Inherited biallelic CSF3R mutations in severe congenital neutropenia

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Key Points

- Biallelic mutations in CSF3R must be considered as a novel genetic subtype in patients with congenital neutropenia.
- The p.Arg308Cys mutation in CSF3R leads to altered G-CSF receptor glycosylation and surface expression and abrogated downstream signaling.

Severe congenital neutropenia (SCN) is characterized by low numbers of peripheral neutrophil granulocytes and a predisposition to life-threatening bacterial infections. We describe a novel genetic SCN type in 2 unrelated families associated with recessively inherited loss-of-function mutations in CSF3R, encoding the granulocyte colony-stimulating factor (G-CSF) receptor. Family A, with 3 affected children, carried a homozygous missense mutation (NM_000760.3:c.922C>T, NP_000751.1:p.Arg308Cys), which resulted in perturbed N-glycosylation and aberrant localization to the cell surface. Family B, with 1 affected infant, carried compound heterozygous deletions provoking framshifts and premature stop codons (NM_000760.3:c.948_963del, NP_000751.1:p.Gly316fsTer322 and NM_000760.3:c.1245del, NP_000751.1:p.Gly415fsTer432). Despite peripheral SCN, all patients had morphologic evidence of full myeloid cell maturation in bone marrow. None of the patients responded to treatment with recombinant human G-CSF. Our study highlights the genetic and morphologic SCN variability and provides evidence both for functional importance and redundancy of G-CSF receptor-mediated signaling in human granulopoiesis. (Blood. 2014;123(24):3811-3817)

Introduction

Severe congenital neutropenia (SCN) is a heterogeneous disease, characterized by an absolute neutrophil count (ANC) <500 cells/μL and recurrent, life-threatening bacterial infections. Autosomal-dominant, autosomal-recessive, X-linked, and sporadic forms have been described, with mutations in several genes, including ELANE,1 GFI1,2 HAX1,3 G6PC3,4 VPS45,5 and WAS.6 The genetic cause in many SCN patients remains unidentified.7 The mainstay of treatment is recombinant human granulocyte colony-stimulating factor (rhG-CSF), which improves quality of life, increases the ANC in most patients, and decreases their infection frequency.8 Despite this, rhG-CSF–treated patients endure 8% and 21% cumulative incidences of sepsis and leukemic conversion, respectively, after 10 years of treatment.9

G-CSF induces proliferation, differentiation, and survival of myeloid progenitors. It signals through the cell-surface receptor G-CSF receptor (G-CSFR), a member of the type I cytokine receptors. The cytoplasmic region has no intrinsic kinase activity, but relies on Janus kinase activation for signal transduction through signal transducer and activator of transcription 3 and 5 (STAT3 and STAT5), as well as the Ras–mitogen-activated protein kinase and phosphatidylinositol 3-kinase–Akt pathways.10,11

CSF3R mutations were originally thought to be a cause of SCN.12 However, subsequent studies13,14 showed that CSF3R mutations found in previously reported SCN patients were somatic or de novo in meiosis rather than inherited from the parents of affected individuals. These noninherited, heterozygous CSF3R mutations typically affect the cytoplasmic region of G-CSFR, lead to a truncated variant, and are associated with a hyperproliferative phenotype in mutated cells and the leukemogenic events seen with SCN.15,16

Rarely, somatic or de novo heterozygous CSF3R mutations in the extracelluar region of the G-CSFR have been described.17,20 These
mutants are characterized by hyporesponsiveness to rhG-CSF and act in a dominant-negative fashion by interfering with proper function of the wild-type (WT) G-CSFR. Additionally, a heterozygous germ-line CSF3R mutation has been associated with neutrophilia.21

Here, we identify recessively inherited, loss-of-function CSF3R mutations in 4 affected children with SCN from 2 unrelated families, with morphologic evidence of full myeloid cell maturation in bone marrow, thus expanding the spectrum of monogenic disorders associated with SCN.

Materials and methods

Patient information and study approval

Patients were referred to the pediatric centers at Erciyes University (Kayseri), the Dr von Hauner Children’s Hospital at Ludwig Maximilian University (Munich), and the Maternal-Infant Hospital Vall d’Hebron (Barcelona).

Informed consent and assent were obtained according to current ethical and legal guidelines. Ethical review boards from involved institutions (Pediatric Centers at Erciyes University [Kayseri], the Dr von Hauner Children’s Hospital at Ludwig Maximilian University [Munich], and the Maternal-Infant Hospital Vall d’Hebron [Barcelona]) approved this study. This study was conducted in accordance with the Declaration of Helsinki.

Genetic analysis

DNA samples of the parents, I-1 and I-2, of family A were genotyped using the Affymetrix Genome-wide Human SNP array 6.0 (GE Platform GPL6801) following the manufacturer’s instructions (Affymetrix). Genomic DNA from patient P2 and her parents was used to construct exome libraries using the SureSelect XT Human All Exon V4 + UTRs kit (Agilent Technologies). Barcoded libraries were sequenced using SOLID 5500 XL next-generation sequencing platform (Life Technologies) to an average coverage depth of 80. Whole-exome sequencing (WES) identified between 110 000 and 120 000 candidate variants for each sample. Further protocol details and primer sequences used to amplify genomic DNA for Sanger sequencing are in the supplemental Methods (see supplemental Data available for details).

Structural analysis of WT G-CSFRWT and mutant G-CSFRArg308Cys proteins

To evaluate possible structural effects of the p.Arg308Cys substitution, we used Swiss PDB viewer22 version 4.01 to visualize and analyze Protein Data Bank (PDB) structure 2D9Q,23 which shows G-CSFR in complex with G-CSF, as described previously for IL21R.24

Cell culture

HeLa cells were maintained in Dulbecco’s Modified Eagle’s Medium (DMEM), supplemented with 10% heat-inactivated fetal bovine serum (FBS), 1% penicillin/streptomycin (Pen/Strep), and 2mM L-Glutamine at 37°C and 5% CO2. For starvation, DMEM containing 0.5% heat-inactivated (FBS), 1% penicillin/streptomycin (Pen/Strep), and 2mM L-Glutamine at 37°C, and incubated on ice for 60 minutes. Cells were centrifuged at 15 000 rpm for 15 minutes at 4°C. Proteins in the supernatant were electrophoretically separated and blotted. Primary antibodies used included rabbit anti-G-CSFR (H-176) (Santa Cruz Biotechnology), rabbit anti-phosphoSTAT3 (Tyr705) (Cell Signaling), rabbit anti-phosphoSTAT5 (Y694), and mouse anti-STAT3 (BD). Secondary antibodies included horseradish peroxidase (HRP)-conjugated anti-rabbit (Cell Signaling) and anti-mouse (BD) immunoglobulin G (IgG). Actin was visualized using HRP-conjugated anti-β-actin by Santa Cruz Biotechnology. For glycosylation studies, protein lysate was divided into 3 aliquots for the following conditions: nontreated, EndoH-treated, and PNGase F-treated. All buffers and enzymes were from New England Biolabs (Ipswich). Protein lysate for the samples not subject to treatment and samples intended for EndoH treatment were first incubated with denaturation buffer (5% sodium dodecyl sulfate [SDS], 0.4 M dithiothreitol) for 10 minutes at 95°C. Samples intended for PNGase F treatment were first incubated with 2.5% SDS for 10 minutes at 95°C. Samples were then treated with EndoH and PNGase F according to the manufacturer’s protocol. All protein lysates were boiled at 95°C for 5 minutes with sample buffer prior to SDS–polyacrylamide gel electrophoresis (PAGE) loading.

Western blot studies

HeLa cell pellets were resuspended in ice-cold 1× cell lysis buffer (Cell Signaling) supplemented with 0.06% protease inhibitor cocktail (Sigma-Aldrich) and 0.005% phenylmethylsulfonyl fluoride (Sigma-Aldrich) and incubated on ice for 60 minutes. Cells were centrifuged at 15 000 rpm for 15 minutes at 4°C. Proteins in the supernatant were electrophoretically separated and blotted. Primary antibodies included rabbit anti-G-CSFR (H-176) (Santa Cruz Biotechnology), rabbit anti-phosphoSTAT3 (Tyr705) (Cell Signaling), rabbit anti-phosphoSTAT5 (Y694), and mouse anti-STAT3 (BD). Secondary antibodies included horseradish peroxidase (HRP)-conjugated anti-rabbit (Cell Signaling) and anti-mouse (BD) immunoglobulin G (IgG). Actin was visualized using HRP-conjugated anti-β-actin by Santa Cruz Biotechnology. For glycosylation studies, protein lysate was divided into 3 aliquots for the following conditions: nontreated, EndoH-treated, and PNGase F-treated. All buffers and enzymes were from New England Biolabs (Ipswich). Protein lysate for the samples not subject to treatment and samples intended for EndoH treatment were first incubated with denaturation buffer (5% sodium dodecyl sulfate [SDS], 0.4 M dithiothreitol) for 10 minutes at 95°C. Samples intended for PNGase F treatment were first incubated with 2.5% SDS for 10 minutes at 95°C. Samples were then treated with EndoH and PNGase F according to the manufacturer’s protocol. All protein lysates were boiled at 95°C for 5 minutes with sample buffer prior to SDS–polyacrylamide gel electrophoresis (PAGE) loading.

Fluorescent microscopy for HeLa cells

HeLa cells were seeded onto glass coverslips, washed, and fixed with 3.7% paraformaldehyde for 20 minutes at room temperature. After washing 3 times with phosphate-buffered saline (PBS), cells were permeabilized by 5-minute incubation with 0.5% Nonidet P40 at room temperature. Cells were washed 3 times followed by incubation with 3% bovine serum albumin for 15 minutes at room temperature. Cells were incubated with mouse anti-Calcinein (1:100; BD) for 45 minutes at 37°C and then washed 3 times with PBS. Coverslips were then incubated with goat-anti-mouse Alexa Fluor 594 (1:150; Invitrogen) for 45 minutes at 37°C. Following 3 5-minute PBS washes, nuclei were stained with 4,6 diamidino-2-phenylindole (DAPI) for 1 minute at room temperature. Cells were washed again and mounted using fluorescent mounting medium (Dako).

For confocal imaging, we used a Fluoview FV1000 (Olympus) microscope with X30 digital camera (Olympus), fitted with a 60× objective. FV10-ASW acquisition software and ImageJ software25 were used for analysis.

Flow cytometry for HeLa cells

Forty-eight hours after transfection, cells were trypsinized, pelleted, and resuspended in ice-cold fluorescence-activated cell sorting (FACS) buffer (2% FBS, 0.1% NaN3 in PBS). Cells were pelleted and resuspended in blocking FACS buffer (10% FBS, 0.1% NaN3 in PBS) for 30 minutes on ice. Cells were washed with FACS buffer and 1 × 106 cells were aliquoted into 5-mL round-bottom Falcons (BD). Cells were stained in 100 μL of FACS buffer with 0.2 μg of phycoerythrin (PE)-conjugated anti-G-CSFR (BD) for 30 minutes on ice in the dark. PE-conjugated IgG1, κ isotype control (BD) was also used. After 3 5-minute washes with FACS buffer, cells were acquired and analyzed using FACScanto (BD) and FACSDiva software (BD). Data were analyzed with FlowJo software (FlowJo). Further details are given in the supplemental Methods.

G-CSF stimulation

To ensure equal transfection efficiency between each time point, cells were transfected in 10-cm dishes. After 24 hours, cells were trypsinized, counted, and equally reseeded in 35-mm dishes. At 5 days post-transfection, cells were trypsinized, pelleted, and resuspended in ice-cold PBS 3 times and harvested by scraping. Cells were washed with ice-cold PBS 3 times and harvested by scraping. Cells were counted, and aliquots were resuspended in ice-cold PBS containing 0.1% NaN3 and 10% FBS.
immediately washed and harvested. Protein lysate was prepared and analyzed by western blot.

Results

Clinical data

Family A consists of a healthy, consanguineous couple of Turkish descent with 3 SCN-affected children (Figure 1A). The index patient (P2) was diagnosed with SCN and dextrocardia at birth and required multiple hospitalizations for pneumonia, purulent otitis media, and fever of unknown origin. Her ANC to date has ranged between 420 and 2180 cells/μL. P1 was also diagnosed at age 2.5 years with SCN, but not dextrocardia. P3 was diagnosed at birth, but died at 3 months of age due to suspected aspiration pneumonia. Bone marrow examinations of P1 and P2 revealed full maturation of all 3 hematopoietic lineages without evidence of myelodysplasia (Figure 1B). P1 and P2 were treated with rhG-CSF (5 mg/kg per day), but did not show a response (Figure 1C). The father and mother of family A showed no signs of neutropenia, with ANCs of 5780 and 7080 cells/μL, respectively.

Index patient P4 of family B was a 9-month-old girl born to a healthy, nonconsanguineous couple of Spanish descent (supplemental Figure 1A). P4 was healthy until 2 months of age, when she was hospitalized for a urinary tract infection and treated with parenteral antibiotics. During this admission, P4 was found to be neutropenic (ANC to date: 200-1000 cells/μL). rhG-CSF was started at 5 μg/kg per day and progressively increased to a maximum dose of 40 μg/kg per day, with an ANC after this dose of 200 cells/μL (supplemental Figure 1B). Bone marrow aspirates revealed no maturation arrest and the presence of mature granulocytes. The father and mother of P4 showed no signs of neutropenia, with ANCs of 5780 and 7080 cells/μL, respectively. Supplemental Table 1 summarizes the clinical and genetic findings for both families.

Identification of CSF3R mutations

We performed homozygosity mapping and WES as described in the supplemental Methods for both parents and the index patient of family A. Homozygosity mapping identified 4 perfectly segregating runs of homozygosity (ROHs) with size above 1 Mb, which overlapped neither HAX1 nor G6PC3. These 4 regions map to chromosome 1 (33280648-40788961, build 37/hg19), chromosome 8 (19664010-2828292), chromosome 17 (13156053-14915602), and chromosome 18 (100949-2900803). WES detected 9 possible candidate variants, among which 2 variants were inside these ROHs (supplemental Results). We focused on a novel missense variant in CSF3R (NM_000760.3:c.922C>T, NP_000751.1:p.Arg308Cys; supplemental Figure 2B). Sanger sequencing of all family A members confirmed that the p.Arg308Cys CSF3R variant segregated with the disease phenotype in an autosomal-recessive inheritance pattern (Figure 2A). WES did not identify potential mutations in any SCN-causing genes (HAX1, G6PC3, ELANE, GFI1, VPS45, WAS) (supplemental Results, supplemental Table 2). Additionally, WES of P2 identified a homozygous variant in SPAG1 (NM_003114.4:c.1733del, NP_003105.2:p.578GlufsTer601), a causative gene for autosomal-recessive ciliary dyskinesia associated with dextrocardia, which was found in P2 (supplemental Table 3). Ciliary dyskinesia is associated with recurrent sinopulmonary infections.26 It is therefore possible that P2 suffers additionally from a ciliary disorder, which may contribute to her recurrent infections.

Sanger sequencing of ELANE, HAX1, G6PC3, GFI1, and SBDS in P4 did not detect mutations. Because of the patient’s hyporesponsiveness to rhG-CSF, CSF3R was sequenced. P4 showed a compound heterozygous mutation in CSF3R. In the first allele, a 16-bp deletion (NM_000760.3:c.948_963del) in exon 8 was found, whereas in the second allele, a 1-bp deletion (NM_000760.3:c.1245del) in exon 10 was found. Both deletions provoke frameshifts and premature stop codons (NP_000751.1:p.Gly316fsTer322 and NP_000751.1:p.Gly415fsTer432, respectively; supplemental Figure 2C). Sanger sequencing of CSF3R revealed that P4’s mother and father
were heterozygous for the mutations in exons 8 and 10, respectively (supplemental Figure 1C).

Structural implications of G-CSFR mutations

The G-CSFR p.Arg308Cys substitution is close to the conserved WSXWS motif at positions 318 to 322. As we recently discovered a similar mutation (p.Arg201Leu) preceding the WSXWS motif (positions 214-218) in the type I cytokine receptor IL21R,24 we searched for similarly located disease-causing mutations in other type I cytokine receptors. Interestingly, we found that IL7R, IL11RA, LEPR, IL12RB1, GHR, and MPL also have disease-causing mutations of an arginine shortly preceding the WSXWS motif that act recessively to cause the phenotype (supplemental Results, supplemental Table 4). In most cases, including G-CSFR, the mutation replaces the positively charged arginine with a neutral amino acid.

Bioinformatic analyses using the SIFT27 and PolyPhen28 algorithms gave scores of 0.02 and 0.998, 1.0 (Polyphen gives 2 scores), respectively, predicting that the p.Arg308Cys variant may be deleterious. Of the 19 possible amino acid replacements at position 308, the change to cysteine was most strongly predicted by SIFT to be deleterious. The SIFT analysis differs from the receptor family algorithms gave scores of 0.02 and 0.998, 1.0 (Polyphen gives 2 scores), respectively, predicting that the p.Arg308Cys variant may be deleterious. Of the 19 possible amino acid replacements at position 308, the change to cysteine was most strongly predicted by SIFT to be deleterious. The SIFT analysis differs from the receptor family predictions of an arginine shortly preceding the WSXWS motif that act recessively to cause the phenotype (supplemental Results, supplemental Table 4). In most cases, including G-CSFR, the mutation replaces the positively charged arginine with a neutral amino acid. Bioinformatic analyses using the SIFT27 and PolyPhen28 algorithms gave scores of 0.02 and 0.998, 1.0 (Polyphen gives 2 scores), respectively, predicting that the p.Arg308Cys variant may be deleterious. Of the 19 possible amino acid replacements at position 308, the change to cysteine was most strongly predicted by SIFT to be deleterious. The SIFT analysis differs from the receptor family predictions of an arginine shortly preceding the WSXWS motif that act recessively to cause the phenotype (supplemental Results, supplemental Table 4). In most cases, including G-CSFR, the mutation replaces the positively charged arginine with a neutral amino acid.

More insight can be gained from a study of the prolactin receptor PRLR,29 which is significantly similar to G-CSFR in sequence (BLAST E-value 6E-16, supplemental Table 4). Dagil et al29 (Figure 3) found that PRLR in the bound state (PDB structure 3D4830) has a very different structure from unbound PRLR (PDB structure 2LFG29). For example, Trp191 in PRLR, which corresponds to Trp318 in G-CSFR, has a very different relative position in the 2 PRLR structures. Residue Arg183 in PRLR corresponds to Arg308 in G-CSFR. In the unbound state, the Trp/Arg ladder does not form among the relevant residues. Therefore, we suggest that Arg308 is essential for the ladder to form and for G-CSFR to bind to its ligand G-CSF, as shown in 2D9Q, but Arg308 may be unimportant for G-CSFR to fold into its inactive state. The distinction between active and inactive states would explain how our antibody immunoblot could detect the mutant G-CSFRArg308Cys in the inactive state, even if G-CSFRArg308Cys cannot fold properly in the active state to receive the G-CSF signal.

The lower panel of Figure 2B shows the mutant protein with the Arg308Cys substitution. Although no preexisting hydrogen bonds appear to be broken by this substitution, an additional hydrogen bond is created. However, the p Arg/Arg ladder does not form among the relevant residues. Therefore, we suggest that Arg308 is essential for the ladder to form and for G-CSFR to bind to its ligand G-CSF, as shown in 2D9Q, but Arg308 may be unimportant for G-CSFR to fold into its inactive state. The distinction between active and inactive states would explain how our antibody immunoblot could detect the mutant G-CSFRArg308Cys in the inactive state, even if G-CSFRArg308Cys cannot fold properly in the active state to receive the G-CSF signal.

Aberrant N-glycosylation pattern of mutant G-CSFRArg308Cys

To further characterize the effect the p.Arg308Cys mutation on the G-CSFR protein, HeLa cells were transfected with plasmids encoding a fusion protein of either G-CSFR WT or G-CSFRArg308Cys fused to C-terminal eGFP. Western blot using an anti-G-CSFR antibody showed that the mutated G-CSFRArg308Cys had a decreased molecular weight in comparison with the WT (Figure 3A). This was also seen with eGFP-tagged human WT or mutant G-CSFRArg308Cys transduced primary myeloid hematopoietic progenitor cells isolated from Csf3r−/− and Csf3r−/− mice; mutant G-CSFRArg308Cys had a clearly reduced molecular weight as compared with WT (supplemental Methods, supplemental Figure 3).
Protein extracts from lysed HeLa cells transfected with eGFP-tagged human WT or mutant G-CSFRArg308Cys were treated with the recombinant glycosidases EndoH (that cleaves mainly high mannose N-glycans) and PNGase F (that removes high mannose, hybrid and complex N-glycans). Subsequent western blotting revealed that G-CSFRWT showed partial insensitivity to EndoH treatment, whereas the mutant G-CSFRArg308Cys was fully sensitive (Figure 3B), suggesting that the N-glycosylation patterns were distinct. In contrast, PNGase F treatment did not reveal differences in cleavage of N-glycans between the WT and mutant proteins.

**Mutant G-CSFR is retained in the ER and showed impaired expression on cell surface**

In type I cytokine receptors both glycosylation and the WSXWS motif appear to be critically involved in the intracellular trafficking of receptors to the cell surface.

To confirm this data in hematopoietic cells, we expressed the eGFP-tagged human WT or mutant G-CSFRArg308Cys in murine bone marrow BaF3 cells by plasmid transfection. In BaF3-cells, G-CSFRWT-eGFP showed speckled expression in the cytoplasm, presumably also at the cell surface, and colocalization with Golgi apparatus marker GM130 (supplemental Figure 4). However, the expression pattern of G-CSFRp.Arg308Cys was markedly different from WT and resembled more of the anti-Calnexin staining (supplemental Figure 5). Furthermore, we used murine primary myeloid Csf3r-knockout hematopoietic progenitor cells transduced by retroviral gene transfer. In murine primary hematopoietic progenitor cells, the human G-CSFRWT-eGFP was expressed in speckles all over the cytoplasm, presumably also at cell surface, and colocalized clearly to the Golgi apparatus, whereas the human G-CSFRp.Arg308Cys-eGFP mutant receptor showed clearly distinct localization pattern throughout the entire cytoplasm (supplemental Figure 6).

Furthermore, we stained nonpermeabilized HeLa cells expressing the above-mentioned eGFP-tagged constructs with a PE-conjugated anti-G-CSFR monoclonal antibody and analyzed with flow cytometry. Comparison of the eGFP signal (total G-CSFR) and PE signal (surface-bound G-CSFR) revealed a linear relationship for cells expressing the WT receptor, but not for the mutant, suggesting that the mutant G-CSFR lacks proper localization at the cell surface (Figure 3D).

The surface expression of G-CSFR (CD114) was studied in P4 of family B, who harbored both the p.Gly316fsTer322 and p.Gly415fsTer432 mutations. The whole peripheral blood samples of 1 healthy unrelated donor and P4 were stained with G-CSFR (CD114) and CD16 antibodies and detected by flow cytometry.
Peripheral blood cells were gated for high CD16 expression and side scatter characteristics, and the expression level of G-CSFR (CD114) was compared between samples. The result showed markedly decreased surface expression of G-CSFR (CD114) in P4 in comparison with a healthy donor (supplemental Figure 7).

**Abrogated signaling of mutant G-CSF receptor**

We then aimed to characterize the functionality of the mutant G-CSFRArg308Cys. Upon binding of its cognate ligand G-CSF, the G-CSF/G-CSFR complex induces phosphorylation of STAT3 on Tyr705 and STAT5 on Tyr694. HeLa cells transiently transfected with the previously mentioned plasmids were serum starved overnight and stimulated with rhG-CSF as described in supplemental Methods. Cell-derived protein lysates were subjected to SDS-PAGE and western blotting to evaluate STAT3 (Tyr705) and STAT5 (Tyr694) phosphorylation. Cells expressing the mutant receptor showed reduced phosphorylation of STAT3 and STAT5, but signal transduction was not completely abrogated (Figure 3E).

**Discussion**

We herein describe a novel genetic subtype of SCN characterized by recessively inherited loss-of-function CSF3R mutations, full myeloid cell maturation in the bone marrow, and refractoriness to in vivo rhG-CSF treatment. This is in contrast to inherited SCN with mutations in ELANE or HAX1, which show a characteristic myeloid maturation arrest at the promyelocyte-to-myelocyte differentiation step and generally respond to rhG-CSF treatment. Although all patients in this study displayed peripheral neutropenia, full maturation of neutrophil granulocytes in the bone marrow was seen in all 3 patients who underwent a bone marrow examination (P1, P2, P4). G-CSF is a critical regulator of granulopoiesis, as shown in several human and murine studies. Consistent with our patients, both Csf3r– and G-CSF–deficient mice showed reduced numbers of neutrophils in peripheral blood. Moreover, no maturation arrest was seen in the bone marrow of the Csf3r– and G-CSF–deficient mice. Our results support the previous findings observed in genetically engineered mice and suggest the presence of G-CSFR–independent signaling pathways controlling granulopoiesis.

Interestingly, although both patients from family A and B were refractory to in vivo rhG-CSF treatment, in vitro assays with the G-CSFRArg308Cys mutation from family A showed residual response to rhG-CSF stimulation. This may be explained by a small cell population able to express receptor on the surface (Figure 3D, supplemental Figure 7). Alternatively, the residual function seen in vitro may be due to heterotopic surface expression of the mutant G-CSFRArg308Cys in HeLa cells.

We also highlight the relevance of the residues interfacing with the WSXWS motif of G-CSFR for proper glycosylation and folding, as well as cell membrane expression. Mutations in or preceding the WSXWS motif in other type I cytokine receptors and cause disease (supplemental Table 4) suggests that the effect in G-CSFR is not Cys-specific.

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**Authorship**

Contribution: A.T., P.M.J., D.M., D.K., N.K., T.R., J.P., D.P., E.M.G., A.A.S., and C.K. designed and performed experiments and analyzed data for family A; and J.I.A., J.L.D.D., C.D.d.H.R., J.S.d.T.C., and J.Y. provided clinical care, designed and performed experiments, and analyzed data for family B; I.P.T. provided bones of Csf3r-/- and Csf3r+/- mice; E.U., M. A.O., T.P., and M.K. provided clinical care and performed experiments for family A; and
E.M.G. and A.A.S. did bioinformatics analysis and the literature search for mutations in other type I cytokine receptors.

Conflict-of-interest disclosure: The authors declare no competing financial interests.

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