Role of FOXC2 and PITX2 rare variants associated with mild functional alterations as modifier factors in congenital glaucoma

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Abstract

Congenital glaucoma (CG) is a severe and inherited childhood optical neuropathy that leads to irreversible visual loss and blindness in children. CG pathogenesis remains largely unexplained in most patients. Herein we have extended our previous studies to evaluate the role of FOXC2 and PITX2 variants in CG. Variants of the proximal promoter and transcribed sequence of these two genes were analyzed by Sanger sequencing in a cohort of 133 CG families. To investigate possible oligogenic inheritance involving FOXC2 or PITX2 and CYP1B1, we also analyzed FOXC2 and PITX2 variants in a group of 25 CG cases who were known to carry CYP1B1 glaucoma-associated genotypes. The functional effect of three identified variants was assessed by transactivation luciferase reporter assays, protein stability and subcellular localization analyses. We found eight probands (6.0%) who carried four rare FOXC2 variants in the heterozygous state. In addition, we found an elevated frequency (8%) of heterozygous and rare PITX2 variants in the group of CG cases who were known to carry CYP1B1 glaucoma-associated genotypes, and one of these PITX2 variants arose de novo. To the best of our knowledge, two of the identified variants (FOXC2: c.1183C>A, p.(H395N); and PITX2: c.535C>A, p.(P179T)) have not been previously identified. Examination of the genotype-phenotype correlation in this group suggests that the presence of the infrequent PITX2 variants increase the severity of the phenotype. Transactivation reporter analyses showed partial functional alteration of three identified amino acid substitutions (FOXC2: p.(C498R) and p.(H395N); PITX2: p.(P179T)). In summary, the increased frequency in PCG patients of rare FOXC2 and PITX2 variants with mild functional alterations, suggests they play a role as putative modifier factors in this disease further supporting that CG is not a simple monogenic disease and provides novel insights into the complex pathological mechanisms that underlie CG.
Primary congenital glaucoma (PCG; MIM# 231300) is a severe and irreversible neonatal or infantile optic neuropathy of uncertain pathogenesis. The immature iridocorneal angle appearance observed clinically indicates that it results from arrested maturation of tissues derived from cranial neural crest cells. This alteration leads to increased aqueous humor (AH) outflow resistance, elevated intraocular pressure (IOP) and optic nerve degeneration. The diagnosis is based in finding isolated trabeculodysgenesis with no other ocular anomalies [1]. PCG is the most common non-syndromic glaucoma in infancy [2], occurring in one of 10,000–20,000 live births in Western countries [3], and it presents increased incidence in consanguineous populations [4, 5]. PCG leads to significant visual loss and blindness in children. Autosomal-recessive inheritance is well documented in this disease, mainly due to CYTOCHROME P450, SUBFAMILY I, POLYPEPTIDE 1 (CYP1B1, MIM# 601771) mutations [6]. Absence of a family history and variable penetrance (ranging from 40% to 100% [6]) is reported in most cases, although a family history is more common in consanguineous families. Classical linkage analysis has identified loss-of-function variants in CYP1B1 [7] and LATENT TRANSFORMING GROWTH FACTOR-β-BINDING PROTEIN 2 (LTBP2, MIM# 602091) [8] as the cause of the disease in some patients. Recently, whole-exome sequencing revealed that mutations in the angiopoietin receptor TEK (TEK, MIM# 600221) underlie PCG with variable expressivity [9] and that rare and hypermorphic G-PATCH DOMAIN-CONTAINING PROTEIN 3 (GPATCH3, MIM# 617486) variants are present in a some congenital glaucoma (CG) cases [10]. Null CYP1B1 mutations are the predominant known genetic cause of this type of glaucoma in different worldwide populations [7, 11–13], and LTBP2 gene alterations have been identified only in a few families [8, 14, 15]. MYOCILIN (MYOC, MIM# 601652) [16, 17] and FORKHEAD BOX C1 (FOXC1, MIM# 601090) alterations [18, 19] have also been found in a small number of CG cases. The genetic heterogeneity present in PCG, along with the frequent incomplete penetrance and variable expressivity, even in patients with null CYP1B1 genotypes [13], strongly indicates the participation of modifier genetic and/or environmental factors in the pathogenicity of this type of glaucoma. Previous studies from our laboratory have provided evidence for the role of rare FOXC1 variants with moderate functional dysregulation as either causative of modifier factors in CG [18–20].

FORKHEAD BOX C2 (FOXC2, MIM# 602402) and PAIRED-LIKE HOMEODOMAIN TRANSCRIPTION FACTOR 2 (PITX2, MIM# 601542), are genes encoding transcription factors that control the expression of other genes and form part of a complex regulatory network involved in ocular development and disease [21]. FOXC2 is structurally and functionally closely related to FOXC1. Their chromosomal localization and genomic organization suggest a common origin from an ancestral gene through chromosomal duplications [22, 23]. The proteins encoded by these two genes share 98% sequence identity at their forkhead domain with only two amino acid differences [24], and present overlapping expression patterns during embryonic development, including similar expression in the periocular mesenchyme and in periocular mesenchyme-derived tissues involved in formation of the AH drainage pathway [25–27]. Either a complete loss or a significant gain of FOXC2 function cause lymphedema-distichiasis syndrome, an autosomal dominant disease that affects the formation of the lymphatic vasculature system and is also characterized by a double row of eyelashes (distichiasis) [28].

The PAIRED-LIKE HOMEODOMAIN TRANSCRIPTION FACTOR 2 (PITX2, MIM# 601542) gene encodes a homeodomain-containing transcription factor. It presents a pattern of expression similar to that of FOXC1 and FOXC2, and it is also present in the periocular mesenchyme. Disruption of PITX2 is one of the major causes of Axenfeld-Rieger syndrome (ARS).
ARS is a clinically and genetically heterogeneous group of developmental dominant disorders, affecting primarily the anterior segment of the eye, frequently associated with secondary glaucoma. Systemic alterations, such as dental defects, mild craniofacial dysmorphism and umbilical anomalies may also be present in ARS patients. ARS is a fully penetrant disease with variable expressivity. PITX2, like FOXC1, is dose sensitive and mutations that alter the level of functional PITX2 protein (either increased or decreased) by different mechanisms are pathogenic. The PITX2 gene is composed of six exons, two alternative promoters located upstream of exons 1 and 4, and two alternative transcription start sites located in exons 2 and 4. The immature mRNA is alternatively spliced to produce four isoforms (PITX2A-D). The first three isoforms share an identical homeodomain and C-terminal region, and they differ in their N-terminal sequences.

In this study we show an increased frequency of rare and functionally altered FOXC2 and PITX2 variants in CG patients, providing further evidence for the role of rare variants of these two genes involved in ocular anterior segment development as putative modifier factors.

Materials and methods

Subjects

One hundred and thirty-three unrelated families affected by PCG participated in this study. The study and informed written consent procedures were approved by the Ethics Committee for Human Research of the Hospital Clínico San Carlos, Madrid (Spain), approval number 13/388-E and followed the tenets of the Declaration of Helsinki. The clinical examination and diagnosis of patients were performed as previously described. Glaucoma was ruled out in 100 control individuals, who were recruited among patients diagnosed with cataracts, floaters, refractive errors or itchy eyes.

Variant screening

The genomic DNA was extracted from peripheral blood, using the QIAamp DNA Blood Mini Kit (Qiagen, Valencia, CA, USA). The DNA sequence variation analyses were carried out by automatic Sanger sequencing. The promoter region (nucleotides -1 to -1500), the coding region and both the 5' and 3' untranslated regions (UTRs) of FOXC2 were amplified by PCR, using the primers and conditions described in S1 Table. The coding region, 5' and 3' UTRs of PITX2 were amplified as previously described. The variants identified in each subject were confirmed by sequencing a new amplification product. The patients with rare FOXC2 variants were screened for LTBP2 mutations using a customized gene panel. Library capture of all exons and 20 bp of intronic boundaries was performed using SureSelect QXT technologies (Agilent Technologies, Santa Clara, CA, United States). Massive sequencing was carried out using Illumina MiSeq or NextSeq 500 platforms running on paired-end mode at a minimum of 450X. Bioinformatic analysis was performed using standard procedures and custom in-house pipelines for mapping, variant calling and annotation.

Cloning and site directed mutagenesis of FOXC2 and PITX2 variants

The FOXC2 variants p.(C498R) (rs61753346) and p.(H395N) were directly amplified and cloned from the genomic DNA of carriers using the following primers, which incorporated the EcoRI or BamHI restriction sites at the 5' end (indicated in bold): FOXC2-Up-EcoRI, 5' -GGGAATTCCGCCGGTCTCTCGTCTCAGG-3' and FOXC2-Dw-BamHI, 5' -GGGGATCCCGTATTTCTGTCGACGCAGAGG-3'. The PCR products were cloned into a modified version of the EcoRI/BamHI restriction sites of the pcDNA3.1(-) vector (C-terminal myc-tagged).
The same approach was used to clone the wild-type FOXC2 coding sequence from DNA of a control subject and the cloned DNA was completely sequenced to rule out the presence of rare FOXC2 variants. The PITX2 coding variants p.(P179T) was obtained by site-directed mutagenesis using the QuickChange Site-Directed Mutagenesis Kit (Agilent, Santa Clara, CA, USA), with the following primers (mutant nucleotides are indicated in bold): F, 5’-CCTAC GACGACATGTACAAGGCTATTCCTACAAC-3’ and R, 5’-GTTGTAGGAATAGCCTGACATGTCGTCGTAAGG-3’. A cDNA encoding the wild-type PITX2C (NP_000316) isoform, cloned into the pcMV6 vector (Origene, NM_000325), was used as a template for the site-directed mutagenesis reaction. The presence of the mutation in the site-directed mutagenesis product was confirmed by Sanger sequencing. The mutagenized cDNA was subcloned into the SgfI and MluI restriction sites of pcMV6 to avoid undesirable mutations in the vector.

**Transactivation luciferase reporter assays**

Transactivation assays were performed using the Luciferase Assay System (Promega) following the instructions of the manufacturer. HEK-293T cells were transfected with the different recombinant versions of FOXC2 and PITX2C, cloned in the expression vector pcMV6 (500 ng each), along with the recombinant pGL3-basic-CXCR4 luciferase reporter vector (50 ng), and the pMirTarget vector (50 ng), which expresses the red fluorescent protein (RFP) as a transfection control. The cells were cultured in Dulbecco’s modified Eagle’s medium (DMEM) supplemented with 10% fetal bovine serum (FBS) and antibiotics (Normocin, Invivogen), at 37˚C in a fully humidified 5% CO₂ atmosphere. Forty-eight hours after transfection the cells were harvested and assayed for firefly luciferase activity using the Luciferase Assay System (Promega) according to the manufacturer’s instructions. FOXC2 and PITX2 proteins were detected by Western blot as expression control and transactivation activity was normalized for these proteins. RFP and endogenous lactate dehydrogenase (LDH) were also assessed by Western blot as transfection and loading controls, respectively.

**Protein stability and half-life calculation**

The protein stability was studied via western blot of transfected cells treated with cycloheximide (300 μg/ml) at different times as previously reported [34]. The FOXC2 and PITX2 protein levels were determined by densitometry and the relative amounts at different time points after cycloheximide treatment were expressed as a percentage of the levels at time 0 h. At least three independent assays for each variant were performed. The transfection efficiency was assessed by co-transfeting the recombinant cDNA constructs encoding the different mutants (500 ng) with the pMirTarget vector (50 ng), which encodes the red fluorescent protein (RFP). The RFP was detected via Western blotting. The loaded samples were normalized for total protein content using the Bradford reagent (Pierce, Rockford, IL, USA). FOXC2 and PITX2 decay followed a first order kinetics. The slope of the decay line was calculated using standard linear regression, and the protein half-life was determined as previously described [35].

**Western blotting and antibodies**

For Western blot analyses the cell lysates were prepared and fractionated using sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE), using the Mini-PROTEAN III Gel Electrophoresis System (BioRad, Hercules, CA, USA). A commercial mouse monoclonal anticyc (sc-40, Santa Cruz Biotechnology, Santa Cruz, CA, USA) was used as the primary antibody, diluted at 1:1000. Horseradish peroxidase-conjugated antibodies against mouse IgG (#32439, Invitrogen) were diluted to 1:1000–1:4000. Chemiluminescence detection and the densitometry for protein band quantification was performed as previously described [36]. As
an additional sample loading control, the endogenous lactate dehydrogenase (LDH) protein was detected in cell extracts using a goat anti-LDH antibody diluted to 1:5000 (AB1222, Chemicon, Temecula, CA, USA) and an anti-goat IgG horse-radish peroxidase-conjugated antibody (sc-2033, Santa Cruz Biotecchnology, diluted to 1:2000). RFP (transfection control) was detected using a rabbit anti-RFP antibody (#AB233, Evrogen, Moscow, Russia), diluted to 1:5000, and an anti-rabbit IgG horse-radish peroxidase-conjugated antibody (#1858415, Pierce), diluted to 1:1000.

Bioinformatic analyses

The deleterious effect of mutations was predicted online with the Sorting Intolerant From Tolerant (SIFT) (Kumar et al., 2009) and PolyPhen-2 (http://genetics.bwh.harvard.edu/pph2/) programs. The MicroSNiPer software was used to identify putative miRNA target sequences (http://cbdb.nimh.nih.gov/microsniper) (Barenboim et al., 2010). Both nucleotide and amino acid sequence alignments were carried out with Clustalw (Thompson et al., 1994). The novel FOXC2 and PITX2 variants were named using directions from Mutalyzer (https://humgenprojects.lumc.nl/trac/mutalyzer), according to RefSeq NM_005251.2 and NM_000325.5, respectively. The novel variants have been submitted to dbSNP (http://www.ncbi.nlm.nih.gov/SNP/) and to ClinVar (https://www.ncbi.nlm.nih.gov/clinvar/). Structure prediction of mutant and wild-type proteins were evaluated by QUARK (https://zhanglab.ccmb.med.umich.edu/QUARK2/) [37] and further visualized using Pymol Molecular Graphics System (Pymol).

Statistical analysis

The statistical comparisons between groups were performed using either the t-test or the one-way analysis of variance (ANOVA). A Bonferroni correction was applied to adjust tests for multiple comparisons. The data were statistically processed by the SigmaStat 2.0 software (SPSS Science Inc., Inc., Chicago, IL, USA).

Results

FOXC2 and PITX2 variants

To further investigate the role in CG of variants of genes involved in ocular development we have extended our previous genetic studies on a large cohort of 133 apparently unrelated PCG families, in which we reported an increased frequency of rare and functionally altered FOXC1 variants [18, 20]. In the present work we evaluated the role of FOXC2 and PITX2 variants in CG using the same group of patients. The main clinical and demographic features of these patients have previously been reported [19]. These patients had no known disease-causing variants in the coding regions of MYOC and CYP1B1. Eight of the patients (6.0%) carried rare FOXC2 variants in the heterozygous state (Table 1 and S1 Fig) and in contrast, none of them presented rare PITX2 variants. Rare FOXC1 and FOXC2 variants were not coinherited in any of these patients. In addition, the presence of pathogenic LTBP2 variants was ruled out in these subjects by Next Generation Sequencing. All the FOXC2 carriers identified in this study had bilateral PCG and most of them were diagnosed before the fourth month of life and required several surgical interventions for adequate control of IOP (Table 2). Segregation analyses of the identified variants did not show a monogenic pattern of co-inheritance with glaucoma, and these variants were not detected in 100 Spanish control individuals (200 chromosomes). The p.(C498R) (rs61753346) substitution was found in two probands (PCG-190 and PCG-143) with different age at diagnosis and evolution of glaucoma (Table 2). Patient PCG-190 was diagnosed with PCG at birth, his left eye was eviscerated and required two surgical operations.
for adequate control of IOP. However, patient PCG-143 was diagnosed at the age of five years and his IOP was well controlled with medication. This variant was located in the inhibitory domain of the protein (Fig 1A) and its predicted effect evaluated with the Ensembl Variant Effect Predictor [38] ranged from moderate to deleterious (0.01) Promoter: modifier/-/-. Moreover, structural prediction using QUARK showed that cysteine 498 is located in a disordered region of the protein and although its substitution by arginine is inferred to cause weak conformational disruption (Fig 2A and Fig 2B), the presence of a positive charge in this position of the mutant polypeptide chain may affect chemical interactions and protein function. Interestingly, this variant and the rest of \textit{FOXC2} nucleotide changes identified in this study, overlapped a regulatory feature (ENSR00000089180), which is annotated in Ensemble and GeneCards databases as a promoter covering the entire \textit{FOXC2} sequence and most of the antisense lncRNA \textit{FOXC2-AS1} gene (Fig 1C). The synonymous change p.(S36S) (rs138318843) was carried by four patients (3%), showing a 4- to 10-fold increased frequency compared with that reported in 1000 Genomes and ExAC databases (0.3% and 0.7%, respectively, Table 1). The variant was associated with variable age at diagnosis and IOPs, ranging from two and a half months to three years and from 22 to 45 mmHg, respectively (Table 2). p.(S36S) carriers required different surgical procedures for correct IOP control (Table 2). This nucleotide substitution also mapped at \textit{FOXC2-AS1}

### Table 1. \textit{FOXC2}, \textit{PITX2} and \textit{CYP1B1} variants identified in congenital glaucoma patients.

| Family | Variant* | %, 1000 Genomes/ExAC (number of homozygotes) | Functional assay | EVEP/Polyphen/\textit{SIFT} Hamiltonian
|--------|----------|-----------------------------------------------|----------------|-----------------------------------------|
| PCG-143 | \textit{FOXC2}: c.1492T>C, p.(C498R) (rs61753346) | 0.03/0.094 (1) | hypomorphic | \textit{FOXC2}: Moderate/Probably damaging (0.987)/deleterious (0.01) Promoter: modifier/-/-
| PCG-190 | \textit{FOXC2}: c.38T>G (rs199552394) | 0.2/0.1 (1) | \textit{FOXC2}: modifier/-/ Promoter: modifier/-/-
| PCG-209 | \textit{FOXC2}: c.108C>T, p.(S36S) (rs138318843) \textit{FOXC2-AS1}: n.145+174G>A | 0.3/0.7 (5) | \textit{FOXC2}: low/-/-; \textit{FOXC2-AS1}: modifier/-/ Promoter: modifier/-/-
| PCG-8 | \textit{PCG-36} | \textit{PCG-163} | \textit{PCG-171} | \textit{PCG-150} | \textit{PCG-133} | \textit{PCG-139}

*All variants were present in the heterozygous state.

Polyphen scores: 0.0–0.15, benign; 0.15–1.0, possibly damaging; 0.85–1.0, damaging.

Threshold for SIFT intolerance is 0.05.

\textit{FOXC2} RefSeq: NM_005251.2.

\textit{Ensembl} transcript ID: ENSR00000089180 (regulatory feature).

\textit{Ensembl} \textit{FOXC2-AS1} transcript ID: ENST00000563280.1.

\textit{PITX2} RefSeq: NM_000325.5; \textit{PITX2C} amino acid sequence was used as a reference to number coding variants (NP_000316).

N.I.: Not Identified; EVEP: Ensembl Variant Effect Predictor. NA: not available. -: not apply.

https://doi.org/10.1371/journal.pone.0211029.t001
intron 1 (n.145+174G>A) and the regulatory feature (promoter) (Fig 1C), and it was inferred to produce a low functional effect on FOXC2 and a modifier outcome on both FOXC2-AS1 and the overlapping promoter. The nucleotide change c.3’8T>G was located in the 3’UTR of the gene (Fig 1A) and was identified in one patient diagnosed at the age of two and a half months, who required several trabeculectomies to correct IOP (Table 2). This variant had a frequency of 0.1–0.2% in the two genome databases used in the study and was bioinformatically classified as a modifier nucleotide substitution (Table 1) in the two overlapping genomic elements (Fig 1C), and according with microSNIPER it does not affect microRNA targets. The last identified FOXC2 nucleotide change was predicted to cause a novel amino acid change (p. (H395N)) located in the second transactivation domain of the protein (Fig 1A) with inferred moderate functional effect (Table 1). The structural prediction showed that the substitution of histidine 395 with asparagine in the polypeptide chain increases the length of an alpha helix.

Table 2. Rare FOXC2 and PITX2 gene variations identified in PCG and associated clinical features.

| Proband (family member) | Nucleotide change | Amino acid change | Laterality/ Sex | Age at diagnosis | IOP (mmHg) at diagnosis (OD/OI) | C/D (OD/OS) | Number and type of surgery (OD/OS) |
|-------------------------|------------------|-------------------|----------------|----------------|---------------------------------|-------------|----------------------------------|
| PCG-143 FOXC2: c.1492T>C (rs61753346) | p.(C498R) | B/F | 5y/F | 21/21 | 0.5/0.1 | No surgery |
| PCG-190 FOXC2: c.1492T>C (rs61753346) | p.(C498R) | B/M | 0m/M | NA/EE | 0.5/EE | No surgery/EE |
| PCG-209 FOXC2: c.38T>G (rs199552394) | - | B/M | 2.5m | 26.5/22 | NA (leukoma) | 3T/2T |
| PCG-8 FOXC2: c.108C>T (rs138318843) FOXC2-AS1: n.145 +174G>A | FOXC2: p.(S36S) | B/M | 2.5m | 22/22 | NA | CTT/CTT |
| PCG-36 FOXC2: c.108C>T (rs138318843) FOXC2-AS1: n.145 +174G>A | FOXC2: p.(S36S) | B/M | 4m | NA | NA | G/G |
| PCG-163 FOXC2: c.108C>T (rs138318843) FOXC2-AS1: n.145 +174G>A | FOXC2: p.(S36S) | B/F | 3m | 45/40 | 0.8/0.7 | G+T/3G |
| PCG-171 FOXC2: c.108C>T (rs138318843) FOXC2-AS1: n.145 +174G>A | FOXC2: p.(S36S) | B/M | 3y | NA | 0.9/1.0 | T/T |
| PCG-150 FOXC2: c.1183C>A | p.(H395N) | B/M | 3m | NA | 0.8/0.5 | G+T/T |
| PCG-133 (II:1) PITX2: c.535C>A (de novo). CYP1B1: c.535delG/c.1159G>A | PITX2: p.(P179T) CYP1B1: A179fs*18/E387K | B/F | 3m | 24/22 | 0.8/0.8 | G+2T+/2G+3T |
| PCG-133 (II:2) CYP1B1: c.535delG/c.1159G>A | CYP1B1: A179fs*18/E387K | B/M | 10y | NA | NA | NA |
| PCG-139 PITX2: c.562G>A (rs77144743). CYP1B1: c.1159G>A | PITX2: p.(A188T) CYP1B1: E387K/E173* | B/M | 0m | NA | 0.4/0.5 | 2G+T/EE |

B/U: bilateral/unilateral; CTT/G/T: combined trabeculotomy-trabeculectomy/goniotomy/trabeculectomy; F/M: female/male; m/y: months/years; NA: not available; OD/OS: right eye/left eye; EE: eviscerated eye 

aFOXC2 RefSeq: NM_005251.2.

bEnsembl FOXC2-AS1 transcript ID: ENST00000563280.1.

cPITX2 RefSeq: NM_000325.5; PITX2C amino acid sequence was used as a reference to number coding variants (NP_000316).

https://doi.org/10.1371/journal.pone.0211029.t002
(Fig 2B and Fig 2C). In addition, the bioinformatic analysis indicated that it was a modifier variant for the overlapping FOXC2-AS1 gene and the promoter (Table 1 and Fig 1C). The patient who carried this variant was diagnosed at the age of three months, showed severe optic nerve excavation and was subjected to one goniotomy in the right eye and two trabeculectomies in the left eye (Table 2).

**Coinheritance of PITX2 and CYP1B1 rare variants**

We also evaluated the possible modifier role of FOXC2 and PITX2 defects on patients with CYP1B1 glaucoma-associated genetic backgrounds. To that end, a group of 25 CG cases who were known to carry CYP1B1 pathogenic genotypes, were selected based on DNA availability from a previously reported cohort of patients [13]. Two heterozygous PITX2 variants were identified in two (8%) of these patients (PCG-133 and PCG-139). As far as we know, one of these variants (c.535C>A, p.(P179T), S1 Fig) was novel, was located in a transcriptional inhibitory domain (Fig 1B) and segregation analysis revealed that although the progenitors were heterozygous carriers for the CYP1B1 variants none of them presented this PITX2 nucleotide change (Fig 3), indicating that it was acquired de novo. The patient who carried this variant (PCG-133) was diagnosed with bilateral PCG at the age of three months and presented advanced optic nerve excavations and was subjected to six surgical operations per eye to control IOP (Table 2). Interestingly, her brother, who inherited the same CYP1B1 genotype and did not present the PITX2 variant, was diagnosed with bilateral infantile glaucoma at the age of three months.
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of ten years. The predicted amino acid substitution was classified as not tolerated by SIFT, although it was considered benign by Polyphen (Table 1). The structural prediction showed no significant conformational changes (Fig 2E and Fig 2F), although the different chemical properties of the mutant amino acid might affect PITX2 molecular interactions. Altogether, the data indicate that this de novo PITX2 variant is a possible genetic modifier of glaucoma onset. The second PITX2 variant identified among CYP1B1 deficient patients was a coding nucleotide substitution (c.562G>A), that resulted in an inferred missense mutation (p.(A188T)), (S1 Fig) also located in the first transcriptional inhibitory domain (Fig 1B). This nucleotide change had a frequency of 0.1–0.36% in the two databases used in this study (Table 1). The variant was classified as benign or tolerated by the bioinformatic analyses and structural prediction indicated that the conformation of the protein is not significantly disrupted by this amino acid change (Fig 2G and Fig 2H). The carrier of this combined genotype (subject PCG-139) was diagnosed with severe bilateral PCG at birth, which required from seven to eight surgical operations per eye to achieve an adequate IOP control. The number of glaucoma surgical operations of the two PITX2 was higher than the average for other patients. These data suggest a modifier effect of this rare PITX2 variant on age at onset and severity in CYP1B1 congenital glaucoma patients.

Fig 2. Prediction of conformational changes induced by the identified coding FOXC2 and PITX2 variants. Predicted model structures of wild type (left) (A, C, E and G) and mutant (right) proteins (B, D, F and H) were obtained with QUARK as described in Materials and Methods.

https://doi.org/10.1371/journal.pone.0211029.g002

Fig 3. Evidence for modifier effect of rare PITX2 variants on age at onset and severity in CYP1B1 congenital glaucoma patients. All mutations were detected in the heterozygous state. Note that subject II:1 in family PCG-133 was diagnosed at the age of 3 months and carried the de novo p.(P179T) PITX2 variant, whereas his brother, who did not carry this variant, was diagnosed at the age of 10 years. The proband in family PCG-139 also carried a rare PITX2 variant (p.(A188T)) and presented glaucoma diagnosed at the age of seven days. Both probands required more surgical operations to control IOP than the rest of patients. Below symbols are indicated genotypes for CYP1B1 and PITX2, age at diagnosis and number or surgical operations per eye, respectively. M1, CYP1B1: p.(A179fs*18). M2, CYP1B1: p.(E387K). M3, CYP1B1: p.(E173*). M4, PITX2: p.(P179T). M5, PITX2: p.(A188T). Arrows show the index cases. +: wild-type allele. The asterisk indicates a de novo PITX2 variant.

https://doi.org/10.1371/journal.pone.0211029.g003
Evolutionary conservation of FOXC2 and PITX2 variants

Evolutionary amino acid or nucleotide sequence conservation analysis were assessed using a multiple sequence alignment of representative orthologous proteins or genes of seven different species, from fish to human. This analysis revealed that most FOXC2 and PITX2 wild-type amino acids or nucleotides were highly conserved (Fig 4). These data indicate that the identified variants affect functionally important positions, and therefore that they may affect the function of these genes and the corresponding proteins.

Transactivation activity, protein stability and subcellular localization

Transactivation activity of two FOXC2 and one PITX2 coding variants was evaluated in HEK-293T cells in culture by transient co-expression of the different variants with luciferase fused to the CXCR4 proximal promoter, which has been reported to be activated by both FOXC1 and FOXC2 proteins [40]. To the best of our knowledge the CXCR4 promoter has not previously been used to assess PITX2 transactivation. However, we obtained a positive response in the initial assays carried out to set up the functional evaluation of PITX2, and we decided to use this promoter in further experiments. The results showed that the two FOXC2 variants p.(H395N) and p.(C498R) were associated with a 22%-28% reduced transactivation (Fig 5A). Only one of the two identified PITX2 variants, p.(P179T), could be cloned for functional evaluation. It showed approximately 130% increased transactivation compared with the wild type (Fig 5B). Control Western blot analysis of RFP (transfection control) and LDH (loading control) indicated that these results were not biased by significant loading or transfection differences (Fig 5).

The effect of the FOXC2 and PITX2 variants on protein stability was also assessed by transient expression in HEK-293T cells and measurement of recombinant protein levels by Western blot at different times after protein synthesis inhibition with cycloheximide. p.(C498R) stability was not significantly different from that of FOXC2 wild-type protein (Fig 6A), with an estimated half-life of 25.4 h (Fig 6B). These data indicate that p.(C498R) alter transcriptional activity independently of protein stability. However, p.(H395N) showed an approximately 20% reduced stability after 24 h of protein synthesis inhibition (Fig 6A), and a 1.5-fold...
decreased half-life compared with the wild-type protein (Fig 6B) which may explain the decreased transactivation observed in the previous functional assay. The PITX2 variant p.(P179T) did not show significant protein stability differences with the wild-type protein (Fig 6C and Fig 6D).

Fluorescence immunocytochemistry of the different recombinant FOXC2 and PITX2 coding variants transiently expressed in HEK-293T cells showed clear nuclear localization. Expression analysis of recombinant myocilin was used as a negative control of nuclear localization, because this protein is known to be absent in the nucleus and present in cytoplasmic secretory

Fig 5. Altered transcriptional activity of FOXC2 and PITX2 variants identified in this study. cDNA constructs encoding the different variants were transiently co-expressed in HEK-293T cells along with cDNAs constructs encoding luciferase coupled to the CXCR4 promoter (transactivation reporter) and the red fluorescent protein (RFP) as a transfection control. The transcriptional activity was normalized to the amount of the corresponding wild type protein and represented as percentage of the luciferase activity of the wild-type protein. Each lane contained 15 μg of total protein obtained from the cell lysates. The protein levels of the different FOXC2 and PITX2 versions present in HEK-293T cells 24 h after transfection were determined by western immunoblot using a monoclonal anti-myc antibody (1:300) (Santa Cruz). RFP was analyzed via western immunoblot using an anti-RFP antibody (1:500) (Evrogen). The sample loading control, endogenous LDH, was also detected via immunoblot using an anti-LDH antibody (1:1000) (Chemicon). Error bars correspond to the SD of two independent experiments carried out in triplicate. Asterisks indicate statistical significance as compared to the control: P<0,01 (**); P<0,001 (***). Significance was calculated by one-way ANOVA followed by Tukey multiple-comparison test and t-student test.

https://doi.org/10.1371/journal.pone.0211029.g005
vesicles and endoplasmic reticulum and [33]. These analyses demonstrated that the identified coding FOXC2 and PITX2 variants do not affect subcellular localization (Fig 7).

Discussion

Elevated frequency of rare FOXC2 and PITX2 variants in congenital glaucoma patients and evidence of non-Mendelian inheritance

It is accepted that CG is caused by developmental abnormalities of the ocular trabecular meshwork [41]. Herein, we have extended our previous investigations on CG genetics to study the role of FOXC2 and PITX2 gene variants in our cohort of patients. As a rationale for these studies we have proposed that different combinations of moderate functional alterations of genes involved in development of the anterior segment of the eye might contribute to development of relatively mild phenotypic alterations, limited to glaucoma, with no other systemic and ocular alterations. This hypothesis implies an oligogenic or polygenic genetic background for this disease, at least in some patients. This and our previous study reveal a higher than expected by chance frequency of rare FOXC2 and FOXC1 [19, 20] variants in the analysed cohort of PCG patients (6% and 7.5%, respectively), with either experimentally assessed or inferred moderate

Fig 6. FOXC2 p.H395N variant decreases protein stability. Time course stability analysis of FOXC2 (A) and PITX2 (C) coding variants found in PCG patients was carried out by transient expression in HEK-293T cells. Transfected cells were treated with the protein synthesis inhibitor cycloheximide and the different recombinant proteins were detected by Western immunoblot using an anti-myc monoclonal antibody (Santa Cruz) at the indicated time points, as explained in Materials and Methods. Relative amounts of FOXC2 are expressed as a percentage of levels at time 0 h. The rate of decay and half-lives of the recombinant FOXC2 (B) and PITX2 (D) coding variants at the indicated time-points were determined from linear regression analysis as described in the Materials and Methods. Error bars correspond to the SD of three independent experiments carried out in triplicate. Asterisks indicate statistical significance compared to the control: p<0.01 (**). Two-way ANOVA followed by Tukey multiple-comparison test.

https://doi.org/10.1371/journal.pone.0211029.g006
FOXC2 and PITX2 rare variants as modifier factors in congenital glaucoma

Table 1. Subcellular localization of recombinant proteins.

| Gene  | Variant | Localization |
|-------|---------|-------------|
| FOXC2 | Wild Type | Membrane |
| FOXC2 | p.(H388N) | Cytoplasm |
| FOXC2 | p.(C498R) | Nucleus |
| PITX2 | Wild Type | Nucleus |
| PITX2 | p.(P179T) | Cytoplasm |
| MYOC  | Wild Type | Cytoplasm |

Fig 7. Coding FOXC2 and PITX2C variants identified in this study do not alter subcellular localization in cells in culture. cDNA constructs encoding the indicated recombinant versions of FOXC2 and PITX2C were transiently expressed in HEK-293T cells. The recombinant proteins were detected by fluorescent immunocytochemistry with an anti-myc antibody (Santa Cruz). Nuclei were visualized by fluorescent DAPI staining. Original magnification: X600.

https://doi.org/10.1371/journal.pone.0211029.g007
functional defects. Segregation analysis of these variants, which were present in the heterozygous state, rule out a monogenic inheritance pattern, indicating according to our earlier reports [19, 20], that these genetic changes might be involved in either oligogenic or complex transmission of the disease. Moreover, the existence of incomplete penetrance, variable expressivity and of a relatively high proportion (close to 20%) of PCG patients with rare heterozygous CYP1B1 variants [13, 34] also suggest non-Mendelian PCG transmission in some cases. In these patients, disease outcome might depend on modifier factors (genetic, stochastic and/or environmental) [13, 42], as will be discussed later.

Functional impact of the rare variants

The two missense FOXC2 variants (p.(H395N) and p.(C498R)) and one of the PITX2 amino acid substitutions (p.(P179T)) were inferred to cause a moderate functional effect at least by one bioinformatic analysis and, experimentally, they were found to be associated with moderately disrupted transactivation. The functional impact of the second PITX2 amino acid substitution, p.(A188T), could not be functionally evaluated due to DNA cloning difficulties. In fact, the two FOXC2 amino acid changes were found to be hypomorphic whereas the PITX2 amino acid substitution (p.(P179T)) behaved experimentally as a hypermorphic variant. Additional structural and functional analysis indicated that p.(H395N) alters polypeptide chain conformation and decreases protein stability, which can explain the associated reduced transactivation. However, our data showed that the transactivation effect of p.(C498R) is not due either to altered protein stability, protein structure or subcellular localization, suggesting that the substitution of a basic (arginine) for a polar (cysteine) amino acid may disrupt protein interactions in the second inhibitory domain of the protein, leading to reduced protein activity. Similarly, our results indicate that increased transactivation associated with the p.P179T PITX2 mutation is not related with altered protein stability, protein conformation or subcellular localization. Proline possesses a hydrophobic side chain, whereas threonine side chain has both hydrophilic and hydrophobic functions. Therefore, this amino acid replacement may affect protein interactions taking place in the transcriptional inhibitory domain where it is located, leading to increased PITX2 activity. In this line, an increased side chain polarity associated with amino acid substitution p.(A188T) could also interfere protein interactions involving the first PITX2 transcriptional inhibitory domain, leading to a functional alteration. Additional studies are required to evaluate these hypotheses. Interestingly, according to Ensembl Regulatory Build, FOXC2 variants p.S36S (synonymous) and c.38T>G (non coding 3’ UTR) also mapped at a promoter, which overlapped with FOXC2 and FOXC2-AS1 genes. FOXC2-AS1 encodes a lncRNA transcribed from the negative strand of FOXC2. Both molecules have been located in the cytoplasm and are concordantly upregulated in doxorubicin-resistant osteosarcoma cell lines and tissues [43]. It has also been suggested that FOXC2-AS1 stabilizes FOXC2 mRNA by forming an RNA-RNA duplex, which subsequently upregulated steady-state FOXC2 expression at both the transcription and post-transcription levels [43]. Therefore, the non-coding nucleotide substitutions and the synonymous variant also have the potential to disrupt gene expression by affecting the promoter. Additionally, p.(S36S) might affect FOXC2 activity through FOXC2-AS1 dysregulation. Further studies are required to confirm these ideas. Finally, the high evolutionary conservation of wild type FOXC2 and PITX2 nucleotide or amino acid positions involved in the variants identified in this study, further indicate that their substitution by infrequent residues might modify their function.

Mild functional alterations of FOXC2 and PITX2 as modifier factors in PCG

Modifier genes are increasingly recognized as an important source of phenotypic variation that may explain the relationship of phenotype to genotype. Individually, genetic modifiers
have small effect size on the phenotypic outcome, leading to reduced penetrance, i.e., not all individuals harbouring a pathogenic variant will exhibit clinical signs of the associated disorder [44]. Normal ocular iridocorneal angle development is the result of highly regulated and intricate gene and protein interactions that can be disrupted by different combinations of subtle functional alteration in multiple genes. In line with this idea, anterior segment abnormalities and CG may manifest in individuals who are heterozygous for functionally altered alleles at multiple genes, resulting in a clinically significant disruption of trabecular meshwork development. This phenomenon is known as synergistic heterozygosity [45]. A representative identified molecular interaction that regulates anterior segment development involves FOXC1 and PITX2A. In fact, it has been reported that PITX2A protein can act as a negative regulator of FOXC1 through formation of PITX2A-FOXC1 heterocomplexes [35]. Therefore, the molecular interaction between these two proteins may affect expression of different target genes. It is also well documented that strict activity levels of these genes are required for normal development of the anterior segment of the eye [29, 46], and that genetic alterations resulting in variable functional disruption (e.g., hypo- and hypermorphic mutations, deletions and duplications of both PITX2 and FOXC1), underlie the phenotypic spectrum of anterior segment associated-diseases [30, 47, 48]. Therefore, activity and/or concentration variations of the individual proteins may influence the phenotypic outcome. The identification of patients in our cohort, as well as in other studies, who combine this type of variants in heterozygosis in two genes (PITX2/CYP1B1, PITX2/FOXC1 [19], FOXC1/CYP1B1 [49] and CYP1B1/GJA1 [50]) provide further evidence for this hypothesis. Digenic inheritance involving CYP1B1 and MYOC has been reported in early-onset congenital glaucoma [17] and PCG [16]. Likewise, in this study we have found two glaucoma patients who carried heterozygous rare PITX2 and CYP1B1 variants and showed very early glaucoma onset and severe disease, characterised by a large number of surgical operations required to control IOP. These data suggest a modifier role of rare PITX2 variants on age at onset and severity on CYP1B1 associated congenital glaucoma. Our data also revealed the presence of one PITX2 de novo mutation in one PCG patient, indicating that this type of genetic variants might also be involved in congenital glaucoma, adding additional genetic complexity to the genetic basis of this disease. We should keep in mind that reduced penetrance, low frequency of these variants and limited number of available patients due to low prevalence of the disease, will also complicate studies to determine the role of this type of genetic variation in CG. Combination of genome, transcriptome and even proteome analysis of individual patients, along with functional studies in animal models will be required to gain new insights into the genetic complexity of congenital glaucoma.

In summary, this study further supports the concept that mild functional alterations associated with variants in genes involved in ocular development may act as modifier factors in non-Mendelian CG cases, providing novel insights to understand the genetic complexity of this disease.

Supporting information

S1 Fig. Electropherograms of FOXC2, PITX2 and CYP1B1 variants identified in this study. Arrows over the peaks indicate the location of mutations. The mutant nucleotides are indicated in red and those in heterozygosis are shown below the corresponding wild type position. The sequences of the different variants were obtained from the corresponding probands in each family and the wild type sequences from control subjects.

(TIF)

S1 Table. The primer sequences and PCR conditions used for FOXC2 gene sequencing.

(DOCX)
Acknowledgments

We are indebted to the patients and their families and the "Asociación de Glaucoma para Afec-
tados y Familiares (AGAF)" for cooperation in this study. We thank Mrs. María-José Cabañero
for excellent technical assistance.

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