G2186E | Laminin G/Laminin G | Exon 32/Exon 32    | Homozygous     | This study |
|           | c.6557G>G         | p.G2186E         | Laminin G/Laminin G |                      |                |           |
| RP1H      | c.4957_4958insA   | p.S1653KfsX2     | Close to coiled-coil | Exon 26             | Heterozygous   | This study |
| RP6H      | c.4957_4958insA   | p.S1653KfsX2     | Close to coiled-coil | Exon 26             | Heterozygous   | This study |
| RP12H     | c.4957_4958insA   | p.S1653KfsX2     | Close to coiled-coil | Exon 26             | Heterozygous   | This study |
| RP51K     | c.4957_4958insA   | p.S1653KfsX2     | Close to coiled-coil | Exon 26             | Heterozygous   | This study |
| RP69H     | c.4957_4958insA   | p.S1653KfsX2     | Close to coiled-coil | Exon 26             | Heterozygous   | This study |
| RP100K    | c.4957_4958insA   | p.S1653KfsX2     | Close to coiled-coil | Exon 26             | Heterozygous   | This study |
| RP8H      | c.8868C>A         | p.Y2956X         | EGF                 | Exon 44             | Heterozygous   | This study |
| RP25H     | c.8868C>A         | p.Y2956X         | EGF                 | Exon 44             | Heterozygous   | This study |
| RP80Kb    | c.8868C>A         | p.Y2956X         | EGF                 | Exon 44             | Heterozygous   | This study |
| RP4H      | c.9272T>C         | p.I3091T         | Laminin G/Laminin G | Exon 32/Exon 32    | Homozygous     | This study |
| RP9H      | c.8875C>A         | p.L2959M         | EGF                 | Exon 44             | Heterozygous   | This study |
| RP49K     | c.9272T>C         | p.I3091T         | Laminin G/Laminin G | Exon 32/Exon 32    | Homozygous     | This study |
| RP53K     | c.5884A>G         | p.T1962A         | Laminin G/Laminin G | Exon 28             | Heterozygous   | This study |
| RP55K     | c.9272T>C         | p.I3091T         | Laminin G/Laminin G | Exon 32/Exon 32    | Homozygous     | This study |
| RP74K     | c.5404C>T         | p.L1802F         | Close to Laminin G  | Exon 26             | Heterozygous   | This study |
| RP79K     | c.77G>A           | p.R26Q           | Close to signal peptide cleavage site | Exon 4 | Heterozygous | This study |
| RP83K     | c.2923T>C         | p.C975R          | EGF                 | Exon 19             | Heterozygous   | This study |

Nucleotide numbering reflects cDNA numbering with +1 corresponding to the A of the ATG translation initiation codon in the reference sequence FM209056, according to the nomenclature recommended by the Human Genome Variation Society (www.hgvs.org/mutnomen). The initiation codon is codon 1. None of these 13 mutations were found in the Japanese controls.

*EYS has a signal peptide, a putative coiled-coil, 29 EGF, and 5 Laminin G domains. See Fig. 3.

**Segregation analysis has been performed. See Fig. 2.

In RP56K and RP81K, 2 pathogenic alleles were considered to be on different chromosomes (compound heterozygous). See Fig. 2.
### Table 2. Summary of the very likely and possible pathogenic mutations identified in 100 Japanese arRP patients.

| Mutation Type | Allele Frequency | Location in Gene | Domain* | Conservation in hu/o/m/ho/d/op/p/c/z/dr | Predicted Effect | Control | Patient | Family ID | Reference | Species | SIFT | PolyPhen2 (HumDiv) | PMut | SNAP |
|---------------|------------------|------------------|---------|---------------------------------------|------------------|---------|---------|----------|-----------|---------|------|------------------|------|------|
| **Very likely** |                  |                  |         |                                       |                  |         |         |          |           |         |      |                  |      |      |
| insertion     |                  |                  |         |                                       |                  |         |         |          |           |         |      |                  |      |      |
| c.2522_2523insA | c.4957_4958insA  |                  |         |                                       |                  |         |         |          |           |         |      |                  |      |      |
| c.2923T>C     |                  |                  |         |                                       |                  |         |         |          |           |         |      |                  |      |      |
| c.3904C>T     |                  |                  |         |                                       |                  |         |         |          |           |         |      |                  |      |      |
| c.8875C>A     |                  |                  |         |                                       |                  |         |         |          |           |         |      |                  |      |      |
| c.9272T>C     |                  |                  |         |                                       |                  |         |         |          |           |         |      |                  |      |      |
| **Nonsense**  |                  |                  |         |                                       |                  |         |         |          |           |         |      |                  |      |      |
| c.8646C>A     |                  |                  |         |                                       |                  |         |         |          |           |         |      |                  |      |      |
| c.7793G>A     |                  |                  |         |                                       |                  |         |         |          |           |         |      |                  |      |      |
| c.8351T>G     |                  |                  |         |                                       |                  |         |         |          |           |         |      |                  |      |      |
| c.9272T>C     |                  |                  |         |                                       |                  |         |         |          |           |         |      |                  |      |      |
| **Deletion**  |                  |                  |         |                                       |                  |         |         |          |           |         |      |                  |      |      |
| Deletion exon 32 | Deletion exon 32 |                  |         |                                       |                  |         |         |          |           |         |      |                  |      |      |
| **Missense**  |                  |                  |         |                                       |                  |         |         |          |           |         |      |                  |      |      |
| c.6557G>A     |                  |                  |         |                                       |                  |         |         |          |           |         |      |                  |      |      |
| c.7793G>A     |                  |                  |         |                                       |                  |         |         |          |           |         |      |                  |      |      |
| c.8351T>G     |                  |                  |         |                                       |                  |         |         |          |           |         |      |                  |      |      |
| c.9272T>C     |                  |                  |         |                                       |                  |         |         |          |           |         |      |                  |      |      |
| **Possible**  |                  |                  |         |                                       |                  |         |         |          |           |         |      |                  |      |      |
| Missense      |                  |                  |         |                                       |                  |         |         |          |           |         |      |                  |      |      |
| c.77G>A       |                  |                  |         |                                       |                  |         |         |          |           |         |      |                  |      |      |
| c.2923T>C     |                  |                  |         |                                       |                  |         |         |          |           |         |      |                  |      |      |
| c.8875C>A     |                  |                  |         |                                       |                  |         |         |          |           |         |      |                  |      |      |
| c.9272T>C     |                  |                  |         |                                       |                  |         |         |          |           |         |      |                  |      |      |

*EYS contains a signal peptide, a putative coiled-coil, 2 EGF, and 5 laminin G domains. See Fig. 3.

*hu/o/m/ho/d/op/p/c/z/dr denotes Human/Orangutan/Marmoset/Horse/Dog/Opossum/Platypus/Chicken/Zebrafish/Drosophila EYS orthologs, respectively. The hyphen (-) indicates that genomic sequence of corresponding region in the species was reported to be unknown [5].

*SIFT, PolyPhen2 (only the HumDiv data are shown), PMut, and SNAP were used as reference data to evaluate the pathogenicity of the missense mutations. c.77G>A, c.2923T>C, c.7793G>A, c.8351T>G, and c.9272T>C were predicted to be pathogenic by a number of different computational prediction programs. In addition, the c.6557G>A mutation, which had been previously reported as disease causing, was classified as pathogenic by the PolyPhen2, PMut, and SNAP programs.

*Homozygous exon 32 deletion mutation was not detected in 200 controls.

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c.4957_4958insA/c.7793T G (p.G2598D), respectively. None of these very likely pathogenic mutations were found in the Japanese controls.

Families with single novel very likely pathogenic mutations

The rest of the patients comprising the group with very likely pathogenic mutations presented only single truncating mutations (Table 1 and Table 2). RP1H, RP6H, RP12H, RP51H, RP96H, and RP100N were heterozygous for c.4957_4958insA. RP8H, RP25H, and RP80K were heterozygous for c.8868C>A. Segregation analysis was performed in patient RP80K. The unaffected father (I-1) demonstrated wild-type alleles, and the unaffected mother (I-2) was heterozygous for the mutation (Fig. 2).

In RP96H, we found very likely pathogenic missense mutation c.8923T>C (p.F2975L), which was not detected in any of the 400 control alleles. However, as c.8923T>C has been described as rs79036642 in the dbSNP database, it was assigned to the group of possible non-pathogenic sequence alterations (Table 3).

Screening of the 2 truncating mutations

We focused on 2 truncating mutations, c.4957_4958insA in exon 26 and c.8868C>A in exon 44, which were identified in 16 separate Japanese arRP patients in this study. The frequency of the 2 mutations was very high in this Japanese arRP cohort. However, we did not detect the 2 mutations in 19 Japanese adRP

![Figure 2. Pedigrees of the families that was available for mutation analysis.](image-url)
patients and 28 LCA patients who were recruited and screened to evaluate the frequency of the mutations. We also recruited 32 unrelated Korean arRP patients and screened for the EYS gene mutations. The c.4957_4958insA mutation was detected in 2 patients and accounted for 3 of 64 Korean patient alleles (4.7%). One patient was homozygous and the other was heterozygous. The c.8868C>A mutation was identified in 1 patient and accounted for 1 of the 64 Korean patient alleles (1.6%).

Clinical findings

Nine Japanese patients with very likely pathogenic EYS gene mutations in both alleles, 9 Japanese patients with single very likely pathogenic changes, and a Korean patient with homozygous c.4957_4958insA mutation demonstrated classic RP with mostly night blindness as the initial symptom, followed by gradual constriction of the visual field. The fundus displayed bone spicules increasing in density with age and attenuated retinal vessels. Electroretinogram responses were not detectable, consistent with severe generalized rod-cone dysfunction. The remaining visual field determined using Goldmann kinetic perimetry with V-4 target ranged from approximately 10° to 60° of the central and inferior visual fields, respectively, in a 74-year-old woman (RP100N) to complete blindness in a 54-year-old man (RP21H).

No remarkable clinical difference was observed between 9 patients with very likely pathogenic EYS gene mutations in both alleles and 9 patients with single very likely pathogenic changes.

Discussion

This study is the first to analyze mutations in the EYS gene among Japanese arRP patients. We detected 67 sequence alterations in the EYS gene, of which 21 were novel. Of these, 7 were very likely pathogenic, 6 were possibly pathogenic, and 54 were possible non-pathogenic sequence alterations (Table 1, Table 2, and Table 3).

Considering only the very likely pathogenic mutations, the minimum observed prevalence of distinct EYS gene mutations in our study is 18% (18/100, 9 patients with 2 very likely pathogenic mutations and 9 with only one such mutations). Additionally, if the possible pathogenic mutations are included in the prevalence estimation, prevalence increases to 26% (26/100, with 17 of 26 patients presenting single mutations). The estimated prevalence in our study may be extremely high compared with those in the previous studies [3–6]. Until recently, mutations in 34 genes have been associated with arRP (http://www.sph.uth.tmc.edu/Retnet/). The most frequently mutated gene is USH2A, accounting for

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**Figure 3. Predicted domain structure and distribution of identified EYS mutations.** SMART (http://smart.embl-heidelberg.de/) and Pfam (http://pfam.sanger.ac.uk/) were used to search protein functional domains. A coiled-coil domain identified by Barragan et al. (2010) between the EGF-like domain and laminin G domain was also indicated. Novel very likely pathogenic mutations, novel possible pathogenic mutations, and a previously described mutation are shown in bold, normal, and italic type, respectively. Six out of 9 missense mutations were found in the EGF or laminin G domains. Furthermore, 7 were located in the latter half of the protein between the putative coiled-coil region and C-terminus.

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**Table 3.** Summary of the possible non-pathogenic sequence alterations in the EYS gene identified in this study.

| Gene exon | Nucleotide change | Predicted effect | Conservation in hu/om/h/o/d/o/p/c/z/dr* | Patient frequency | Control frequency | SNP ID | Reference |
|-----------|-------------------|------------------|----------------------------------------|------------------|------------------|--------|-----------|
| Exon 1  | c.500A>G          |                  |                                        | 13/200           |                  |        | Abd El-Aziz et al., 2010 |
| Exon 4  | c.334G>A          | p.V112I          | V/I/I/I/V/V/-/-/-/E                     | 1/200            | 0/192            | rs112609906 |                      |
| Exon 19 | c.2980C           |                  |                                        | 1/200            | 0/192            | rs12193967 | Audo et al., 2010; Abd El-Aziz et al., 2010 |
| Exon 22 | c.3568            |                  |                                        | 12/200           |                  |        | Abd El-Aziz et al., 2010 |
| Exon 23 | c.3568_3570delTGA |                  |                                        | 12/200           |                  |        | Abd El-Aziz et al., 2010 |
| Exon 26 | c.3936A           | p.K120M          | T/T/T/T/A/-/-/-/-                      | 137/200          |                  |        | Abd El-Aziz et al., 2010 |
| Exon 27 | c.4026C           | p.L1342S         | S/S/S/S/-/-/-/-                       | 10/200           |                  |        | Abd El-Aziz et al., 2010 |
| Exon 28 | c.4081A>G         | p.I1361V         | I/I/T/V/-/-/-/-                     | 12/200           |                  |        | Abd El-Aziz et al., 2010 |
| Exon 29 | c.4256T>C         | p.L1419S         | L/S/S/S/L/S/V/Q/V                    | 137/200          |                  |        | Abd El-Aziz et al., 2010 |
| Exon 30 | c.4325T>C         | p.L1419S         | L/S/S/S/L/S/V/Q/V                    | 137/200          |                  |        | Abd El-Aziz et al., 2010 |
| Exon 31 | c.4543C>T         | p.R1515W         | R/R/R/R/R/-/-/-/-                     | 36/200           |                  |        | Abd El-Aziz et al., 2010 |
| Exon 32 | c.4593G>A         | p.S1517G         | S/G/D/T/T/-/-/-/-                     | 36/200           |                  |        | Abd El-Aziz et al., 2010 |
| Exon 33 | c.5244A>C         | p.L1748F         | L/L/L/L/L/-/-/-/-                     | 8/200            |                  |        | Abd El-Aziz et al., 2010 |
| Exon 34 | c.6834T>G         | p.L1873V         | L/L/L/P/-/-/-/-/-                    | 38/200           |                  |        | Abd El-Aziz et al., 2010 |

*Note: The table provides a summary of non-pathogenic sequence alterations identified in the EYS gene, including nucleotide changes, predicted effects, conservation scores, and patient/control frequencies. The table also includes references for the identified alterations.*
approximately 7% of arRP cases [9,10], whereas most other genes contribute to only 1% to 2% of arRP cases [11]. The estimated prevalence of very likely and possible pathogenic mutations of the EYS gene in our study was 26%, suggesting its major involvement in the pathogenesis of arRP in the Japanese population.

We found that 16% of Japanese arRP patients displayed at least one c.4957_4958insA or c.8868C>A mutation, which accounted for 57.1% (15/53) of the mutated alleles. Thus, these mutations seem to be frequent among Japanese arRP patients. Previous studies employing Indonesian, Pakistani, Chinese, Israeli, Spanish, French, Britisch, Dutch, and Palestinian RP patient populations have not detected them [3-6,12-15]. Since the Japanese were divided into small semi-closed population groups among which intercommunication was quite less until the mid-20th century, obvious or latent consanguineous marriages were carried out more frequently, leading to relatively high inbreeding levels in those populations. The frequency of the c.4957_4958insA and c.8868C>A mutations may result from a founder effect like that of the 2299delG USH2A gene mutation, which accounts for 44% of disease alleles in Danish and Norwegian patients with Usher syndrome type II [16].

We detected 13 different very likely and possible pathogenic mutations. Three were truncating mutations and accounted for 60% (21/35) of mutated alleles. Likewise, previous studies reported that most pathogenic mutations were truncated type (nonsense, deletion, insertion, or splicing) [3-6,12-15]. Furthermore, c.6557G>A was the only mutation that was common between the Japanese and other populations. This mutation has been found in Korean/American and Chinese patients [3,6]. These results indicate that the EYS gene mutation spectrum among Japanese patients largely differs from that among the previously mentioned non-Asian populations. The Japanese and Korean mutation spectrum may resemble each other, but an accurate comparison could not be made, because further EYS gene analysis of Korean RP patients is required to clarify this possibility.

A second mutant allele could not be detected by direct sequencing in 17 of 26 patients in our study. Previous studies reported 7 of 10 [3] and 9 of 17 [5] patients with heterozygous EYS gene mutation, implying that this finding could be due to relatively large heterozygous deletions [15]. The second mutation in these families may also have been located within the gene regulatory elements or unknown exons including alternative splicing areas.

Although rare, a single EYS mutation in combination with another mutation on a second gene could also explain this phenotype [3]. The c.4957_4958insA and c.8868C>A mutations were not detected in Japanese patients with adRP or with LCA. Abd El-Aziz et al. reported that EYS gene mutation screening did not reveal any pathogenic mutations in 95 British and Chinese adRP patients [3]. Bandah-Rosenfeld et al. reported that no mutation was found in 2 Oriental Jewish and Israeli Muslim LCA patients who had a large homozygous region harboring the EYS gene [12]. Although further analysis of all EYS gene exons is required, EYS gene mutations may not be detected in Japanese patients with adRP and LCA. The c.4957_4958insA and c.8868C>A mutations were also detected in Korean patients with arRP and accounted for 6.3% (4/64 alleles) of the disease alleles. Similar to Japanese arRP results, the c.4957_4958insA mutation was more frequently detected than the c.8868C>A mutation. The fact that both c.4957_4958insA and c.8868C>A mutations were also detected in Korean patients suggests the possibility that the mutations occurred in an ancient common ancestor and spread throughout East Asia.

RP is a highly heterogeneous disease, with a reported prevalence rate of 1 in 4,000-8,000 people in Japan. Given the population of Japan, approximately a 100 million, the number of patients with RP can be estimated to be 12,500–25,000. The relative frequencies of RP inheritance patterns in Japanese patients were estimated as 25.2% for autosomal recessive, 16.9% for autosomal dominant, 1.6% for X-linked, and 56.3% for simplex, indicating that most Japanese RP patients represent adRP or isolated cases [17]. Autosomal recessive and simplex cases account over 80% of RP cases in Japan (approximately 10,000–20,000 people). Our results indicate that c.4957_4958insA and c.8868C>A mutations are possibly present in 1,600–3,200 Japanese patients with RP. These 2 novel mutations will be very useful for genetic diagnosis and counseling, and analysis of the mutated proteins may be helpful in the development of effective therapies for RP in Japan and Korea.

In conclusion, mutation screening of the EYS gene in 100 Japanese patients revealed 13 different pathogenic mutations, confirming that the mutation spectrum in Japanese patients differs from the previously reported spectrum in patients of non-Asian populations. Among these 13 mutations, 2 truncating mutations, c.4957_4958insA and c.8868C>A, were detected in at least one mutated allele in 16% of Japanese arRP patients and may be the
most frequent mutations causing RP in the Japanese populations. Screening for c.4957_4958insA and c.8868C>A mutations in the EYS gene is, therefore, very effective for the genetic testing and counseling of RP patients in Japan. Further analysis is necessary to obtain a more precise mutation spectrum and to identify other frequent mutations in other East Asian populations.

Supporting Information

Table S1  PCR primer sequences for human EYS.  

(DOC)

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Author Contributions

Conceived and designed the experiments: KH MT SY MK YH. Performed the experiments: KH CI YZ. Analyzed the data: KH CI. Contributed reagents/materials/analysis tools: MT DHP YH HS SU TY AH TF SN JPS ITK SY NA HT MS MK YH. Wrote the paper: KH SM YH.