

GNAT1 Associated with Autosomal Recessive Congenital Stationary Night Blindness

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PURPOSE. Congenital stationary night blindness is a nonprogressive retinal disorder manifesting as impaired night vision and is generally associated with other ocular symptoms, such as nystagmus, myopia, and strabismus. This study was conducted to further investigate the genetic basis of CSNB in a consanguineous Pakistani family.

METHODS. A consanguineous family with multiple individuals manifesting cardinal symptoms of congenital stationary night blindness was ascertained. All family members underwent detailed ophthalmic examination, including fundus photographic examination and electroretinography. Blood samples were collected and genomic DNA was extracted. Exclusion and genome-wide linkage analyses were completed and two-point LOD scores were calculated. Bidirectional sequencing of *GNAT1* was completed, and quantitative expression of *Gnat1* transcript levels were investigated in ocular tissues at different postnatal intervals.

RESULTS. The results of ophthalmic examinations were suggestive of early-onset stationary night blindness with no extraocular anomalies. The genome-wide scan localized the critical interval to chromosome 3, region p22.1-p14.3, with maximum two-point LOD scores of 3.09 at $\theta = 0$, flanked by markers *D3S3522* and *D3S1289*. Subsequently, a missense mutation in *GNAT1*, p.D129G, was identified, which segregated within the family, consistent with an autosomal recessive mode of inher-

itance, and was not present in 192 ethnically matched control chromosomes. Expression analysis suggested that *Gnat1* is expressed at approximately postnatal day (P)7 and is predominantly expressed in the retina.

CONCLUSIONS. These data suggest that a homozygous missense mutation in *GNAT1* is associated with autosomal recessive stationary night blindness. (*Invest Ophthalmol Vis Sci.* 2012; 53:1353–1361) DOI:10.1167/iovs.11-8026

Congenital stationary night blindness (CSNB) is a clinically heterogeneous group of nonprogressive retinal disorders, characterized by impaired night vision, decreased visual acuity, nystagmus, myopia, and strabismus.¹ CSNB can be classified into two groups based on the electroretinographic recordings that exhibit waveforms in response to flashes of light that correspond to changes in the polarization of the photoreceptor and bipolar cells.^{2,3} The Schubert-Bornschein type is characterized by an electronegative electroretinogram (ERG) at the scotopic bright flash, in which the amplitude of the b-wave is smaller than that of the a-wave,² while the Riggs type is defined by proportionally reduced a- and b-waves.³ The Schubert-Bornschein- and Riggs-type CSNB patients not only differ electrophysiologically, but manifest different clinical characteristics. Decreased visual acuity, myopia, and nystagmus can be associated with Schubert-Bornschein CSNB, whereas patients with Riggs-type CSNB have visual acuity within normal range with no symptoms of myopia and/or nystagmus.¹

Familial cases of CSNB with autosomal dominant, autosomal recessive as well as X-linked inheritance have been reported. Mutations in *RHO*, *PDE6B*, and *GNAT1* have been associated with autosomal dominant CSNB, and patients with mutations in these genes exhibit the Riggs- or Schubert-Bornschein-type ERG.^{4–10} The Schubert-Bornschein form can be further subdivided into complete and incomplete types. Complete CSNB is characterized by reduced rod b-wave response, whereas the incomplete CSNB is characterized by both a reduced rod b-wave and substantially reduced flicker cone responses, due to both ON and OFF bipolar cell dysfunction.¹¹ Mutations in *GRM6* and *TRPM1* lead to autosomal recessive CSNB, whereas mutations in *NYX* to X-linked complete CSNB.^{12–18} Mutations in *CABP4* have been associated with autosomal recessive CSNB and mutations in *CACNA1F* with X-linked incomplete CSNB.^{19–21} Recently, *SLC24A1* was implicated in the pathogenesis of autosomal recessive CSNB with Riggs-type ERG response.²²

Transducin is a trimeric G protein that is expressed in the retina and consists of three subunits.²³ The α -subunit binds GTP and activates cyclic GMP phosphodiesterase (PDE), whereas the β and γ subunits form a complex that is necessary to interact with rhodopsin.²⁴ The α -subunit expressed in the rod cells is encoded by *GNAT1*, whereas α -subunit expressed

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in the cone cells is encoded by *GNAT2*.^{25,26} Transducin is activated by conformational changes in rhodopsin due to the absorption of light, which causes a GDP bound α subunit to be exchanged with GTP and subsequent dissociation from the β -gamma complex.^{27,28}

Previously, a missense mutation c.113G>A was identified in the Nougaret form of autosomal dominant CSNB.⁹ This mutation (Gly38Arg) affects part of small loop of α subunit of rod transducin and is believed to affect GTPase activity.⁹ Recently, Szabo et al.¹⁰ identified a heterozygous mutation; c.598C>G, in exon 6 of *GNAT1* in individuals affected with autosomal dominant CSNB. This mutation (Gln200Glu) resides in a domain that plays an important role in binding and hydrolyzing GTP, and the ERG recordings were suggestive of a nonprogressive presynaptic defect in rod phototransduction.¹⁰

Here, we report a consanguineous family with multiple individuals diagnosed with CSNB that segregated within the family in an autosomal recessive fashion. A genome-wide link-

age scan localized the disease phenotype to chromosome 3, region p22.1-p14.3, harboring *GNAT1*, a gene previously associated with autosomal dominant CSNB. Bidirectional sequencing identified a homozygous missense mutation in *GNAT1* that segregated with the disease phenotype in the family and was not present in ethnically matched control chromosomes. To the best of our knowledge, this is the first report to implicate *GNAT1* in the pathogenesis of autosomal recessive CSNB.

MATERIALS AND METHODS

Clinical Assessment

A consanguineous Pakistani family with four affected individuals with a history of night blindness was recruited to participate in study to investigate autosomal recessive CSNB. Institutional review board (IRB) approval for this study was obtained from the National Centre of Excellence in Molecular Biology and the National Eye Institute. The

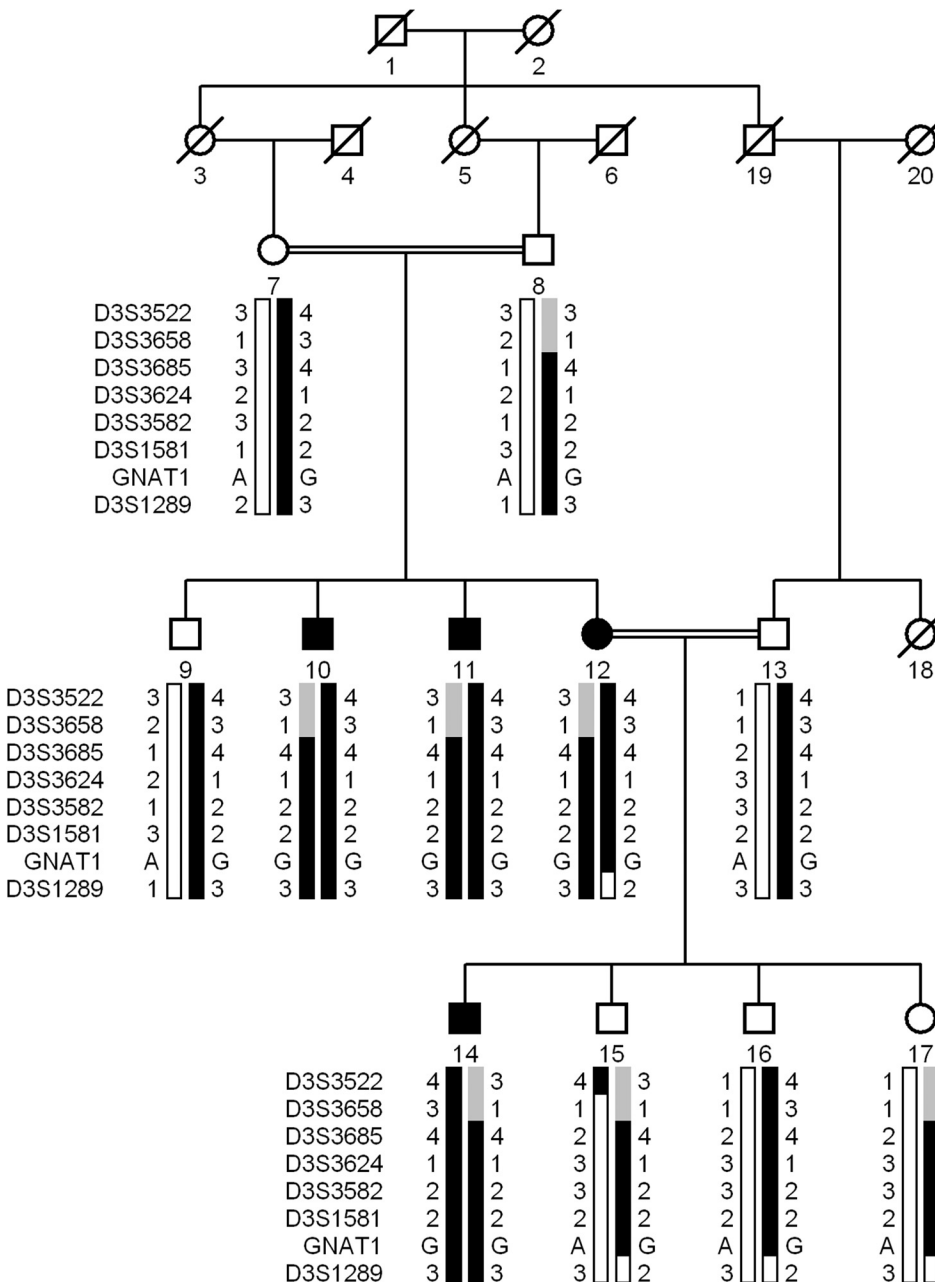


FIGURE 1. Pedigree of PKRP130 with a haplotype formed from alleles of the short arm of chromosome 3 microsatellite markers. Alleles forming the risk haplotype (black), alleles cosegregating with CSNB but not showing homozygosity (gray), and alleles not cosegregating with CSNB (white) are shown. Squares: males; circles: females; filled symbols: affected individuals; double line between individuals: consanguinity; and diagonal line through a symbol: deceased member.

participating subjects gave informed written consent consistent with the tenets of the Declaration of Helsinki. Funduscopy was performed at Layton Rehmatullah Benevolent Trust (LRBT) Hospital (Lahore, Pakistan). Electroretinography (ERG) measurements were recorded by using equipment manufactured by LKC (Gaithersburg, MD). Rod responses were determined through incident flash attenuated by -25 dB, and the rod-cone response was measured at 0 dB. Isolated cone responses were recorded at 0 dB with a 30-Hz flicker to a background illumination of 17 to 34 cd/m^2 . Color vision was assessed through Ishihara charts. Blood samples were collected from affected and unaffected family members and genomic DNA was extracted by the non-organic method as described previously.²⁹

Genotype Analysis

A genome-wide scan was performed with 382 highly polymorphic fluorescent markers from a linkage mapping with (PRISM MD-10;

Applied Biosystems, Inc. [ABI], Foster City, CA) having an average spacing of 10 centimorgans (cM). Multiplex polymerase chain reactions were performed, as follows (9700 PCR System; ABI): Each reaction was performed in a 5- μL mixture containing 40 ng genomic DNA, various combinations of 10 μM dye-labeled primers pairs, 0.5 μL 10 \times PCR buffer (GeneAmp; ABI), 1 mM dNTP mix, 2.5 mM MgCl_2 , and 0.2 U of *Taq* DNA polymerase. Initial denaturation was performed for 5 minutes at 95°C, followed by 10 cycles of 15 seconds at 94°C, 15 seconds at 55°C, and 30 seconds at 72°C and then 20 cycles of 15 seconds at 89°C, 15 seconds at 55°C, and 30 seconds at 72°C. The final extension was performed for 10 minutes at 72°C, followed by a final hold at 4°C. PCR products from each DNA sample were pooled and mixed with size standards (HD-400; ABI). The resulting PCR products were separated in a DNA analyzer (model 3100; ABI), and alleles were assigned (GeneMapper Software ver. 4.0; ABI).

Linkage Analysis

Two-point linkage analyses were performed using the FASTLINK version of MLINK, and the maximum LOD score was calculated with ILINK from the LINKAGE program.^{30,31} Autosomal recessive CSNB was analyzed as a fully penetrant trait with an affected allele frequency of 0.001. The marker order and distances between the markers were obtained from the Marshfield database and the National Center for Biotechnology Information (NCBI) chromosome 3 sequence maps. For the initial genome scan, equal allele frequencies were assumed, whereas for fine mapping, allele frequencies were estimated from 96 unrelated and unaffected individuals from the Punjab province of Pakistan.

Mutation Screening

Primer pairs for individual exons were designed with Primer3 software. The sequences and annealing temperatures are available on request. Amplifications were performed in 25 μL reactions containing 50 ng of genomic DNA, 8 picomoles of each primer, 250 μM dNTP, 2.5 mM MgCl_2 , and 0.2U *Taq* DNA polymerase in the standard 1 \times PCR buffer provided by the manufacturer (ABI). PCR amplification consisted of a denaturation step at 96°C for 5 minutes, followed by 40 cycles, each consisting of 96°C for 30 seconds followed by 57°C (or specific annealing temperature of the primer pair) for 30 seconds and at 72°C for 1 minute. PCR products were analyzed on 2% agarose gel, precipitated, and purified by ethanol precipitation. The PCR primers for each exon were used for bidirectional sequencing by means of dye termination chemistry, according to the manufacturer's instructions (Big Dye Terminator Ready Reaction Mix; ABI). Sequencing products were resuspended in 10 μL of formamide (ABI) and denatured at 95°C for 5 minutes. Sequencing was performed on an automated system (PRISM 3100; ABI). The sequencing results were assembled with sequencing software and analyzed (SeqScape software, ver. 3.7; ABI).

Prediction Analysis

The degree of evolutionary conservation of positions at Asp129 and the possible impact of an amino acid substitution on the structure of GNAT1 protein was examined with the SIFT and PolyPhen tools, available online. Evolutionary conservation of the mutated amino acids in other *GNAT* orthologs was examined using the UCSC genome browser.

Comparative Homology Modeling

Comparative homology modeling was performed using crystal coordinates for the human *GNAT1*. The wild and mutated protein sequences were loaded on to the Swiss PDB Viewer, and homology models were created. Hydrogen bonding was incorporated in the Swiss PDB Viewer.

Quantitative Expression of *Gnat1* in the Eye

C57BL/6 mice were maintained in a 12 hour dark-light cycle according to the ARVO Statement for the Use of Animals in Ophthalmic and

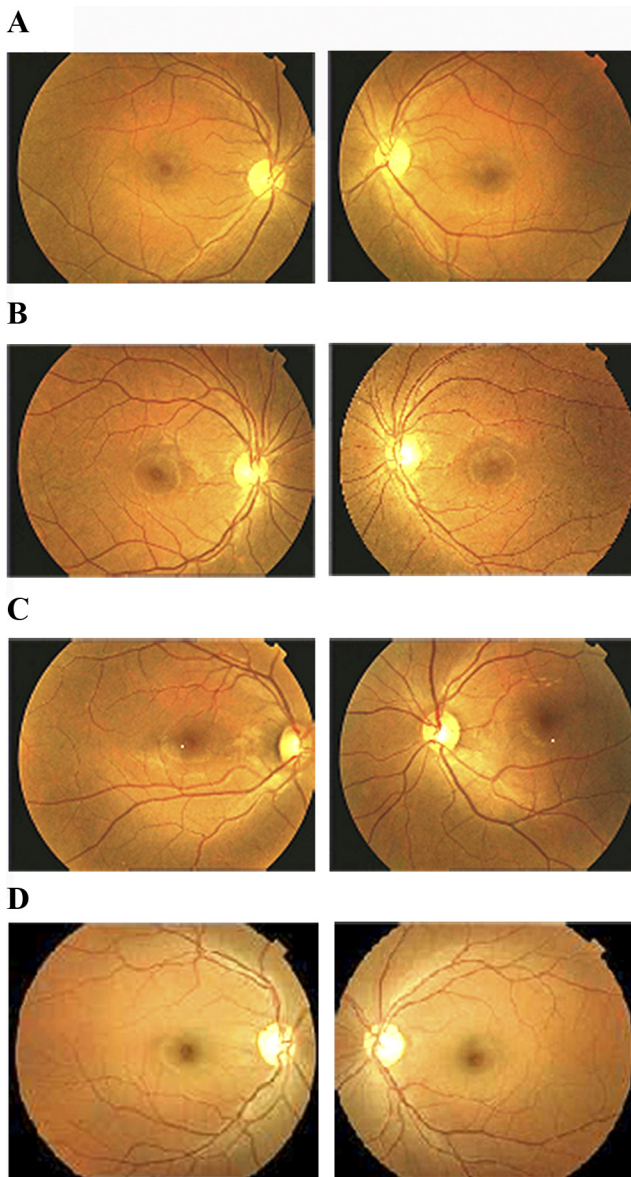


FIGURE 2. Fundus photographs of family PKRP130. (A) OD and OS of affected individual 10, (B) OD and OS of affected individual 12, (C) OD and OS of affected individual 14, and (D) OD and OS of unaffected individual 9. Fundus photographs of affected individuals 10, 12, and 14 revealed no signs of retinal arteries attenuation or bone spicule pigmentations in the retina.

Vision Research and with protocols approved by UCSD Institutional Animal Care and Use Committee. The eye tissues were collected from C57BL/6 mice that were euthanized at predetermined time points at the end of the dark cycle by CO₂ asphyxiation followed by cervical dislocation. Quantitative RT-PCR was performed, and data were analyzed according to previously published protocols.³² The expression of *Gnat1* was determined using GACGCTGTCACCGACATTATCATC and GGCTGCAGGGCTACAGAATCTC and analyzing *Rpl19* as a housekeeping gene to normalize *Gnat1* gene expression, as described.³⁵ Expression levels (\pm SEM) were calculated by analyzing at least three independent samples with replica reactions and presented on an arbitrary scale that represents the expression over the housekeeping gene. Rhodopsin mRNA levels served as a molecular marker for photoreceptor development and were measured in the same set of samples.

Internet Sources

The following web sites were used in the study: ILink from LINKAGE, Rockefeller University, New York, NY, <http://linkage.rockefeller.edu/soft/>; the FASTLINK version of MLINK and ILink, NCBI, National Institutes of Health, Bethesda MD, <http://www.ncbi.nlm.nih.gov/>; Marshfield Database, Marshfield Clinic, Marshfield, WI, <http://research.marshfieldclinic.org/>; Primer3 PCR Primer Design Tool; <http://primer3.sourceforge.net/>; PolyPhen: Prediction of Functional Effect of Human nsSNPs, Brigham and Women's Hospital, Harvard Medical School, Boston, MA, <http://genetics.bwh.harvard.edu/pph/>; SIFT database, J. Craig Venter Institute, Rockville, MD, <http://sift.jcvi.org/>; Human Genome Browser, University of California, Santa Cruz, Santa Cruz, CA, <http://genome.ucsc.edu/>; and Swiss PDB-Viewer, Swiss Institute of Bioinformatics, Geneva, Switzerland, <http://us.expasy.org/spdbv/>.

primer3.sourceforge.net; PolyPhen: Prediction of Functional Effect of Human nsSNPs, Brigham and Women's Hospital, Harvard Medical School, Boston, MA, <http://genetics.bwh.harvard.edu/pph/>; SIFT database, J. Craig Venter Institute, Rockville, MD, <http://sift.jcvi.org/>; Human Genome Browser, University of California, Santa Cruz, Santa Cruz, CA, <http://genome.ucsc.edu/>; and Swiss PDB-Viewer, Swiss Institute of Bioinformatics, Geneva, Switzerland, <http://us.expasy.org/spdbv/>.

RESULTS

The family PKRP130 was recruited from the Punjab province of Pakistan (Fig. 1). The medical records available to us suggested that all four affected individuals of the family had reported an inability to see at night from the early years of their lives. Fundus photographs of affected individuals 10, 12, and 14 reveal no signs of retinal artery attenuation or bone spicule pigmentation (Fig. 2). The ERGs of affected individuals, 10, 12, and 14 show reduced or nearly absent a- and b-waves, even after 4 hours of dark adaptation (Fig. 3). The 30-Hz flicker recordings representing isolated cone responses were normal, although the amplitude of these recordings were reduced for affected individuals 10 and 12, when compared with those in

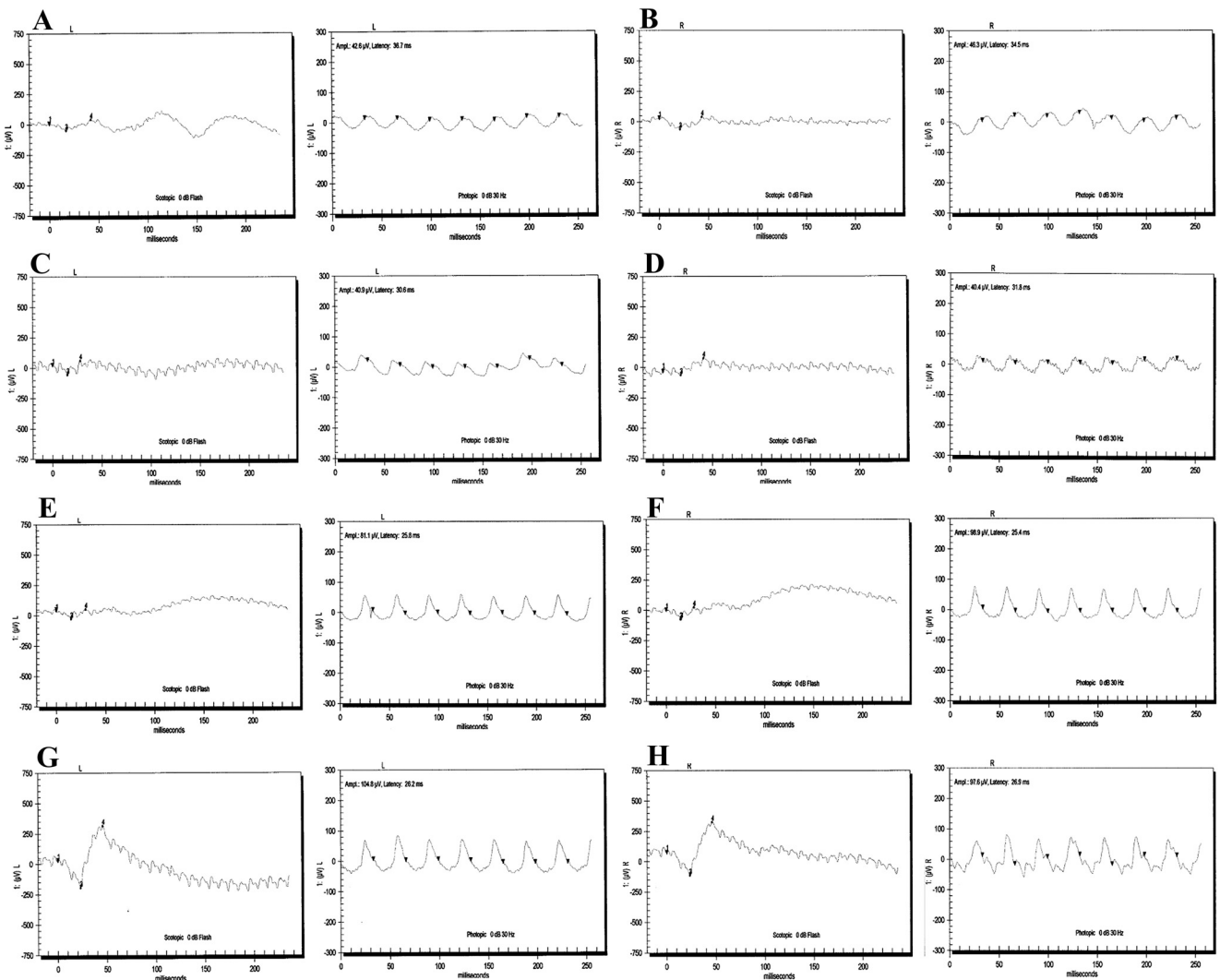


FIGURE 3. ERG recordings of family PKRP130. Combined rod and cone response recorded at 0 dB and cone responses recorded with 30-Hz flicker of affected individual 10: (A) OS and (B) OD; affected individual 12: (C) OS and (D) OD; affected individual 14: (E) OS and (F) OD; and unaffected individual 9: (G) OS and (H) OD; respectively. All affected individuals showed reduced amplitude and shortened implicit times in response to white flashes, suggesting an absence of rod function, whereas the unaffected individual had normal rod response in both eyes.

unaffected individual 9 (Fig. 3). Similarly, the scotopic ERG recordings measured at -25 dB were absent in all three affected individuals, when compared with those in unaffected individual 9 (Fig. 4). Taken together, these clinical data are suggestive of stationary night blindness in PKRP130 (Table 1).

Initially, *CABP4*, *GRM6*, and *SLC24A1* loci were excluded with closely spaced STR markers (data not shown). A genome-wide scan with four affected individuals (10, 11, 12, and 14) and five unaffected individuals (9, 13, 15, 16, and 17) was completed. A LOD score of 3.09 was obtained at *D3S3582* during the genome-wide scan. Additional markers from the

Marshfield database were selected to refine the critical interval, and two-point LOD scores of 3.07, 3.12, and 2.03 were obtained with markers *D3S3685*, *D3S3624*, and *D3S1581*, respectively, at $\theta = 0$ (Table 2). No suggestive or significant two-point LOD scores were obtained during the genome-wide scan other than *D3S3582*.

Haplotype analysis confirmed the linkage results and localized the critical interval to chromosome 3, region p22.1-p14.3. Recombination in unaffected individual 15 at marker *D3S3522* defines the proximal boundary, whereas recombination in affected individual 12 at marker *D3S1289* defines the distal

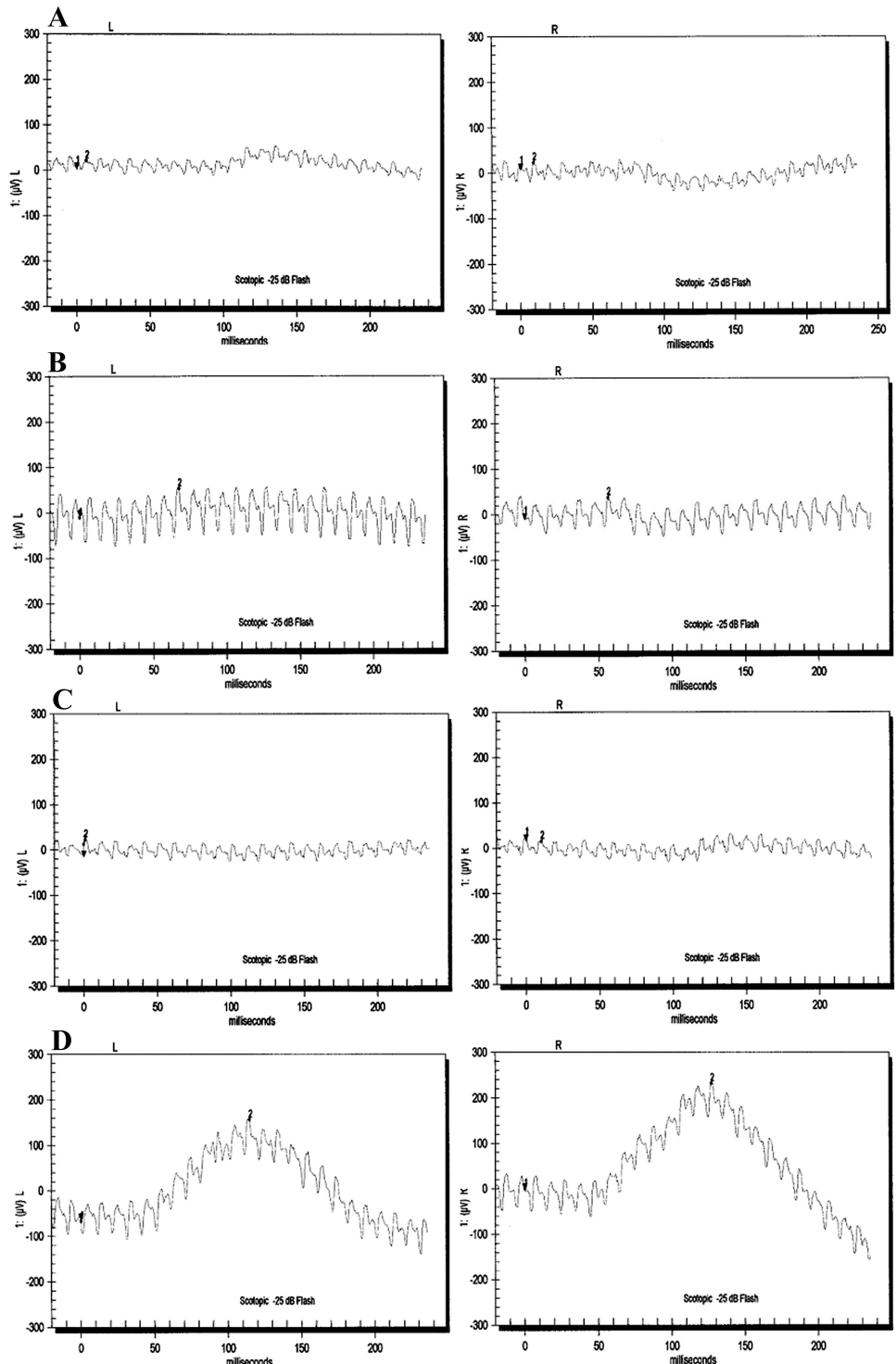


FIGURE 4. Scotopic ERG recordings of family PKRP130 recorded at -25 dB after 4 hours after dark adaptation. (A) OS and OD of individual 10; (B) OS and OD of individual 12; (C) OS and OD of individual 14; and (D) OS and OD of individual 19. All affected individuals show no rod response, whereas the unaffected individual has normal rod response in both eyes.

TABLE 1. Clinical Characteristics of Affected Individuals of Pedigree PKRP130 Diagnosed with Autosomal Recessive CSNB

Individual ID	First Symptoms	Visual Acuity		Color Vision	Fundus Findings	ERG Characteristics
		OD	OS			
10	Night blindness since early childhood	6/6	6/6	Normal	No macular atrophy, no pigment deposition, and no vascular attenuation.	The a- and b-waves are absent in the scotopic condition, whereas the cone responses are somewhat reduced in the photopic condition.
11	Night blindness since early childhood	6/6	6/6	Normal	ND	ND
12	Night blindness since early childhood	6/6	6/6	Normal	No macular atrophy, no pigment deposition, and no vascular attenuation.	The a- and b-waves are absent in the scotopic condition with normal cone responses in the photopic condition.
14	Night blindness since early childhood	6/6	6/6	Normal	No macular atrophy, no pigment deposition, and no vascular attenuation.	The a- and b-waves are absent in the scotopic condition with normal cone responses in the photopic condition.

ND, not determined.

boundary (Fig. 1). In addition, lack of homozygosity in affected individuals for alleles of markers *D3S3522* and *D3S3658* suggest that the pathogenic mutation lies distal to *D3S3658* (Fig. 1). Taken together, this places the critical interval in a 6.15 cM (13.57 Mb) region flanked by markers *D3S3658* proximally and *D3S1289* distally.

The critical interval harbors *GNAT1* which has been associated with autosomal dominant CSNB. Sequencing of all the coding exons of *GNAT1* in one affected and an unaffected individual of PKRP130 identified a novel homozygous missense mutation; c.386A>G that results in aspartic acid-to-glycine substitution; p.D129G (Fig. 5A). All affected individuals were homozygous for this substitution, whereas the unaffected individuals were heterozygous carriers of the mutant allele (Fig. 1). This mutation was not present in 192 ethnically matched control chromosomes.

To investigate the possible impact of p.D129G substitution on the GNAT1 protein, we used SIFT and PolyPhen software. SIFT prediction based on evolutionary conservation suggested that p.D129G substitution will not be tolerated by the native three dimensional structure of GNAT1 protein. Likewise, the position-specific score difference of 2.742 obtained from PolyPhen strongly suggested that the D129G substitution could have a deleterious effect on GNAT1 protein structure. We further analyzed the evolutionary conservation of the amino acids around the missense mutation (121–136 amino acids) by aligning *GNAT1* orthologs. The results strongly suggest that not only Asp129 but amino acids in the immediate neighborhood are well conserved among *GNAT1* orthologs (Fig. 5B). To further understand the effect of the p.D129G substitution on GNAT1 structure, we compared the three-dimensional models

of wild-type and mutant protein. Our results suggest that the wild-type sequence of D129 forms hydrogen bonds with the surrounding amino acids, but these interactions are compromised with G129 substitution (Figs. 5C, 5D).

Next, we investigated the quantitative expression of *Gnat1* transcript levels in ocular tissues at different postnatal intervals. Rhodopsin mRNA levels were measured in the same set of samples and served as a molecular marker for photoreceptor development. Both *Gnat1* and rhodopsin expression were not detected before P5; however, they increased exponentially from P7 indicating that *Gnat1* is present in the photoreceptors (Fig. 6A). The expression profile of *Gnat1* is similar to *Slc24a1*, a gene recently implicated in the pathogenesis of CSNB.²² The level of rhodopsin expression in the retina showed a steady increase with a maximum level at P30 and is in agreement with the profile of rhodopsin expression reported previously.^{32,33} Comparison of the expression levels of *Gnat1* in different eye tissues strongly suggested that *GNAT1* is highly expressed in the retina followed by the ciliary body, iris, and retinal pigment epithelium (Fig. 6B).

DISCUSSION

Herein, we report a consanguineous Pakistani family with CSNB that was recruited from the Punjab province of Pakistan. Clinical evaluation consisting of fundus photography and ERG responses confirmed the disease phenotype, and linkage analyses localized the critical interval to chromosome 3, region p22.1-p14.3, with statistically significant LOD scores. Sequential analyses identified a novel mutation, p.D129G, which seg-

TABLE 2. Two-Point LOD Scores of 3p Markers for Pedigree PKRP130

Marker	cM	Mb	0	0.01	0.05	0.09	0.1	0.2	0.3	z_{\max}	θ_{\max}
<i>D3S3522</i> *	65.26	40.78	-1.32	0.08	0.65	0.79	0.72	0.69	0.39	0.79	0.09
<i>D3S3658</i>	65.26	40.90	-1.54	0.07	0.60	0.74	0.71	0.59	0.35	0.74	0.09
<i>D3S3685</i>	67.94	42.46	3.07	3.03	2.77	2.51	2.44	1.76	1.08	3.07	0.0
<i>D3S3624</i>	68.47	44.61	3.12	3.08	2.81	2.56	2.48	1.81	1.12	3.12	0.0
<i>D3S3582</i> *	69.19	45.39	3.09	3.06	2.78	2.53	2.46	1.78	1.10	3.09	0.0
<i>D3S1581</i>	70.61	48.59	2.03	1.98	1.79	1.6	1.56	1.08	0.64	2.03	0.0
<i>D3S1289</i> *	71.41	54.47	−∞	−0.01	0.54	0.64	0.65	0.58	0.4	0.65	0.1

* Markers used in the genome-wide scan.

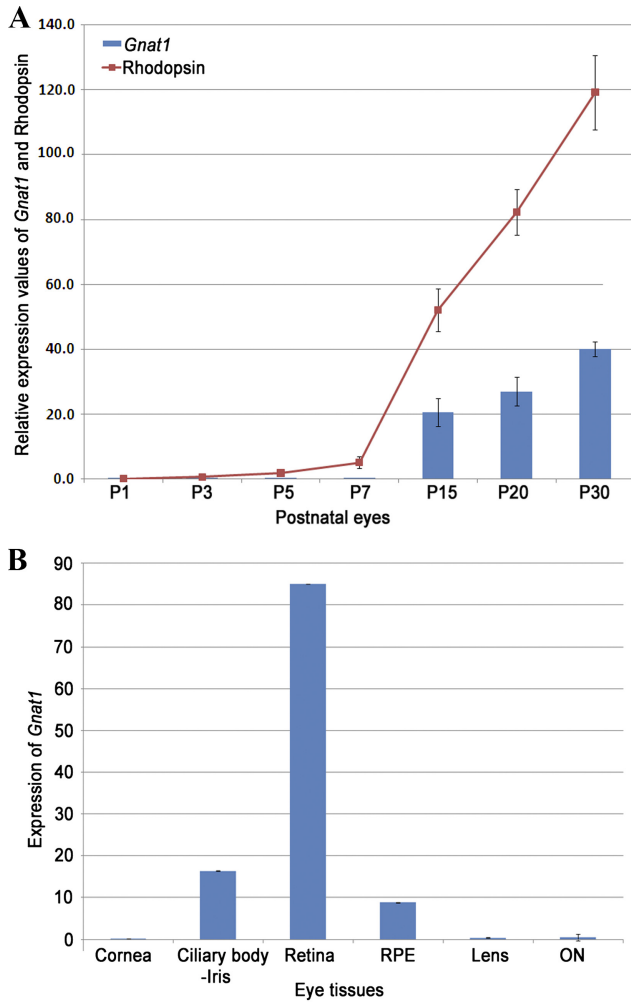


FIGURE 6. Expression pattern of *Gnat1* mRNA in postnatal mouse eyes. **(A)** Expression of *Gnat1* and *rhodopsin* postnatal mouse eyes at postnatal day (P)1, P3, P5, P10, P15, and P30. *Gnat1* expression is presented as bars and *rhodopsin* expression is presented as a line, with the relative scale on the y-axis. **(B)** Quantitative expression of *Gnat1* in adult mouse eye tissue (180 days) presented as an arbitrary scale. All the values are the mean (\pm SEM) of three independent observations after normalization with the control gene (*Rpl19*) expression. RPE, retinal pigment epithelium; ON, optic nerve.

been associated with autosomal dominant CSNB, this is the first report to implicate *GNAT1* in the pathogenesis of autosomal recessive CSNB.

The phenomenon of a gene associated with a certain phenotype but manifesting different modes of inheritance has been reported. Previously, we identified three homozygous pathogenic mutations in *RPI*, a gene implicated in the pathogenicity of autosomal dominant retinitis pigmentosa^{34–36} that segregated with the disease phenotype in their respective families with heterozygous carriers of these causal lesions clinically unaffected.³⁷ Likewise, causal mutations in crystalline β B1 have been associated with both autosomal dominant and autosomal recessive congenital cataracts.^{38,39} The mechanistic details of some mutations manifesting an autosomal dominant mode of inheritance, while other mutations in the same gene result in a recessive phenotype, are not yet completely understood. One possible explanation is that proteins harboring dominant mutations exhibit a negative effect on their wild-type counterparts; thus, one mutant allele is sufficient to manifest the disease phenotype. In contrast recessive mutations dem-

onstrate a local effect on the native structure of the mutant protein, without affecting the wild-type gene product that subsequently requires two mutant alleles to exhibit the affection status.

Identification of novel mutations and new genes associated with autosomal recessive CSNB will help us better understand the pathophysiology of the disease at a molecular level and will lead to development of better treatments.

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