Human miR223 Promoter as a Novel Myelo-Specific Promoter for Chronic Granulomatous Disease Gene Therapy

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Abstract

Targeting transgene expression to specific hematopoietic cell lineages could contribute to the safety of retroviral vectors in gene therapeutic applications. Chronic granulomatous disease (CGD), a defect of phagocytic cells, can be managed by gene therapy, using retroviral vectors with targeted expression to myeloid cells. In this context, we analyzed the myelosppecificity of the human miR223 promoter, which is known to be strongly upregulated during myeloid differentiation, to drive myeloid-restricted expression of p47^{phox} and gp91^{phox} in mouse models of CGD and in primary patient-derived cells. The miR223 promoter restricted the expression of p47^{phox}, gp91^{phox}, and green fluorescent protein (GFP) within self-inactivating (SIN) gamma- and lentiviral vectors to granulocytes and macrophages, with only marginal expression in lymphocytes or hematopoietic stem and progenitor cells. Furthermore, gene transfer into primary CD34+ cells derived from a p47^{phox} patient followed by *ex vivo* differentiation to neutrophils resulted in restoration of *Escherichia coli* killing activity by miR223 promoter-mediated p47^{phox} expression. These results indicate that the miR223 promoter as an internal promoter within SIN gene therapy vectors is able to efficiently correct the CGD phenotype with negligible activity in hematopoietic progenitors, thereby limiting the risk of insertional oncogenesis and development of clonal dominance.

Introduction

CHRONIC GRANULOMATOUS DISEASE (CGD) comprises a group of immunodeficiencies characterized by malfunctioning of the phagocytic nicotinamide adenine dinucleotide phosphate (NADPH) oxidase, resulting in defective reactive oxygen species (ROS)-dependent killing of microbes and consequently leading to recurrent severe bacterial and fungal infections (Seger, 2008). About 67% of CGD cases result from mutations within the X-linked *CYBB* gene encoding the gp91^{phox} subunit of NADPH oxidase (X-CGD), whereas 16% result from malfunction of the p47^{phox} subunit (van den Berg *et al.*, 2009).

The first clinically successful gene therapy (GT) trial for primary immunodeficiency was reported in 2000 (Cavazzana-Calvo *et al.*, 2000). Since then, the list of indications for which

GT treatment led to permanent or temporary benefit for treated patients has been continuously growing (Aiuti *et al.*, 2002; Mavilio *et al.*, 2006; Morgan *et al.*, 2006; Ott *et al.*, 2006; Kaplitt *et al.*, 2007; Bainbridge *et al.*, 2008; Cartier *et al.*, 2009; Boztug *et al.*, 2010; Cavazzana-Calvo *et al.*, 2010; Porter *et al.*, 2011).

For CGD, the first clinically successful X-CGD GT trial with proven restoration of ROS production and resolution of treatment refractory *Aspergillus* infections was initiated in 2004 in Frankfurt and Zürich. In this study, a gammaretroviral long terminal repeat (LTR)-driven vector containing a strong spleen focus-forming virus (SFFV) promoter/enhancer was used to express gp91^{phox}. However, vector-mediated transactivation of the *EVII* proto-oncogene in progenitor cells resulted in clonal dominance progressing to genetic instability and myelodysplastic syndrome (MDS)

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(Stein *et al.*, 2010). These severe side effects revealed the need of vector redesign to avoid vector-mediated genotoxic effects. This could be achieved by using vectors with self-inactivating (SIN) configuration and cell-type-specific internal enhancer/promoters, thereby restricting transgene expression to the deficient cell compartment (i.e., to granulocytes, monocytes, and macrophages in CGD) and limiting the risk of proto-oncogene transactivation to cells with limited life span (reviewed in Toscano *et al.*, 2011). Herewith, we tested the human miR223 regulatory region, which is highly active only in myeloid cells, in the context of SIN gamma- and lentiviral vectors (Fukao *et al.*, 2007) for its expression specificity, stability over time, and its ability to restore NADPH oxidase function and mediate bacterial killing.

Materials and Methods

Antibodies and reagents

Anti-human CD14-FITC, anti-human CD206-FITC (clone 19.2), anti-Ly6G-FITC (clone 1A8), anti-Gr1-PE (clone RB6-8C5), anti-B220-FITC or APC-Cy7 (clone RA3-6B2), anti-CD3-PE (clone 17A2), anti-CD11b-PE (clone M1/70), anti-ScaI-PE (clone E13-161.7), mouse Fc-block (clone 2.462), and Cytofix/Cytoperm Kit were from Beckton Dickinson (Basel, Switzerland), as was anti-p47^{phox} (clone 1)-APC labeled by Beckton Dickinson services. Anti-CD11b-PerCP/Cy5.5, APC or PE-Cy7 (clone M1/70), anti-F4/80-PerCP/Cy5.5 (clone BM8) and anti-CD117-FITC or APC (clone 2B8), CD45.1-PE or PE-Cy7 (clone A20), CD45.2-PerCP-Cy5.5 (clone 104), and ScaI-PE-Cy7 (clone D7) were from eBiosciences (Vienna, Austria). Anti-CD11C-PerCP (clone N418) came from Biolegend (Uithoorn, The Netherlands). Anti-ΔLNGFR-APC, human and mouse Lineage Cell Depletion Kits, CD14 microbeads, and Gr1-VioBlue (clone RB6-8C5) were from Miltenyi Biotec GmbH (Bergisch Gladbach, Germany). For intracellular p47^{phox} staining combined with surface staining, cells were blocked for 10 min at 4°C, incubated with surface staining antibodies for 1 hr, and washed twice followed by intracellular staining using Cytofix/Cytoperm according to manufacturer's description.

Animals

p47^{phox}-/- (B6[Cg]-Ncf1^{m1J}/J) and gp91^{phox}-/- (B6.129S-Cybb^{tm1Din}/J) mouse husbandry and animal experiments were carried out in compliance with the local animal protection law.

Vector construction

The 5' regulatory region of the human miRNA223 promoter (782 bp, chrX:65,234,612-65,235,394 GRCh37/hg19 assembly) was amplified from PLB985 gDNA using the following primers: forward, 5'-CATATGACTTGTACAGCTTCACAGGGCTCC (*NdeI* linker); reverse, 5'-GGATCCCTGT CAGTGGAGTGGTGC (*BamHI* linker). This element was cloned into lentiviral vectors via the *NdeI*-*BamHI* sites to replace the MRP8 promoter in the MRP8-gp91-wPRE (MgpW) vector (Brendel *et al.*, 2011) or the corresponding GFP-containing MEW vector. Gammaretroviral constructs were derived from the pSER11M8ΔN91s vector (M. Grez) by exchanging the transgene against a synthetic human p47^{phox} reading frame (Geneart, Regensburg, Germany) and by ex-

changing the internal promoter against promoter candidates (Supplementary Fig. S1; Supplementary Material available online at www.liebertonline.com/hgtb). The gammaretroviral vector encoding SFFV:ΔLNGFR-2A-p47^{phox} was already described (Wohlgensinger *et al.*, 2010). In this construct, the SFFV promoter was exchanged against the miR223 promoter. Lentiviral vectors encoding miR223 p47^{phox} were derived from the pCCLCHIM.GP91^{phox}.WPRES (Santilli *et al.*, 2011) by exchanging the promoter and the transgene.

Virus production

Virus supernatants were produced in 293T cells upon cotransfection with pMD2.VSV.G with CMVΔR8.91 for lentiviral vectors or pUMVS plus the gene transfer vector for gammaretroviral vectors (Supplementary Fig. S1), using TransIT-293 transfection reagent (Mirus Bio LLC, Madison, WI) or calcium phosphate-mediated transfection. Culture supernatants were collected after 24 and 48 hr, sterile-filtered, concentrated in Ultracal-100K Centrifugal Filters (Millipore, Zug, Switzerland) or by ultracentrifugation (2 hr at 50,000 g at 4°C), and stored at -80°C.

Transduction and ex vivo granulocytic differentiation

Murine lineage-negative (Lin-) bone marrow (BM) cells were isolated from femur and tibiae of 6-12-week-old mice using the Miltenyi lineage cell depletion kit (Miltenyi Biotec GmbH). The cells were prestimulated for 24 hr in StemSpan SFEM (StemCell Technologies, Grenoble, France) supplemented with mouse stem cell factor (mSCF) (10 ng/ml), mouse thrombopoietin (mTPO) (20 ng/ml), mouse insulin-like growth factor 2 (mIGF-2) (20 ng/ml), and human fibroblast growth factor 1 (hFGF1) (10 ng/ml) (all from Peprotech, Hamburg, Germany) at 0.5×10^6 to 1×10^6 cells per milliliter. Transduction was carried out at 20-50 multiplicity of infections (MOIs) in the presence of protamine sulfate (6 μg/ml) for 24 hr in 24-well plates. The cells were cultivated for another 24 h before transplantation. For granulocytic differentiation *in vitro*, the cells were incubated in RetroNectin (Takara Bio Europe/SAS, Saint-Germain-en-Laye, France) plates in differentiation medium 1 (RPMI) supplemented with 20% fetal calf serum (FCS) and 25 ng/ml mouse granulocyte colony-stimulating factor (mG-CSF) for 3 days up to 2 weeks.

After informed consent, human Lin- cells were isolated from BM aspirates of a p47^{phox} CGD patient, using the Lineage Cell Depletion Kit (Miltenyi Biotec GmbH). Lin- BM cells were expanded for 3 days on 24-well plates coated with RetroNectin in X-Vivo 10, penicillin/streptomycin, 2 mM L-glutamine, 50 ng/ml human interleukin-3 (hIL-3), 150 ng/ml human stem cell factor (hSCF), 150 ng/ml human FMS-like tyrosine kinase 3 (flt3) ligand (hFlt3L), 50 ng/ml human thrombopoietin (hTPO), and 2% FCS (or 1% bovine serum albumin). For retroviral transduction, cells were harvested and centrifuged onto virus preloaded RetroNectin plates.

For *ex vivo* differentiation, cells were harvested again and seeded on RPMI (10% FCS, penicillin/streptomycin, 50 ng/ml hSCF, 10 ng/ml human granulocyte colony-stimulating factor (hG-CSF), 10 ng/ml hFLT3L) at day 1. At day 10, cells were shifted to differentiation medium 2 (differentiation medium 1 without SCF) and incubated for another 10 days, followed by

FACS sorting using a BD FACSAria Flow Cytometer (Beckton Dickinson, Basel, Switzerland).

Animal GT

p47^{phox}^{-/-} and gp91^{phox}^{-/-} mice received lethal (9.5 Gy or a split dose of 2×5.5 Gy) irradiation. About 1×10⁶ to 2×10⁶ gene-modified Lin⁻ BM cells were injected intravenously. Animals were sacrificed after 7–23 weeks, and blood, spleen, BM, and peritoneal macrophages were analyzed for transgene expression.

Escherichia coli killing assay

The assay was conducted as described recently (Ott *et al.*, 2006) with minor modification. Granulocytes were co-incubated with opsonized *E. coli* at a ratio of 1:10 for 30 min at 32°C. Essentially, neutrophils were incubated with the *E. coli* strain ML-35. This particular strain lacks the membrane transport protein lactose permease and constitutively expresses β -galactosidase (β -Gal). Engulfment of *E. coli* ML-35 by neutrophils is followed by perforation of the bacterial cell wall and accessibility to β -Gal, which is subsequently inactivated by ROS (Hamers *et al.*, 1984). Briefly, 2×10⁹ *E. coli*/ml were opsonized with 20% (v/v) Octaplas (Octapharma AG, Lachen, Switzerland) for 5 min at 37°C. Opsonized *E. coli* (final concentration, 0.9×10⁸/ml) were added to granulocytes (0.9×10⁷/ml). At defined time points, granulocytes were lysed with 0.05% saponin (Calbiochem, Darmstadt, Germany) and the samples were incubated with 1 mM ortho-nitrophenyl- β D-galactopyranoside (Sigma-Aldrich, Seelze, Germany) at 37°C for 30 min. β -Gal activity was followed by standard procedures at 420 nm.

Human monocytes/macrophage culture

After informed consent, human monocytes were isolated using CD14 microbeads. Cryostorage in CryoSto CS10 (StemCell Technologies) resulted in 30% survival upon thawing. Monocytes were cultured, transduced at a high MOI, and differentiated to macrophages in RPMI, 10% Octaplas, 2 mM L-Gln, 10 mM HEPES, 2 mM sodium pyruvate, and M-CSF 100 ng/ml.

Nitroblue tetrazolium test

Macrophages were incubated in 1 mg/ml nitroblue tetrazolium (NBT; Sigma-Aldrich Chemie GmbH, Buchs, Switzerland) in phosphate buffered saline (PBS) containing 200 ng/ml phorbol myristate acetate (PMA) plus 0.5 nmol/ml N-Formyl-Met-Leu-Phe (fMLP) (both from Sigma-Aldrich Chemie GmbH) for 1 hr at 37°C and fixed thereafter in PBS and 2% formaldehyde (Merck [Schweiz] AG, Zug, Switzerland). Photographs were taken on a Leica DM-IL microscope with a DFC-480 camera (Leica Microsystems [Schweiz] AG, Heerbrugg, Switzerland).

Dihydrorhodamine 123 assay

About 10⁶ granulocytes in 1 ml of Hanks Balanced Salt Solution (HBSS) with Ca/Mg, 0.5% human albumin, 1000 U catalase, 7.5 mM glucose, and antibodies were stimulated with 200 ng PMA and incubated for 5 min at 37°C. Then, dihydrorhodamine 123 (DHR) was added to a final concentration of 0.435 μ M, incubated for further 15 min at 37°C, kept on

ice for 30 min, and analyzed using FACS Calibur (Beckton Dickinson, Basel, Switzerland) or FACS Canto II (Beckton Dickinson, Heidelberg, Germany).

Results

The miR223 promoter induces transgene expression upon granulocytic differentiation in vitro

To identify a myeloid-restricted promoter adequate for the gene therapy of CGD, we analyzed the promoter activity of a 782-bp DNA fragment derived from the human miR223 promoter within SIN gamma- and lentiviral vector constructs, as earlier reports have indicated strong and specific upregulation of miR223 gene expression during myeloid differentiation (Fukao *et al.*, 2007; Landgraf *et al.*, 2007; Johnnidis *et al.*, 2008). We tested the miR223 promoter in conjunction with the gp91^{phox} and the p47^{phox} subunits of the phagocytic NADPH oxidase as well as with a Δ LNGBFR-2A-p47^{phox} fusion construct (Wohlgensinger *et al.*, 2010) (Supplementary Fig. S1).

Initially, we transduced murine Lin⁻ BM cells obtained from p47^{phox}^{-/-} animals with SIN gammaretroviral vectors harboring p47^{phox} under control of the miR223 promoter, and compared this with the specificity of other promoters derived from myeloid-restricted genes such as *cFes* (Heydemann *et al.*, 2000) and *MRP8* (Brendel *et al.*, 2011), or synthetic promoters like the p40/48 (a 250-nt chimeric promoter consisting of two PU.1 transcription factor binding sites from the p40 and one PU1 site from the p47 promoter), or the Sp107 (He *et al.*, 2006) promoters. We used the SFFV promoter as a benchmark for high expression levels in myeloid cells (Supplementary Fig. S1). p47^{phox} expression was analyzed by FACS upon intracellular p47^{phox} staining 2 days after transduction in Sca1⁺ cells as well as in *ex vivo* differentiated Ly6G⁺ granulocytes generated thereof (Supplementary Fig. S2A). The mean fluorescence intensity (MFI) ratio of p47^{phox} expression in differentiated versus undifferentiated cells was used to estimate the induction of transgene expression upon myeloid differentiation. In this screen, the miR223 promoter showed the strongest induction of p47^{phox} expression among the promoters tested (Supplementary Fig. S2C). In addition, differentiated cells were analyzed for ROS production in a DHR oxidation assay. We observed a consistent increase in DHR⁺ granulocytes from *cFes*, p40/48, and the miR223-corrected cells, but only the miR223-corrected population showed a statistically significant increase of DHR⁺ cells compared with the mock control (Supplementary Fig. S2B and D). Next, we transduced the human myelomonocytic cell line X-CGD PLB985 with a lentiviral SIN vector encoding gp91^{phox} under control of the miR223 promoter (223gpW). Transductions at MOIs of 2 and 0.2 resulted in moderate gp91^{phox} expression levels in 51% and 10% of the cells, respectively. Upon differentiation, the percentage of gp91^{phox}-expressing cells increased to 70% and 19%, respectively, in parallel to a significant increase in expression levels (Fig. 1A). Restoration of ROS production could be confirmed in 57% of the cells transduced at an MOI of 2, with DHR MFIs comparable to those of differentiated wild-type PLB-985 cells (Fig. 1B).

We generated single clones of PLB-985 cells transduced with a lentiviral vector encoding GFP under control of the miR223 promoter in order to quantify gene expression

FIG. 1. Performance of miR223 promoter-containing lentiviral vectors in the human myelomonocytic X-CGD cell line PLB985. **(A)** FACS analysis of gp91^{phox} expression in undifferentiated (gray) and differentiated (black) X-CGD-PLB985 cells transduced with 223gpW at MOIs 2 and 0.2 along with nontransduced controls. **(B)** NADPH oxidase activity in 223gpW-transduced X-CGD-PLB985 cells assessed by the DHR assay. **(C)** Correlation of VCN and GFP expression levels in undifferentiated (diamonds) and differentiated (circles) PLB985 clonal cell populations transduced with 223GFP. **(D)** Fold upregulation of GFP expression levels during granulocytic differentiation in clonal populations derived from 223GFP-transduced PLB985 cells. Variation of expression levels is given as percentage of coefficient of variation (%CV). **(E)** Cumulative long-term stability of GFP expression in 223GFP-transduced PLB985 cells during a 4-month culture period in differentiated and undifferentiated clonal populations. CGD, chronic granulomatous disease; DHR, dihydrorhodamine 123; GFP, green fluorescent protein; MOI, multiplicity of infection; NADPH, nicotinamide adenine dinucleotide phosphate; VCN, vector copy number.

without variations resulting from antibody staining. The analysis revealed a good correlation between vector copy number (VCN) and transgene expression particularly in differentiated cells ($R^2 = 0.94$), whereas a reduced correlation was evident in undifferentiated cells ($R^2 = 0.69$) (Fig. 1C). In 53 monoclonal populations, the mean fold induction of GFP expression upon granulocytic differentiation was 3.5 ± 0.88 (Fig. 1D) with a coefficient of variation of only 25.7%. Moreover, GFP signals in these clones remained stable during the total observation period of 4 months (Fig. 1E).

The induction of transgene expression and reconstitution of NADPH oxidase activity could also be confirmed in granulocytes obtained upon *ex vivo* differentiation of transduced murine Lin⁻ gp91^{phox}/BM cells. Lentiviral transduction with the vector harboring gp91^{phox} under SFFV promoter control (SgpW) resulted in similar expression of gp91^{phox} in hematopoietic progenitors (cKit⁺ Sca1⁺) and granulocytes (CD11b⁺ Gr1⁺) (Fig. 2A) with good reconstitution of ROS production in differentiated CD11b⁺ Gr1⁺ cells (Fig. 2B). On the other hand, the miR223 promoter

FIG. 2. Expression of gp91^{phox} and NADPH oxidase activity after *in vitro* culture of 223gpW-transduced murine X-CGD Lin⁻ BM cells. **(A)** 223gpW-transduced murine Lin⁻ cells were analyzed for gp91^{phox} expression in stem and progenitor cells (lower quadrants) and in granulocytes (upper quadrants). **(B)** Functional reconstitution of NADPH oxidase activity in X-CGD neutrophils mediated by the recombinant expression of gp91^{phox} as assessed by the DHR assay. BM, bone marrow; Lin⁻, lineage-negative.

(223gpW) mediated weak gp91^{phox} expression in a small fraction (6%) of the hematopoietic progenitors (cKit⁺ Sca1⁺ CD11b⁻ Gr1⁻), whereas 62% of the CD11b⁺ Gr1⁺ cells showed significant amounts of gp91^{phox} expression, confirming induction of miR223 promoter activity upon granulocytic differentiation. As in PLB-985 cell-derived granulocytes (Fig. 1B), miR223-driven gp91^{phox} expression also mediated high levels of ROS production in granulocytes derived from transduced primary murine Lin⁻ cells with ROS levels similar to those observed in granulocytes derived from wild-type (C57Bl6) or SgpW-transduced X-CGD cells (Fig. 2B).

Collectively, these results strongly indicate an induction of miR223 promoter activity upon granulocytic differentiation regardless of transgene identity or the vector system utilized.

The miR223 promoter drives myelo-specific transgene expression in CGD mouse models in vivo

To characterize the miR223 promoter activity *in vivo*, we monitored transgene expression patterns in gp91^{-/-} and p47^{-/-} mouse models of CGD. After lentiviral transduction of X-CGD Lin⁻ BM cells with 223gpW and reinfusion into lethally irradiated recipient mice, gp91^{phox} expression was analyzed in donor cells up to 23 weeks after transplantation (Fig. 3A and B; Supplementary Fig. S3). Expression in BM LSK (Lin⁻ Sca1⁺ cKit⁺) cells and lymphoid cells (CD3⁺ or B220⁺) was almost undetectable, while increasing amounts of gp91^{phox} were detected during differentiation from progenitor LSK cells (Lin⁻ Sca1⁻ cKit⁺) to monocytes (CD11b⁺ Gr1⁻) and to mature granulocytes (CD11b⁺ Gr1⁺). Gp91^{phox} expression in granulocytes led to the reconstitution of NADPH oxidase activity and ROS production as analyzed by the DHR assay. The percentages of DHR⁺ granulocytes

correlated with the percentages of gp91^{phox} expression in individual animals, and DHR MFIs indicate the production of significant amounts of ROS at moderate VCNs between 0.7 and 1.5 (Fig. 3B).

In the p47^{-/-} animal model, an miR223 p47^{phox} gamma-retroviral vector (Supplementary Fig. S1) was used to transduce syngeneic Lin⁻ BM cells, and cell populations were analyzed for p47^{phox} expression by intracellular staining 7 weeks after reinfusion (Fig. 3C). Very low expression of p47^{phox} was observed in BM cKit⁺ Sca1⁺ or splenic T and B cells (data not shown), and expression levels in monocytes remained under the detection limit. Expression levels in granulocytes were below 11% at an average of 4%, indicating one copy per cell (Fehse *et al.*, 2004). Although MFIs of p47^{phox}⁺ granulocyte signals were moderate, ROS production was restored in all animals (Fig. 3D). The percentages of DHR⁺ granulocytes closely correlated with the percentages of p47^{phox} FACS⁺ granulocytes ($R^2 = 0.9872$). No adverse events regarding animal health, organ size, and appearance were observed during these studies. Furthermore, there was no decline of expression over time, which might indicate the absence of epigenetic silencing.

The miR223 promoter is active in p47^{phox}-deficient macrophages

As NADPH oxidase function is required in granulocytes and in macrophages, we tested transgene expression in peritoneal macrophages of NADPH oxidase-deficient animals. Lethally irradiated p47^{phox}^{-/-} mice were reconstituted with Lin⁻ cells transduced with a gammaretroviral vector encoding the Δ LNGBR-2A-p47^{phox} fusion construct under miR223 control, allowing for indirect p47^{phox} detection by Δ LNGBR surface staining (Wohlgensinger *et al.*, 2010).

FIG. 3. *In vivo* expression profile of miR223 promoter-driven gp91^{phox} or p47^{phox} vectors. (A) Lentivirally (223gpW) transduced Lin⁻ BM cells from gp91^{phox}^{-/-} (CD45.2) mice were transplanted into irradiated SJL mice (CD45.1). Expression of gp91^{phox} in different donor-derived hematopoietic lineages was determined 23 weeks after transplantation by flow cytometry. Shown in overlay histograms: gray, gp91-X-CGD-PLB985 cells; black, transduced sample. (B) Analysis of the NADPH oxidase activity in peripheral blood granulocytes derived from the mice shown in (A). (C) Lin⁻ BM cells from p47^{-/-} mice were gammaretrovirally (miR223 p47^{phox}) transduced and transplanted into congenic recipient mice. Analysis of p47^{phox} expression and reconstitution of NADPH oxidase activity (D) was performed 8 weeks after transplantation.

FIG. 4. miR223 promoter activity in macrophages. Lin⁺BM cells from p47^{phox}^{-/-} mice were gammaretrovirally transduced (miR223-ΔLNGFR-2A-p47^{phox}) and reinfused into lethally irradiated p47^{phox}^{-/-} mice. FACS analysis of (A) ΔLNGFR expression in peritoneal macrophages and (B) T, B, and dendritic cells after 8 weeks revealed preferential expression in peritoneal macrophages. (C) Frozen blood monocytes from a p47^{phox} CGD patient were taken into culture overnight and lentivirally (miR223-p47^{phox}) transduced. The cells were kept in culture for 7 days, followed by FACS analysis of p47^{phox} expression in macrophages (CD206+ CD14+). (D) Functional reconstitution of ROS production was analyzed in macrophages by nitroblue tetrazolium assay. ROS, reactive oxygen species.

Significant ΔLNGFR expression was observed in F4/80+ peritoneal macrophages in all treated animals, indicating miR223 promoter activity in macrophages (Fig. 4A), while ΔLNGFR detection signals in splenic T and B cells and blood dendritic cells were in the range of background staining (Fig. 4B). Next, we analyzed the miR223 promoter activity in primary human macrophages obtained from blood monocytes of a p47^{phox}-deficient CGD patient. Monocytes were transduced with lentiviral vectors encoding p47^{phox} under control of the SFFV or the miR223 promoter and differentiated to macrophages for 7 days, resulting in high levels of transduction for both vectors (Fig. 4C). ROS production in primary macrophages was confirmed in an NBT reduction assay upon PMA/fMLP stimulation (Fig. 4D), clearly indicating miR223 promoter activity in this cell type.

The miR223 promoter-controlled p47^{phox} expression restores E. coli killing activity in primary human granulocytes

To analyze whether miR223 promoter-driven p47^{phox} expression restores granulocyte anti-microbial activity, we transduced Lin⁺ BM cells from a p47^{phox}-deficient CGD patient with the gammaretroviral vector encoding the ΔLNGFR-2A-p47^{phox} fusion construct under miR223 or SFFV transcriptional control followed by granulocytic differentiation *ex vivo*. ΔLNGFR surface staining circumvents intracellular staining for p47^{phox} for functional analyses and allows for the enrichment of gene-modified cells. In differentiated myeloid cells, gene marking rates of 7%–10% were observed (Fig. 5A). Stimulation of these cells with either PMA or opsonized *E. coli* resulted in DHR oxidation exclusively in ΔLNGFR+ cells, indicating restoration of ROS production in a p47^{phox}-dependent manner (Fig. 5B). ΔLNGFR surface

marking of neutrophils, together with CD11b marking of differentiated granulocytes, was used to FACS-sort mock-transduced p47^{phox}+ (ΔLNGFR+) and p47^{phox}- (ΔLNGFR-) granulocytes (all CD11b+) derived from γRetro-SIN miR223-ΔLNGFR-2A-p47^{phox} and γRetro-SIN SFFV-ΔLNGFR-2A-p47^{phox}-transduced cells. Sorted cell populations were then analyzed in an *E. coli* killing assay (Ott *et al.*, 2006). In this assay, an increase in optical density at 420 nm indicates effective bacterial killing. Both ΔLNGFR+ populations showed effective restoration of *E. coli* killing activity, whereas the CD11b+/ΔLNGFR- populations derived from the same cultures were unable to kill *E. coli* (Fig. 5C). Thus, antimicrobial function was restored in neutrophils expressing p47^{phox} under control of the miR223 (as well as SFFV) promoter. These data reinforce the potential utility of the miR223 promoter for high and myeloid-restricted transgene expression.

Discussion

Transactivation of proliferation-promoting genes as the primary cause of clonal dominance in gene therapy studies can potentially be avoided by the use of SIN vectors driving transgene expression from an internal tissue-specific promoter rather than from strong constitutively active promoters such as the viral LTR (Zychlinski *et al.*, 2008; Modlich *et al.*, 2009; Montini *et al.*, 2009). Another potential risk arises from transcriptional read-through, that is, the extension of viral transcripts through the virus-genome border into the genomic sequence or splicing of vector transcripts into cellular exons. Both features have contributed to leukemia formation in animal models and clinical studies (Cesana *et al.*, 2012). For CGD, a myelo-specific promoter driving transgene expression preferentially in phagocytes could be an option to

FIG. 5. *E. coli* killing activity of *ex vivo* differentiated human $p47^{\text{phox-/-}}$ granulocytes after gene transfer. Lin⁻ cells from a $p47^{\text{phox-/-}}$ CGD patient were mock transduced or gammaretrovirally transduced with the self-inactivating SFFV- Δ LNGFR-2A- $p47^{\text{phox}}$ or the self-inactivating miR223- Δ LNGFR-2A- $p47^{\text{phox}}$ vectors and propagated to granulocytes in liquid cell culture. (A) Expression of the surrogate marker Δ LNGFR reflecting $p47^{\text{phox}}$ expression in myeloid differentiated (CD11b⁺) and undifferentiated (CD11b⁻) cells. (B) Within the CD11b/ Δ LNGFR double-positive populations, 0.1%, 48.7%, and 49.1% responded by ROS production upon stimulation with phorbol myristate acetate as revealed in the DHR assay (upper panels). Stimulation with opsonized *E. coli* (lower panels) resulted in ROS production in 1.3%, 32.1%, and 35.5% of the double-positive cells. (C) Transduced CD11b⁺ cells were separated by FACS as indicated by the Δ LNGFR gates shown in (A) (purity: SFFV, 87%; miR223, 93%). The isolated populations were analyzed for their *E. coli* killing capacity in a standard killing assay using the *E. coli* strain ML35. An increase in optical density at 420 nm indicates bacterial killing activity.

limit transgene expression to short-lived cells, thereby theoretically limiting the risks associated with enhancer- or read-through-mediated activation of proto-oncogenes (Kustikova *et al.*, 2009; Toscano *et al.*, 2011). The study was motivated by the results obtained during the long-term follow-up of CGD patients treated by gene therapy revealing transactivation of *EVI1* as primary cause of clonal dominance with development of MDS, and promoter silencing leading to treatment failure (Stein *et al.*, 2010).

In our study, we evaluated the properties of a DNA fragment derived from the human miR223 regulatory region as an internal promoter within SIN retroviral vectors for the gene therapy of CGD, as the hsa-miR223 gene was reported to be transcribed nearly exclusively within the hematopoietic compartment (Landgraf *et al.*, 2007) in a highly myeloid-specific manner (Fukao *et al.*, 2007; Johnnidis *et al.*, 2008).

In the $gp91^{\text{phox}}$ -deficient mouse model, miR223-driven $gp91^{\text{phox}}$ expression resulted in up to 43% $gp91^{\text{phox}}$ granulocytes, but the same resulted in only 0.3%–2.2% $gp91^{\text{phox}}$ early stem and progenitor cells (LSK cells) and in a maximum of 1.8% positive T and B cells, as assessed by intracellular staining despite mean VCNs of 0.7–1.5 (Fig. 3A). The lack of $gp91^{\text{phox}}$ expression in immature primitive progenitors and B and T cells could be attributed to a detection bias, as $gp91^{\text{phox}}$ is not stable in cells lacking $p22^{\text{phox}}$, the second component of the cytochrome b558 complex. However, $p22^{\text{phox}}$ is ubiquitously expressed (Chambers *et al.*, 2007) and $gp91^{\text{phox}}$ can be easily detected in LSK cells transduced with SgpW, a lentiviral vector containing the SFFV enhancer/promoter element driving $gp91^{\text{phox}}$ expression (Fig. 2A) (Brendel *et al.*, 2011). Similarly, in the $p47^{\text{phox}}$ model, miR223-

mediated $p47^{\text{phox}}$ expression was weak in BM cKit⁺/ScaI⁺ cells and undetectable in T and B cells, while neutrophils were up to 10.3% $p47^{\text{phox}}$. Although the analyses of both models were performed at different time points, hematopoiesis after 7 or more weeks is mainly maintained by mid- and long-term repopulating multipotent progenitors (Dykstra *et al.*, 2007; Gerrits *et al.*, 2010). Therefore, lineage-specific expression is unlikely to originate from the transduction of lineage-restricted progenitors but rather truly reflects the specificity of the miR223 promoter element. The expression distribution observed in our experiments resembles the expression pattern of the *miR223* gene in mice (Johnnidis *et al.*, 2008) and humans (Allantaz *et al.*, 2012). Moreover, our *in vitro* experiments show that the miR223 promoter can direct $gp91^{\text{phox}}$ expression in a VCN-dependent manner and generates levels of superoxide equivalent to those observed in differentiated wild-type human myeloid cell lines and *in vitro* differentiated murine neutrophils (Figs. 1 and 2).

Myeloid-restricted expression of the *miR223* gene has been attributed to the action of a complex regulatory loop including the coordinate binding of CEBP and PU.1 to a region upstream of the major pri-miR223 transcription start site (MTSS) and to the binding of C/EBP α and NFI-A to a non-conserved responsive element 2710 bp downstream of the MTSS, which corresponds to an intron of the pri-miR223 transcript (Fukao *et al.*, 2007). Although the detailed interplay of these two regulatory regions in mediating tissue specificity has not been clarified, evidence exists that the intronic regulatory region contributes to miR223 expression specificity (Fazi *et al.*, 2007). The DNA fragment used in our studies comprises the MTSS and includes the CEBP and PU.1

sites but lacks the binding sites for C/EBP α and NF- κ B. Despite this, a myeloid-restricted expression pattern was observed *in vitro* and *in vivo*.

A recent study characterized a myelo-specific chimeric promoter generated by the fusion of cathepsin G and *c-Fes* minimal 5'-flanking regions driving gp91^{phox} expression within a lentiviral SIN vector (Santilli *et al.*, 2011). In comparison to this vector, the 223gpW exhibits even lower leakiness in lymphocytes and more importantly in LSK cells. One possible explanation for this is the lack of SP1-binding sites in the miR223 DNA fragment used in this study, which are present in the chimeric promoter and which may account for the low levels of gene expression seen in primitive progenitors transduced with SIN vectors containing the chimeric promoter (Santilli *et al.*, 2011).

Alternatively, miRNA target sequences can also be used to restrict transgene expression to the desired target population (Brown *et al.*, 2007). Best example of this is the use of the miR126 target sequences to detarget expression of an otherwise toxic gene product (galactocerebrosidase) from HSCs in the context of gene therapy for globoid cell leukodystrophy (Gentner *et al.*, 2010).

Although targeting gene expression to myeloid cells in the context of CGD is a reasonable approach, there is no experimental evidence for a toxic effect of gp91^{phox} expression in hematopoietic primitive progenitor cells (Grez *et al.*, 2011). Thus, the relevant issue for CGD vectors is the absolute reconstitution of superoxide levels, as the analysis of 227 CGD patients revealed a close correlation between superoxide production levels and long-term survival (Kuhns *et al.*, 2010).

Unfortunately, a restricted tissue-specific transgene expression frequently yields weak expression levels. The question is whether, despite the high specificity, miR223-mediated expression levels are sufficient to correct the CGD phenotype.

In our hands, FACS-sorted p47^{phox}+ neutrophils derived from the BM cells of transduced Lin[−] p47^{phox}-deficient patients clearly demonstrated *E. coli* killing activity *ex vivo* for both SFFV- and miR223-containing vectors compared with the fraction of untransduced neutrophils, indicating that the level of miR223-mediated p47^{phox} expression in neutrophils is sufficient to restore the physiological antimicrobial function of these cells (Fig. 5).

ROS production seems to possess a triggering role for antimicrobial defense mechanisms, and it is a prerequisite for NET formation and extracellular killing (Bianchi *et al.*, 2009). Already, very low amounts of granulocytic ROS production result in significant changes in membrane potential, K⁺ efflux, and bacterial killing (Rada *et al.*, 2004). This indicates that low levels of gp91^{phox} or p47^{phox} expression in CGD phagocytes could result in significant physiological improvement in the clinical status of treated patients. In mice, already 4%–8% of wild-type granulocytes or 11%–21% gene-modified neutrophils were sufficient to protect X-CGD mice against respiratory challenge with *Aspergillus fumigatus* conidia, although a significant higher number of gene-corrected neutrophils was required to improve host defense of X-CGD animals against *Staphylococcus aureus* and *Burkholderia cepacia* (Dinauer *et al.*, 2001; Urban *et al.*, 2009; Bianchi *et al.*, 2011). We expect our tissue-specific miR223 vectors to show an improved safety profile compared to vectors with ubiquitously active promoters, as our highly myelo-specific expression pattern strongly limits the promoter activity in

stem and progenitor cells and thus may prevent the occurrence of side effects while restoring the physiological antimicrobial function of gene-corrected neutrophils.

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Author Disclosure Statement

No competing financial interests exist.

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