Cyba-deficient mice display an increase in hematopoietic stem cells and an overproduction of immunoglobulins

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ABSTRACT

The regulation of protein function by reversible oxidation is increasingly recognized as a key mechanism for the control of cellular signaling, modulating crucial biological processes such as cell differentiation. In this scenario, NADPH oxidases must occupy a prominent position. Our results show that hematopoietic stem and progenitor cells express three p22phox-dependent NADPH oxidase members (NOX1, NOX2 and NOX4). By deleting the p22phox coding gene (Cyba), here we have analyzed the importance of this family of enzymes during in vivo hematopoiesis. Cyba−/− mice show a myeloid bias, and an enrichment of hematopoietic stem cell populations. By means of hematopoietic transplant experiments we have also tried to dissect the specific role of the NADPH oxidases. While the absence of NOX1 or NOX2 provides a higher level of reconstitution, a lack of NOX4 rendered the opposite result, suggesting a functional specificity among the different NADPH oxidases. Cyba−/− cells showed a hampered activation of AKT1 and a sharp decrease in STAT5 protein. This is in line with the diminished response to IL-7 shown by our results, which could explain the overproduction of immunoglobulins observed in Cyba−/− mice.

Introduction

From being considered harmful metabolic by-products, reactive oxygen species (ROS) have turned out to be important regulators of cellular biology, by acting as bona fide secondary messengers.1–3 NADPH oxidases are the only cellular system specialized in the production of ROS.4 The founding member of this family is phagocyte oxidase, a multiprotein complex that produces huge amounts of superoxide during respiratory burst, which is required for the elimination of pathogens.4 The complex consists of two integral membrane proteins (p22phox, and the catalytic subunit, named gp91phox or NOX2), three cytosolic subunits (p40phox, p47phox, p67phox), and the Rac GTPase.4 Inactivating mutations affecting the complex produce chronic granulomatous disease (CGD), a hematological disorder characterized by the occurrence of repetitive infections.5

For quite some time the NADPH oxidase from phagocytes seemed to be the only oxidase. However, in 1999 cloning and characterization of NOX1 was reported,7 which was followed by the discovery of other homologues oxidases. Nowadays the family comprises seven members, which can be classified into two groups, those dependent on p22phox (NOX1, NOX2, NOX3, NOX4), and those p22phox-independent that can be activated by calcium (NOX5, DUOX1 and DUOX2).6 These enzymes are present in all eukaryotic cells, including unicellular organisms,7 and several members of the family are commonly expressed simultaneously.4 This broad distribution and its regulation by extracellular signaling make NADPH oxidases a key element in redox signaling. ROS and NADPH oxidases are involved in the control of cell fate.2,9–11
Hematopoiesis is a paradigmatic example of cell differentiation, because hematopoietic stem cells (HSC) must produce all mature blood lineages. There is increasing evidence suggesting the importance of redox signaling for hematopoietic differentiation, as well as for the contribution of an elevated level of ROS in the development of leukemia.10,11 Our previous data has shown the requirement of NADPH oxidase-produced ROS for in vitro megakaryocytic differentiation.12 However, even though NADPH oxidases were discovered in the hematopoietic system, knowledge of the importance of these enzymes for hematopoiesis in vivo is scarce. This analysis has surely been hampered by the fact that hematopoietic cells express several NADPH oxidase isoforms (our unpublished data and13).

Here we have analyzed the relevance of NADPH oxidases during in vivo hematopoiesis. Mouse hematopoietic progenitor cells express NOX1, NOX2 and NOX4, all of them p22phox-dependent. In order to assess the importance of NADPH oxidase activity for in vivo hematopoiesis we have generated p22phox-deficient mice (Cyba−/−). The lack of p22phox induces a myeloid bias. Moreover, there is an increased proportion of HSC in the bone marrow (BM). In competitive transplant experiments, cells deficient in NOX1 (Nox1−/−), NOX2 (Cybb−/−) and p22hox (Cyba−/−) showed an increased reconstitution ability with respect to control cells. However, NOX4 (Nox4−/−) deficient cells did not show this effect. The response of Cyba−/− cells to IL-7 is severely impaired, which can be explained by a hampered activation of AKT1 and by the reduction in STAT5 protein levels. These redox signaling alterations could be the cause of the increased production of immunoglobulins observed in Cyba−/− mice.

**Methods**

**Animals**

C57BL/6 mice were from the University of Salamanca Animal Facility Unit. Albino C57BL/6 mice (B6.Cg-Tyr+129), Nox1 (B6.129X1- Nox1tm1Kkr/J), Nox2 (B6.129S2- Cybbtm1Din/J) and Nox4 (B6.129- Nos4tm1Kkr/J) deficient mice were from Jackson Laboratory (Bar Harbor, ME, USA). Cyba−/− mice were generated as detailed in the Online Supplementary Materials and Methods. All procedures were approved by the Bioethics Committee at the University of Salamanca.

**ROS detection**

Peripheral blood (PB) was lysed to eliminate red blood cells. ROS levels were detected by cell staining with 10 μM DCFDA (2',7-dichlorodihydrofluorescein diacetate)12 or with 100 μM luminol after stimulation with 2 μM PMA (phorbol-12-myristate-13-acetate).

**Colony forming unit (CFU) assays**

Cells were grown in methylcellulose semisolid medium supplemented with a cocktail of cytokines (50 ng/ml SCF, 20 ng/ml IL-3, 20 ng/ml GM-CSF and 3 U/ml EPO), or with individual cytokines (IL-3, GM-CSF, and IL-7) at 20 ng/ml. 200,000 spleen and 10,000 BM cells (200,000 with IL-7) were used. Cells were grown at 37°C and 5% CO2 for 12 days.

**BM transplantation**

C57BL/6 recipient mice were lethally irradiated with two doses of 5 Gy from a Cs source (Gammarcell 1,000 Elite, Nucleus Facility, University of Salamanca) as previously described.15 Recipient mice were injected intravenously through the lateral tail vein with 3x106 BM cells from ES Cyba−/− or from wild-type C57BL/6 mice. Competitive transplant experiments were performed by injecting 1.5x106 BM cells from NADPH oxidase deficient mice (Nox1−/−, Cybb−/−, Nox4−/− or ES Cyba−/−) or from C57BL/6 control mice, together with 1.5x106 BM competitor cells from wild-type C57BL/6 mice. Cell origin was assessed by the differential expression of the CD45.1 and CD45.2 isotypes. Secondary transplants were carried out 18 weeks post-first transplant, by injecting 3x106 BM cells from the primary transplanted mice into lethally irradiated C57BL/6 mice.

**In vivo bromodeoxyuridine incorporation assay**

Mice were intraperitoneally treated with 100 mg/kg 5-bromo-2’-deoxyuridine (BrdU) and sacrificed 14 hours post-injection. The percentage of BrdU proliferating cells was analyzed by flow cytometry as previously described.16

**RNA sequencing analysis**

cDNA libraries were compiled using the Illumina TruSeq RNA Library Preparation Kit v2, with 2 μg of total RNA from immunopurified Lin− cells.17 Single-end 150 nt length sequencing was performed on an Illumina NextSeq 500 System using a Mid Output kit v2.5 from Illumina. Reads obtained were compared against the GRCm38 mouse genome using the bioinformatic facility tool of the University of Salamanca (https://ranaseq.eu/home). GEO accession: GSE131725.

**Immunoblotting**

Immunoblotting and quantification of bands were performed as previously described,18 and using fluorescently labelled secondary antibodies with an Odyssey Infrared Imaging System (Li-Cor). VINCULIN was used as a loading control.

**Statistical analyses and data report**

Data are expressed as mean values ± standard deviation. In dot graphs, each dot represents an individual value and the horizontal line denotes the mean value. Data were analyzed with SPSS 23 software. Two-tailed unpaired Student’s t test was used, and differences were considered statistically significant when P<0.05 (*), P<0.01 (**), and P<0.001 (**).
Figure 1. The lack of Cyba gene induces a myeloid bias. (A) NADPH oxidase expression in Lin progenitor cells analyzed by RT-PCR. The tissues indicated in the figure were used as positive controls (n=2). (B) Representative blots of p22phox protein expression in wild-type, ES Cyba-/− and CR Cyba-/− mice (n=4). (C) Peripheral blood cells were stimulated with 2 μM phorbol 12-myristate 13-acetate (PMA) for 30 min. The level of reactive oxygen species (ROS) was analyzed in the granulocyte population. A representative flow cytometry experiment and the mean ± standard deviation (SD) of the ROS fold-increase of seven different experiments are shown. (D) 200,000 peripheral blood (PB) cells were incubated with 100 μM luminol, and chemiluminescence was monitored for 15 min at 37°C after addition of 2 μM PMA. A representative experiment showing the RLU (relative luminescence units) and the mean ± SD of the chemiluminescence fold-increase at 10 min of three different experiments are shown. (E) Analysis of hematopoietic lineages in ES Cyba-/− and control mice. (F) Analysis of hematopoietic lineages in CR Cyba−/− and control mice.
Figure 2. Bone marrow cells from Cyba<sup>-/-</sup> mice are enriched in hematopoietic stem cells. (A) Representative flow cytometry analysis of the hematopoietic stem cells (HSC) populations in CR Cyba<sup>-/-</sup> and control mice. (B) Percentages of the different stem and progenitor populations in CR Cyba<sup>-/-</sup> and control mice. (C) Percentages of the different stem and progenitor populations in ES Cyba<sup>-/-</sup> and control mice. (D) Analysis of stem and progenitor cell populations at 20 weeks post-transplantation in lethally irradiated mice transplanted with wild-type or ES Cyba<sup>-/-</sup> cells. LK: lineage-committed progenitors; LSK: Lin-<sup>Sca-1</sup>-<sup>c-Kit</sup> hematopoietic stem cells; LMPP: lymphoid-primed multipotent progenitors; LT-HSC: long-term HSC; ST-HSC: short-term HSC; GMP: granulocyte-monocyte progenitors; MEP: megakaryocyte-erythrocyte progenitors; CMP: common myeloid progenitors, CLP: common lymphoid progenitors.
is suggested to be due to the requirement of NOX3 activity for otoconia synthesis.18

The lack of NADPH oxidase activity in Cyba−/− mice was checked by measuring the levels of extracellular and intracellular ROS after stimulating NADPH oxidase with PMA (Figure 1C-D and Online Supplementary Figure S3).

ES Cyba−/− mice showed a significant increase in CD11b+ myeloid cells in PB and spleen. A general increase of CD11b+ myeloid cells is also observed in the BM, though not statistically significant (Figure 1E). These results support the existence of a myeloid bias in ES Cyba−/− mice, a notion that is further supported by analysis of CR Cyba−/− mice, which shows a significant increase in myeloid cells in BM (CD11b+Gr1− cells) and in spleen (CD11b+Gr1− cells) (Figure 1F).

Cyba−/− mice show an enrichment of hematopoietic stem progenitor cells in the BM

In the BM, no differences in cellularity or the percentage of Lin− progenitor cells were observed between knockout (ES Cyba−/− or CR Cyba−/−) and wild-type mice (data not shown). Within the Lin− cell population we observed an enrichment in HSC (multipotent long-term HSC [LT-HSC] and short-term HSC [ST-HSC]) in both CR Cyba−/− (Figure 2A-B) and ES Cyba−/− mice (Figure 2C). This increase was around two-fold for both LT- and ST-HSC. No significant differences were found in more-committed myeloid progenitor cells (megakaryocyte-erythrocyte progenitor [MEP], granulocyte-monocyte progenitor [GMP] and common myeloid progenitor [CMP] cells) (Figure 2A-C).

In order to test whether the enrichment of HSC in Cyba−/− mice is cell-autonomous or determined by the niche, BM cells from ES Cyba−/− and wild-type mice were transplanted into lethally irradiated wild-type mice. There was an enrichment of hematopoietic stem progenitor cells (HSPC) populations in animals transplanted with ES Cyba−/− BM cells (LT-HSC, ST-HSC, lymphoid-primed multipotent progenitor [LMPP], MEP and common lymphoid progenitor [CLP] cells) (Figure 2D). These results strongly suggest that the higher proportion of HSC in the Cyba−/− mice does not depend on the influence of the niche, instead it is a cell-autonomous phenomenon.

Cyba−/− HSC show an increased proliferation capacity

In vivo BrdU incorporation experiments were performed

![Figure 3. Bone marrow progenitor cells from Cyba−/− mice proliferate more than control cells.](image)

CR Cyba−/− and control mice were injected intraperitoneally (IP) with 100 mg/kg 5-bromo-2'-deoxyuridine (BrdU), and sacrificed 14 hours later. BrdU incorporation in the hematopoietic stem cells (HSC) bone marrow (BM) populations was analyzed by flow cytometry. (A) A representative experiment of the BrdU incorporation in hematopoietic stem cells (LSK cells) is shown. (B) Percentage of BrdU+ cells in the different HSC populations. LMPP: lymphoid-primed multipotent progenitors; LT-HSC: long-term HSC; ST-HSC: short-term HSC.

![Figure 4. Colony forming assays with bone marrow and spleen cells from Cyba−/− mice.](image)

The colony forming unit (CFU) ability of bone marrow (BM) and spleen cells in response to complete medium or individual cytokines was tested. (A) CFU ability of BM and spleen cells in response to complete medium (n=6 and n=9, respectively). (B) BM CFU in the presence of IL-3, GM-CSF or IL-7 (n=6). The mean ± standard deviation (SD) of the number of colonies normalized with respect to the control is shown.
to analyze the proliferation capacity of HSPC from CR Cyba° and control mice. Our results showed a significant increase in BrdU incorporation in Lin−Sca-1−c-Kit+ hematopoietic stem cells (LSK cells), mainly in ST-HSC and LMPP, but also a general increase in LT-HSC (Figure 3). This higher proliferation potential would explain the enrichment of HSC in Cyba° mice.

**Cyba° cells show a lower clonogenic capacity in response to IL-7**

We next analyzed the self-renewal potential of BM and spleen cells in response to different cytokines by CFU assays. In the presence of complete media (SCF, IL-3, GM-CSF and EPO) BM CR Cyba° cells produced a slightly higher number of colonies, though not statistically significant, while spleen CR Cyba° cells produced a significantly higher number (Figure 4A). BM cells from CR Cyba° mice showed a general trend towards a higher clonogenic capacity in the presence of cytokines that induce myeloid differentiation (IL-3 and GM-CSF), while there was a significant decrease in the number of colonies in the presence of IL-7, a cytokine that drives B-cell differentiation in mice19 (Figure 4B). These results are consistent with the myeloid bias observed in vivo.

**Nox1, Cybb and Cyba deficient cells show an increased hematopoietic reconstitution capacity**

The results so far support the relevance of NADPH oxidase activity for hematopoiesis. Therefore, an interesting angle would be to delineate the importance of the different NADPH oxidase enzymes. Thus, through competitive transplantation experiments we analyzed the

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**Figure 5. The deficiency of NOX1 and NOX2 increases hematopoietic reconstitution capacity in competitive transplant experiments.** The percentage of chimerism throughout time in peripheral blood (PB) for the different primary and secondary transplants is shown: (A) Primary and (B) secondary transplants for Nox1° and control mice (n=11 and n=6, respectively), (C) Primary and (D) secondary transplants for Cybb° and control mice (n=11 and n=6, respectively), (E) Primary and (F) secondary transplants for Nox4° and control mice (n=10 and n=6, respectively). Graphics show mean ± standard deviation (SD).
hematopoietic reconstitution ability of cells lacking NOX1 (Nox1-/-), NOX2 (Cybb-/-) and NOX4 (Nox4-/-). BM cells from these mice strains were challenged with the same amount of BM cells from wild-type mice, and were transplanted into lethally irradiated recipient wild-type mice (Figure 5).

The lack of NOX1 (Nox1-/-) and NOX2 (Cybb-/-) conferred a mild but significant increase in reconstitution compared to wild-type cells in primary recipients (Figure 5A, C). Interestingly, Nox4-/- cells showed the opposite, with a slightly lower reconstitution capacity in primary transplants (Figure 5E). In order to test this issue further, secondary transplants were performed in the three settings. While Nox1-/- and Cybb-/- once again showed a higher reconstitution capacity than wild-type cells (Figure 5B, D), Nox4-/- displayed the same percentage of chimerism (50%) as in the primary transplant (Figure 5F). These results validate those obtained in the primary transplants and suggest a differential role for each Nox isoform during in vivo hematopoiesis.

In agreement with the results described for Nox1-/- and Cybb-/-, competitive transplant experiments performed with ES Cyba-/- mice and wild-type littermates offered a very subtle enhancement of reconstitution of Cyba-/- cells (Figure 6A-B), which could also be observed in secondary transplants (Online Supplementary Figures 4A-B).

When we analyzed the lineages within the ES Cyba-/- transplanted cells, there was a significant increase in CD11b+ Gr1+ and CD3+ cells, and a decrease in CD19+ and B220+ cells (Figure 6C). These features were also evidenced in secondary transplants (Online Supplementary Figure S4C). Moreover, there was also an enrichment of LT-HSC and lineage-committed progenitors (LK) and a general increase in LMPP and LSK in the ES Cyba-/- transplanted cells (Figure 6D). These features are much like those found in the ES Cyba-/- mice (Figure 1E and Figure 2C), strongly suggesting once again that the hematopoietic alteration we have found in Cyba-/- mice is the result of cell-endogenous causes and not due to the influence of the niche.

Unlike Cyba-/- cells, in the transplant experiments performed with Nox1-/- and Cybb-/- deficient cells, no clear differences were found in regard to the different hematopoietic lineages (Online Supplementary Figure S5A). No signifi-
Figure 7. Transcriptome analysis of Lin- cells from Cyba-/- and control mice. Lin cells were immunopurified from the bone marrow (BM) of two CR Cyba-/- and two control mice. The gene expression profile was analyzed by RNA sequencing. (A) Heat-map showing Z-scores of the expression values (normalized as transcript per million [TPM] values) of genes with significantly altered expression in Cyba-/- cells. (B) Network of functional analysis results for processes. Each node represents a functional category. The percentage of significantly differentially-expressed genes annotated by the different categories on the GO database is represented by colors, as shown by the scale in the figure. Connecting lines show the relationships between different functional categories; the width of the lines is proportional to the genes in common. (C) Network of functional analysis results for pathways.
Figure 8. p22phox deficiency increases myeloid gene expression, immunoglobulin serum levels and induces the downregulation of STAT5 protein. (A) The expression of several myeloid genes that were upregulated according to RNA sequencing data was analyzed by quantitative RT-PCR (qRT-PCR) in Lin- cells. Results show the relative expression of these genes in CR Cyba-/- mice with respect to control mice. The discontinuous line represents expression in control mice. (B) Relative expression of Cybb in Lin- cells from CR Cyba-/- mice with respect to control mice analyzed by qRT-PCR. (C) Immunoglobulin protein levels in the bone marrow (BM) and spleen of CR Cyba-/- and control mice (n=4). (D) Immunoglobulin serum levels in CR Cyba-/- and control mice. Levels of IgA, IgG and IgM in serum were analyzed by ELISA. Results are shown as mg/dL. The steady state level in BM and spleen of the following proteins was analyzed: (E) STAT5 and CRKL (n=4), (F) STAT3 (n=3). (G) Stat5a and Stat5b expression was analyzed by qRT-PCR in Lin- cells. Results show the relative expression of these genes in CR Cyba-/- mice with respect to control mice. (H) MYC protein levels in BM and spleen from CR Cyba-/- and control mice (n=3).
cant differences were found regarding HSPC, though Cyba<sup>-/-</sup> cells showed a general increase in the LT-HSC and ST-HSC populations (Online Supplementary Figure S5B).

Interestingly, lineage analysis of Nox<sup>-/-</sup> cells offered some similarities to the differences described in the transplants performed with ES Cyba<sup>-/-</sup> mice, such as, for example, a significant increase in CD11b<sup>+</sup> Gr1<sup>-</sup> myeloid cells in BM, and the same changes in CD3<sup>+</sup> and CD19<sup>+</sup> cells (Online Supplementary Figure S5C).

Cyba deficiency increases the expression of immunoglobulin genes

In order to gain some insight regarding the previous results we performed transcriptome profile analyses in Lin<sup>-</sup> cells from control and CR Cyba<sup>-/-</sup> mice by RNA sequencing. We found 47 upregulated and five downregulated genes in Cyba<sup>-/-</sup> cells (Figure 7A). In line with the myeloid bias described above, among the upregulated genes there were several genes encoding proteins highly expressed in myeloid cells, such as Il6st (CD11b), Pirb (CD68A), Lilrβ4 (CD85K), Ccrf1 (CD191) and Lrg1. Intriguingly, there was also an upregulation of Cybb mRNA expression in the Cyba<sup>-/-</sup> cells. These results were corroborated by quantitative RT-PCR (qRT-PCR) (Figure 8A-B).

However, the most striking result was the upregulation of 18 immunoglobulin genes of both heavy variable (IgVh) and κ light variable (IgKv) chains. In line with this, we detected an increased expression of immunoglobulins in the BM and spleen of CR Cyba<sup>-/-</sup> mice (Figure 8C), suggesting an increased production of immunoglobulins by CR Cyba<sup>-/-</sup> mice. Confirming this hypothesis, CR Cyba<sup>-/-</sup> mice showed a significant increase in Igα, in IgG, in serum, and a general increase in IgM that did not reach statistical significance (Figure 8D). All in all, these results are in agreement with an exacerbated production of immunoglobulins by CR Cyba<sup>-/-</sup> mice.

Functional analyses of these results (Figure 7B) suggested that the innate immune response and immunoglobulin production processes are severely affected by the lack of Cyba, with a high proportion of genes involved in these processes among the miss-regulated genes in Cyba<sup>-/-</sup> mice. Accordingly, the response to bacteria and defense against viruses were among the altered processes. Moreover, as expected, processes related to superoxide anion generation are also hampered. Of note, other fundamental processes such as cell migration and cell adhesion also seem to be affected by the lack of Cyba (Figure 7B). In the same line, pathway analyses also suggest the alteration of the innate immune response, and pathways related with cell adhesion and migration (Figure 7C). Signaling pathways such as VEGF, Rho GTPases and PI3K-AKT-mTOR appear among the potentially altered pathways (Figure 7C). In support of this, we observed that activation of AKT1 in response to several hematopoietic cytokines was somewhat impaired in Cyba<sup>-/-</sup> cells (Online Supplementary Figure S6A).

Cyba deficiency leads to the downregulation of STAT5

Given the possible implication of NADPH oxidases in the regulation of the signaling pathways governing hematopoiesis<sup>22-24</sup> we analyzed the activation of several signaling pathways by immunoblotting. No differences were found in the activation of ERK or in levels of β-CATENIN (Online Supplementary Figures 6B-C). However, our analyses revealed a sharp decrease in STAT5 protein in BM and spleen Cyba<sup>-/-</sup> cells (Figure 8E and Online Supplementary Figure 6D). In contrast, no differences were found regarding the expression of STAT3 protein (Figure 8F), which highlights the specificity of the decrease in STAT5 in Cyba<sup>-/-</sup> mice. No differences in Stat5a and Stat5b mRNA between Cyba<sup>-/-</sup> and control mice were detected in the RNA-seq analyses (Figure 7), which was corroborated by qRT-PCR (Figure 8G). Therefore, this strongly suggests that the decrease in STAT5 in Cyba<sup>-/-</sup> mice occurs at the protein level. Moreover, the levels of CRKL, a protein that can form functional heterodimers with STAT5<sup>22,23</sup> were also downregulated in Cyba<sup>-/-</sup> mice (Figure 8E). Finally, the levels of MYC, another transcription factor important for hematopoiesis<sup>21</sup> were also upregulated in Cyba<sup>-/-</sup> mice (Figure 8H).

Discussion

There is accumulating evidence supporting the importance of ROS and redox signaling in hematopoietic differentiation. NADPH oxidases would be ideal candidates as an adjustable source of ROS during hematopoiesis, given that they can be activated by extracellular signals, including hematopoietic cytokines.<sup>22-25</sup> Our previous results support this working hypothesis, since we have shown the relevance of ROS production by NADPH oxidases for regulating in vitro megakaryocytic differentiation.<sup>12</sup>

Therefore, we questioned whether NADPH oxidase activity would be required for in vivo hematopoiesis. The difficulty of this study lies in the fact that many cells express several NADPH oxidases simultaneously. Mouse hematopoietic progenitors express three different p27<sup>moes</sup>-dependent NOX (NOX1, NOX2 and NOX4), therefore, we started by generating Cyba<sup>-/-</sup> mice.

Flow cytometry analyses supported the existence of a myeloid bias in ES Cyba<sup>-/-</sup> mice, which could also be observed in the CR model, though at a weaker level. Nevertheless, we found the upregulation of myeloid genes in Lin<sup>-</sup> cells from CR Cyba<sup>-/-</sup> mice, also supporting the existence of a myeloid bias in the CR Cyba<sup>-/-</sup> mice. Although the genetic background of the ES and CR models (C57BL/6N and C57BL/6J respectively) are very closely related,<sup>26</sup> we cannot rule out that this might account for the differences regarding the myeloid bias. Moreover, the weaker phenotype in the CR Cyba<sup>-/-</sup> mice could be in line with phenomena associated with the CRISPR/Cas9 tool, such as the activation of genetic compensation mechanisms, and the residual expression of the targeted protein.<sup>28</sup>

A remarkable increase in HSC populations is observed in Cyba<sup>-/-</sup> mice, which correlates with a higher cellular proliferation capacity, as shown by the in vivo BrdU incorporation assays. It has been suggested before that MYC upregulation can lead to the expansion of HSPC<sup>29</sup> so the increased expression of MYC in BM Cyba<sup>-/-</sup> mice could explain this effect in our cells. Thrombopoietin (TPO) has recently been identified as a crucial cytokine for the control of HSC quiescence.<sup>30</sup> TPO signaling depends on the activation of STAT5<sup>30</sup> in turn, the importance of STAT5 for maintaining HSC quiescence has also been reported.<sup>31</sup> Considering the dramatic decrease in STAT5 protein levels in Cyba<sup>-/-</sup> mice, hampered TPO signaling as a consequence of diminished STAT5 activation could shift the balance
from quiescence towards an increase in cell proliferation. In line with this hypothesis NOX2 and NOX4 ROS production seems to be required for maintaining the stemness of induced pluripotent stem cells.32

We have addressed the relevance of NOX isoforms for hematopoiesis through competitive transplantation experiments. Cells lacking NOXI (Nox1+) and NOX2 (Cybb−) had a higher level of hematopoietic reconstitution with respect to wild-type cells. The hematopoietic reconstitution ability of Cybb− cells has been reported through competitive transplant experiments, showing an enhanced hematopoietic reconstitution of adult BM Cybb− cells at the beginning of the experiment, which is lost at later time points.31 Our results would be similar to those reported by Weisser et al.32 since at short time points we also found a slight reconstitution advantage in Cybb− cells, though in our case said advantage is kept throughout time in the primary transplant. Moreover, we performed secondary transplants that confirmed the greater reconstitution ability of not only Cybb− cells, but also of Nox1+ and Cyba− cells. These results support those obtained by us in primary transplants. While Weisser et al.32 used a limited number of purified HSC that were modified by lentivirus infection before the transplant, we used 3x10^6 whole BM cells for the transplant. We wondered whether the different progenitor cell numbers or the presence of more mature progenitors in our assays could be responsible for the sustained advantage of reconstitution observed. Another interesting area may be the upregulation of IL-1β signaling caused by the hyperinflammation linked to the CGD. According to the data presented by Weisser et al.33 IL-1β signaling can lead to the impaired reconstitution capacity of the HSC. The group comments in their methods section that animals with overt infections were not included in their study, which suggests they may have used animals with some kind of inflammatory pathology. In our case, none of the animals used showed any sign of inflammation or disease, so it is likely that a different level of IL-1 signaling could explain the differences in our results. While preparing our manuscript, another article was published reporting a lower reconstitution ability for Cybb− cells in primary transplants.34 Adane et al.35 also used whole BM for their assays, but again a more limited cell number was used, and unlike us, their findings were not confirmed by secondary transplants.

Together these results show a certain variability in the outcome of these assays, which is also supported by the fact that Weisser et al.35 did not find any difference in the transplants when using Cybb− fetal liver cells.33 Despite these differences, Weisser et al.35 report that Cybb− mice show an increased number of HPC, with a higher proliferation rate. They also show an increase in the formation of myeloid CFU by Cybb− BM cells. All these results are in line with our findings for Cyba− mice, and combined would support the importance of NOX2 and p22phox for the control of HSC function.36

Nox4− transplanted cells showed certain lineage alterations, similar to those observed in the Cyba− mice. All this considered, we hypothesize that NOX1 and NOX2 could be involved in the regulation of HSC homeostasis, while NOX4 might be important for lineage decisions given its implication in cell differentiation.7

The lack of phagocyte oxidase activity in Cyba− mice would lead to a defective immune response, as suggested by analysis of our RNA-seq results. Moreover, alterations of the NOD-like receptor signaling pathway, involved in the recognition of pathogen derived patterns,36 or the alteration of antigen processing and presentation pathways46 suggests that adaptive immunity could also be compromised in Cyba− mice. Additionally, we detected an exacerbated production of immunoglobulins in Cyba− mice, in agreement with the high level of IgG in CGD human subjects.37 An interesting question is the molecular mechanism leading to this scenario. IL-7 signaling drives B cell lymphopoiesis in mice.38 STAT5 and the PI3K/AKT pathway are the downstream effectors of IL-7 during B cell development.39 The lack of Stat5a and Stat5b blocks B cell development.40 STAT5 is not only required for maintaining cell survival during B-cell differentiation,41 but also for regulating Ig rearrangement,42 so much so that IL-7 signaling prevents premature Igκ rearrangement through STAT5 activation.43 Moreover, there are studies suggesting that STAT5b deficiency in humans leads to an increased production of immunoglobulins E32 and G.44 Therefore, we hypothesize that STAT5 downregulation and the reduced response to IL-7 must be key factors for the upregulation of immunoglobulins observed in Cyba− mice.

Analyses of STAT5 protein stability in the literature are rather scarce; STAT5 can be degraded by calpain44 and caspase-3.45 Moreover, it has also been shown that the redox sensitive phosphatase DUSP4 can trigger STAT5 protein degradation through proteasome and lysosome pathways.46 In line with that, it could be hypothesized that a lower level of ROS in Cyba− mice may allow a higher level of DUSP4 activity, thus leading to STAT5 protein degradation.

In summary, by downregulating NADPH oxidase activity in hematopoietic cells through the deletion of the Cyba gene, we show the relevance of such activity for in vivo hematopoiesis. The lack of Cyba induces a myeloid bias and promotes the enrichment of HSC populations, together with an increased proliferation potential. Moreover, the lack of p22phox hinders the activation of the PI3K-AKT pathway, and induces STAT5 downregulation. This would jeopardize the activation of IL-7 signaling, which would explain the increased immunoglobulin production in Cyba− mice.

Disclosures
No conflicts of interest to disclose

Contributions
RPB performed experiments, analyzed data, assembled figures, revised the manuscript. MRG and APF performed experiments and revised the manuscript. IGT and MSM generated the Cyba− mice. AHH conceived and designed experiments, performed experiments, analyzed data and wrote the manuscript.

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