Ex vivo Improvement of a von Willebrand Disease Type 2A Phenotype Using an Allele-Specific Small-Interfering RNA

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Introduction

Von Willebrand disease (VWD) is the most common inherited bleeding disorder and is mainly caused by dominant-negative mutations in the multimeric protein von Willebrand factor (VWF). These mutations may either result in quantitative or qualitative defects in VWF. VWF is an endothelial protein that is secreted to the circulation upon endothelial activation. Once secreted, VWF multimers bind platelets and chaperone coagulation factor VIII in the circulation. Treatment of VWD focuses on increasing VWF plasma levels, but production and secretion of mutant VWF remain uninterrupted. Presence of circulating mutant VWF might, however, still affect normal hemostasis or functionalities of VWF beyond hemostasis. We hypothesized that inhibition of the production of mutant VWF improves the function of VWF overall and ameliorates VWD phenotypes. We previously proposed the use of allele-specific small-interfering RNAs (siRNAs) that target frequent VWF single nucleotide polymorphisms to inhibit mutant VWF. The aim of this study is to prove the functionality of these allele-specific siRNAs in endothelial colony-forming cells (ECFCs). We isolated ECFCs from a VWD type 2A patient with an intracellular multimerization defect, reduced VWF collagen binding, and a defective processing of proVWF to VWF. After transfection of an allele-specific siRNA that specifically inhibited expression of mutant VWF, we showed amelioration of the laboratory phenotype, with normalization of the VWF collagen binding, improvement in VWF multimers, and enhanced VWF processing. Altogether, we prove that allele-specific inhibition of the production of mutant VWF by siRNAs is a promising therapeutic strategy to improve VWD phenotypes.
HEK293 cells are a good model to prove the principle of allele-specific VWF inhibition and to select for efficient and specific siRNA candidates; however, HEK293 cells do not endogenously produce VWF. We therefore aim in this study to test the approach of allele-specific VWF inhibition by SNP-targeted siRNAs in endothelial colony-forming cells (ECFCs; previously called blood outgrowth endothelial cells or BOECs). ECFCs are cultured endothelial cells that can be isolated from peripheral blood.19,20 We show that siRNAs are able to selectively inhibit VWF alleles in healthy control ECFCs and that allele-specific siRNAs can improve the VWD phenotype of ECFCs that were isolated from a VWD type 2A patient with the VWF p.Cys1190Tyr mutation.

Methods

Patients and Controls
A VWD type 2A patient with the VWF p.Cys1190Tyr (c.3569G > A) mutation and five healthy control subjects were included in the study. A total of 50 to 70 mL of peripheral blood was drawn for ECFC, plasma, and genomic DNA isolation. The patient’s plasma was analyzed for VWF antigen (VWF:Ag), VWF activity, and FVIII activity. VWF:Ag in plasma was determined using the STA LIA VWF:Ag test (Stago, Leiden, the Netherlands) and was analyzed on the Sta-R Max analyzer (Stago) with the HemosIL AcuStar VWF:RCo reagent (Werfen IL, Breda, the Netherlands). Samples were analyzed on the BIO-FLASH (Werfen) and a commercial calibrator (supplied with the HemosIL AcuStar VWF:RCo) was used as the reference. The FVIII activity was determined using an automated one-stage clotting assay on the STA-RA MAX analyzer (Stago) with Sta-immunodef VIII (Stago) and STA-CK Pres 5 (APTT) (Stago) reagents. Commercial normal pool plasma (STA Uni-Calibrator, Stago) was used as the reference. Plasma characteristics of the patient and controls included in the study are summarized in Table 1.

Genomic DNA was used to confirm the mutation in the VWD patient and to determine the genotypes in all subjects for four VWF SNPs: rs1800378, rs1063856, rs1063857, and rs1800380. Genotypes were determined by Taqman SNP

Table 1 Plasma phenotype of included subjects

|       | VWF:Ag (IU/dL) | VWF:GPIbR (IU/dL) | FVIII:C (IU/dL) | VWF:GPIbR/ VWF:Ag |
|-------|----------------|-------------------|----------------|-------------------|
| ECFC C1 | 105            | 73                | 97             | 0.70             |
| ECFC C2 | 80             | 68                | 98             | 0.85             |
| ECFC C3 | 101            | 87                | 136            | 0.86             |
| ECFC C4 | 108            | 95                | 149            | 0.88             |
| ECFC C5 | 85             | 80                | 144            | 0.94             |
| ECFC 2A | 107            | 27                | 78             | 0.25             |

Abbreviations: VWF, von Willebrand factor; VWF:Ag, VWF antigen; VWF:GPIbR, VWF ristocetin-triggered GPIb binding assay.
genotyping assays (Thermo Fisher Scientific, Carlsbad, California, United States).

The study protocol was approved by the institutional ethical review board. Informed consent was obtained from all subjects in accordance with the Declaration of Helsinki.

SNP Phasing
PacBio long-read single-molecule sequencing was used for linkage analysis of the heterozygous SNP and the p.Cys1190-Tyr mutation. RNA was isolated from the ECFC cell pellet using the RNeasy Micro Kit (Qiagen, Venlo, the Netherlands). Complementary DNA (cDNA) was synthesized using SuperScript II Reverse Transcriptase (Thermo Fisher Scientific) with primers designed to span a region of 2.8 kb (forward primer: CACCTTCAGTGGGATCTGCC; reverse primer: TTTCAGACCTCGCTGGTGG). VWF-specific products were amplified using the KAPA HiFi HotStart ReadyMix PCR kit (Roche Diagnostics, Mannheim, Germany). Amplicons were barcoded and SMRTbell adapters were added using the SMRTbell Barcoded Adapter Complete Prep Kit (Pacific Biosciences, Menlo Park, California, United States). Barcoded amplicons were sequenced using a P6-C4 SMRT cell on a Pacific Biosciences RSII sequencer. Error-free circular consensus reads were mapped to the reference, followed by variant calling and resolution of variant phase using WhatSThap suit.

ECFC Isolation and Culture
ECFCs were isolated from peripheral blood that was drawn in Lithium Heparin tubes (ECFC C1, C2, and C4; BD Biosciences, Erembodegem, Belgium) or in Sodium Heparin CPT Mononuclear Cell Preparation tubes (ECFC C3, C5, and 2A; BD Biosciences). Blood was diluted 1:1 in phosphate buffered saline (PBS) and layered over Ficoll-Paque (LUMC Pharmacy, Leiden, the Netherlands), followed by centrifugation. The mononuclear cell fraction was washed twice in PBS supplemented with 10% fetal bovine serum (FBS), after which cells were taken up in Lonza EGM-2 BulletKit medium. ECFCs were cultured in EGM-2 BulletKit (Lonza, Breda, the Netherlands) or in FBS (GIBCO, Carlsbad, California, United States), and 7 mL Antibiotic-Antimycotic solution (Sigma-Aldrich) was used as a substrate and dissolved in 11 mL buffer (PBS containing 1% bovine serum albumin (BSA)) to generate rabbit anti-VWF-IgG-Biotin. ELISA plates were coated with unprocessed proVWF in lysates was quantitated with the modification that samples were diluted in PBS containing 0.1% Tween 20 (PBS-tween), or in PBS containing 1% bovine serum albumin (BSA) when the VWF:Ag ELISA was performed simultaneously with the VWF collagen binding (VWF:CB) assay. The amount of unprocessed proVWF in lysates was quantified by ELISA.

VWF Protein Analysis
VWF:Ag levels in conditioned medium and protein lysates were measured by enzyme-linked-immunosorbent serologic assay (ELISA) as described before, with the modification that samples were diluted in PBS containing 0.1% Tween 20 (PBS-tween), or in PBS containing 1% bovine serum albumin (BSA) when the VWF:Ag ELISA was performed simultaneously with the VWF collagen binding (VWF:CB) assay. The amount of unprocessed proVWF in lysates was quantified by ELISA.

siRNA Transfections
Custom Silencer Select siRNAs (Life Technologies, Bleiswijk, the Netherlands), a negative control siRNA (s1NEG, 4404020, Life Technologies), or an siRNA against VWF (s1VWF, s14834, Life Technologies) were transfected in ECFCs using DharmaFECT duo transfection reagent (Dharmacon, Lafayette, Colorado, United States). For transfections, 100,000 to 125,000 cells per well were plated on rat tail collagen (50 µg/mL; BD biosciences) coated wells of a 24-well plate. siRNAs were transfected into ECFCs 24 hours after plating the cells. Before transfection, cells were washed twice with Hanks’ Balanced Salt solution (Thermo Fisher Scientific) and once with PromoCell EGM2 culture medium. siRNAs and DharmaFECT duo were separately diluted in Optimem1 (Thermo Fisher Scientific) and mixed together by pipetting. After 20 minutes of incubation, the siRNA–DharmaFECT mixture was complemented with Optimem1 supplemented with 4% FBS. Cells were incubated with 150 µL transfection mixture containing a high concentration of siRNA for 3 hours. After the 3-hour incubation, 250 µL culture medium was added to obtain a final siRNA concentration of 20 nM (unless otherwise stated). The medium was refreshed 24 hours after transfection and every other day thereafter. Six days after transfection, the medium was harvested and cells were lysed with protein lysis buffer containing Optimem1, 0.1% Triton X-100 (Sigma-Aldrich) and a tablet of Complete protease inhibitor cocktail with EDTA (Roche Diagnostics, Mannheim, Germany).
with protein lysates diluted in PBS-tween. Mouse anti-VWFpp conjugated to horseradish peroxidase (CLB-Pro 14.3, Sanquin, Amsterdam, the Netherlands) was used as a detection antibody and ELISA plates were incubated with the detection antibody diluted in PBS-tween. o-Phenylenediamine dihydrochloride (Sigma-Aldrich) was used as a substrate and dissolved in substrate buffer with the addition of 11 μL 30% H₂O₂. The enzymatic reaction was followed kinetically for 5 minutes. The ratio of unprocessed VWF over the total VWF:Ag levels was used as a measure for the processing of VWF. The average of control samples was set to 1.

VWF multimers were visualized using agarose gel electrophoresis under nonreducing conditions as described before. VWF multimers were visualized using ECL Western Blotting Substrate (Promega, Madison, Wisconsin, United States).

VWF monomers were visualized by western blot under reducing conditions. Protein lysates were reduced using NuPAGE Sample Reducing Agent (Thermo Fisher Scientific) and proteins were separated on a 4 to 12% Bis-Tris gel (Thermo Fisher Scientific). Proteins were transferred to a PVDF membrane (BioRad, Veenendaal, the Netherlands) using a Trans-Blot Turbo Transfer System (BioRad, 1.5A, 15 minutes). Rabbit anti-VWF conjugated to horseradish peroxidase (DAKO, P0226) was used as the detection antibody. The membrane was incubated with the detection antibody diluted in PBS-tween containing 5% nonfat milk (Nutricia, Zoetermeer, the Netherlands). ECL Western Blotting Substrate was used to visualize VWF. Densitometry images were generated and quantified by ImageJ (ImageJ 1.51h, Bethesda, Maryland, United States).

RNA Analysis
RNA isolation, cDNA synthesis, and quantitative polymerized chain reaction (qPCR) analysis with GAPDH as the endogenous reference gene were performed as described before. qPCR was performed to quantify VWF, but also separate VWF alleles. Allele-specific qPCR primers were designed, containing the SNPs on the second last position of the forward primer. Primer sequences and annealing temperatures are shown in Supplementary Table S1 (available in the online version).

Immunofluorescence Staining
ECFCs were fixed using ice-cold methanol and stained as described before. Sheep anti-VWF (Abcam, ab11713), rabbit anti-protein disulphide isomerase (PDI; Stressgen Biotechnologies, San Diego, California, United States), and mouse anti-Golgi matrix protein of 130 kDa (GM130, BD Biosciences) were used as primary antibodies.

Statistical Analysis
Statistical analyses were performed using GraphPad Prism 8 (GraphPad Software, La Jolla, California, United States). Mann–Whitney U-tests were performed to determine significance between specific and aspecific inhibition. One-way ANOVA with Dunnett’s multiple comparisons test was used to determine significance between three or four groups. p < 0.05 was considered significant.

Results

Time Course of VWF Inhibition by siVWF in ECFCs
siRNAs are used in this study to inhibit VWF (alleles) in ECFCs. To assess the moment of strongest VWF inhibition on a protein level, but also the duration of VWF inhibition, we transfected ECFCs with an siRNA-targeting total VWF (siVWF) or siNEG and followed the VWF secreted in 24 hours for up to 28 days. Relative VWF inhibition by siVWF was determined by normalization of the VWF:Ag secreted in siVWF-treated cells over the VWF:Ag levels in siNEG-treated cells. The experiment was performed under different conditions, but for all tested conditions the strongest VWF inhibition was observed 6 days after siVWF transfection (Fig. 1 and Supplementary Fig. S1 [available in the online version]). Furthermore, only after 28 days the VWF levels in siVWF-transfected ECFCs were at the same level as siNEG-transfected ECFCs. Since the strongest effects were observed 6 days after transfection, we decided to consistently measure the effects of the siRNAs on the protein level, 6 days after transfection.

Allele-Specific siRNA Inhibition in Healthy Control ECFCs
We previously tested the efficiency and specificity of allele-specific siRNAs that target four common VWF SNPs in VWF-overexpressing HEK293 cells. For the current study, we selected the most effective allele-specific siRNA per SNP target from the previous study. To evaluate the efficiency and specificity of the selected siRNAs on a protein level in ECFCs, siRNAs were transfected in ECFCs that were homozygous for specific SNPs (Table 2). For all siRNAs tested, no or only minor reduction of the production of the untargeted VWF allele was observed as is shown by the relative VWF protein expression. The relative VWF protein expression was followed over time. The experiments under these conditions were performed three times in duplicate. Separate experiments are shown by different colored circles. Lowest relative VWF protein expression was observed 6 days after transfection. Similar results were observed under slightly different conditions (i.e., cell and DharmaFECT concentration; Supplementary Fig. 1). ECFC, endothelial colony-forming cell; nM, nanomolar; siNEG, negative control siRNA; siVWF, siRNA against VWF; VWF, von Willebrand factor; VWF:Ag, VWF antigen.
protein expression in ECFCs that did not harbor the SNP variant corresponding to the transfected siRNA (►Fig. 2A). On the other hand, a strong VWF knockdown of the targeted allele was observed in ECFCs that contained the SNP variant corresponding to the transfected siRNA (►Fig. 2A). Knockdown was especially strong and specific for si1451A and si1451G. Results of siRNA transfections with siRNA concentrations of 10 and 50 nM are shown in ►Supplementary Fig. S2 (available in the online version).

The allele-specific siRNAs have been designed with the aim to target heterozygous SNPs located on the same allele as a dominant-negative mutation and thereby improve disease phenotypes. However, the ability of the siRNAs to inhibit VWF alleles in heterozygous ECFCs can only be assessed on the RNA level. We therefore tested the allele-specific siRNAs in an ECFC line that is heterozygous for all four SNPs (ECFC C5; ►Table 2) and performed allele-specific qPCR to determine the relative RNA expression of either of the VWF alleles. On the RNA level, a stronger overall knockdown of VWF was observed compared with what was observed on the protein level, both of the targeted and untargeted alleles (►Fig. 2B). However, for most siRNAs clear specificity for its targeted allele remained. Only si2365A and si2365G showed minor specificity for its targeted allele. This is in line with results in HEK293 cells.17

Correction of a VWD Type 2A Multimerization Phenotype by an Allele-Specific siRNA

After proving that allele-specific siRNAs can discriminate between VWF alleles in healthy control ECFCs, we investigated

**Table 2** ECFCs used per siRNA for testing siRNA efficiency and specificity

| siRNA   | Protein analysis | RNA analysis             |
|---------|-----------------|--------------------------|
| si1451A | C3 (1451AA)     | C1 (1451GG)              |
|         | to test siRNA   | C5 (1451AG)              |
|         | efficiency      |                          |
| si1451G | C1 (1451GG)     | C3 (1451AA)              |
|         | to test siRNA   |                          |
|         | efficiency      |                          |
| si2365A | C3 (2365AA)     | C2 (2365GG)              |
|         | to test siRNA   | C5 (2365AG)              |
|         | efficiency      |                          |
| si2365G | C2 (2365GG)     | C3 (2365AA)              |
|         | to test siRNA   |                          |
|         | efficiency      |                          |
| si2385T | C3 (2385TT)     | C2 (2385CC)              |
|         | to test siRNA   | C5 (2385TC)              |
|         | efficiency      |                          |
| si2385C | C2 (2385CC)     | C3 (2385TT)              |
|         | to test siRNA   |                          |
|         | efficiency      |                          |
| si2880A | C4 (2880AA)     | C1 (2880GG)              |
|         | to test siRNA   | C5 (2880AG)              |
|         | efficiency      |                          |
| si2880G | C1 (2880GG)     | C4 (2880AA)              |

Abbreviations: ECFC, endothelial colony-forming cell; siRNA, small interfering RNA.
Note: Genotype is indicated between brackets.

**Fig. 2** Allele-specific inhibition of VWF in healthy control ECFCs. (A) Relative VWF protein expression of ECFCs transfected with allele-specific siRNAs. Shown are the total VWF:Ag levels (conditioned medium + protein lysates) measured in ECFCs transfected with specific siRNAs, normalized to the total VWF:Ag levels measured in the same ECFCs transfected with siNEG. Shown are the mean ± 1 SD of three independent experiments performed in duplicate. Mann–Whitney, ‘p < 0.05, ‘‘p < 0.01. (B) Relative VWF RNA expression of ECFC C5 transfected with allele-specific siRNAs. Shown are the RNA expression levels of VWF alleles of ECFC C5 transfected with specific siRNAs, normalized to the expression level of the same allele measured in ECFC C5 transfected with siNEG. Shown are the mean ± 1 SD of two independent experiments performed in duplicate. Mann–Whitney, ‘p < 0.05. ECFC, endothelial colony-forming cell; nM, nanomolar; qPCR, quantitative PCR; siNEG, negative control siRNA; SD, standard deviation; si1451G, indicates “small interfering RNA against VWF c.1451G,” all siRNAs are indicated according to this principle; VWF, von Willebrand factor; VWF:Ag, VWF antigen.
whether allele-specific VWF inhibition could also correct a VWD phenotype in ECFCs. ECFCs from a VWD type 2A patient with the VWF p.Cys1190Tyr mutation and reduced high molecular weight (HMW) multimers were successfully isolated (ECFC 2A; plasma phenotype in Table 1). Heterozygous SNPs are essential for allele-specific inhibition of mutant VWF, and genotyping of genomic DNA revealed that the patient is heterozygous for one of the four selected SNPs: VWF c.1451A\(\rightarrow\)G. PacBio long-read sequencing was performed to identify the phasing of this SNP, and showed that VWF c.1451A was located on the same allele as the dominant-negative mutation p.Cys1190Tyr (c.3569A) (Fig. 3A). Therefore, ECFCs should be transfected with si1451A to inhibit the mutant allele and correct for the VWD type 2A phenotype. Treatment of ECFC 2A with si1451G should reduce expression of the wild-type allele and is expected to deteriorate the phenotype.

Treatment of ECFC 2A with either si1451G or si1451A resulted in allele-specific VWF inhibition, as is shown by the allelic VWF expression determined in RNA, 48 hours after transfection (Fig. 3B). Inhibition of the mutant allele resulted in a 53 ± 6% reduction of total VWF secreted in the conditioned medium, 6 days after transfection. To assess whether allele-specific inhibition of mutant VWF resulted in phenotypic improvements, we first subjected protein lysates and conditioned medium to the VWF:CB assay, an assay that is able to detect VWF multimerization defects.21 As expected, we observed both in protein lysates and in conditioned medium of ECFC 2A a lower VWF:CB/VWF:Ag ratio compared to samples derived from healthy control ECFCs (Fig. 3C, D). This VWF:CB defect was almost corrected in protein lysates of ECFC 2A transfected with si1451A. Contrarily, inhibition of expression of wild-type VWF by si1451G clearly worsened the VWF:CB phenotype (Fig. 3C). Remarkably, in the conditioned medium of ECFC 2A transfected with si1451A, no correction of the VWF:CB defect was observed. Nevertheless, inhibition of the wild-type allele by si1451G did result in significantly lower VWF:CB/VWF:Ag as compared with siNEG-transfected ECFC 2A (Fig. 3D). We wondered why inhibition of mutant VWF did not result in correction of VWF:CB/VWF:Ag in the conditioned medium as it did in protein lysates. We hypothesized that downregulation of VWF in general leads to reduced VWF:CB/VWF:Ag in the conditioned medium. To assess this hypothesis, VWF:CB/VWF:Ag was determined in the conditioned medium and protein lysates of ECFC C1, C2, and C3 transfected with allele-specific siRNAs, siNEG, and siVWF. We observed in the conditioned medium, but not in protein lysates, that inhibition of VWF production by (allele-specific) siRNAs resulted in a gradual decrease of VWF:CB/VWF:Ag (n = 6 for each siRNA in control ECFCs; Fig. 3E, F). When the VWF:CB/VWF:Ag ratios of ECFC 2A transfected with siNEG, si1451A, or si1451G were plotted against the VWF:CB/VWF:Ag ratios of healthy control ECFCs, we observed that the VWF:CB/VWF:Ag ratio of ECFC 2A transfected with si1451A (inhibition of mutant allele) shifted toward the reference line of the healthy control ECFCs (n = 9 for ECFC 2A). Whereas inhibition of the normal allele by si1451G resulted in further deterioration of VWF:CB/VWF:Ag (in the conditioned medium as well as in protein lysates, Fig. 3E, F).

Although multimerization defects can be detected by the VWF:CB ELISA, the assay gives no information on the exact multimerization pattern. To assess the multimerization pattern, plasma, conditioned medium, and protein lysates were subjected to the VWF multimer analysis. In the conditioned medium, a slight decrease of HMW VWF was observed in the patient-derived ECFCs compared with control ECFCs (Fig. 3I). This was comparable to the multimerization defect observed in the patient’s plasma (Fig. 3H). Inhibition of mutant VWF resulted in a slight increase in HMW VWF.

Inhibition of production of wild-type VWF clearly worsened the multimerization pattern in the conditioned medium (Fig. 3I). No clear decrease of HMW VWF was observed in the protein lysates of ECFC 2A compared with the protein lysates of healthy control ECFCs. However, a clear increase in the intensity of the dimer band was apparent (Fig. 3G). Also, a slight change in the running pattern is visible, i.e., VWF from ECFC 2A seems to migrate slightly slower than the VWF from the control ECFC. After inhibition of mutant VWF by si1451A, the intensity of the dimer band clearly decreased; however, this coincided with a small decrease in HMW VWF. Furthermore, VWF migration shifted toward the pattern of the control ECFC. When the wild-type allele was inhibited by si1451G, the multimerization pattern deteriorated with a decrease in HMW VWF and an increase in the intensity of the dimer band.

**Improved VWF Processing after Allele-Specific Inhibition of Mutant VWF in ECFC 2A**

The multimerization pattern of protein lysates of ECFC 2A showed an increase in the intensity of the dimer band. This suggests that the processing of VWF is affected. Dimerization of VWF takes place in the endoplasmic reticulum (ER), after which VWF multimers are formed in the Golgi.22 To test whether VWF is retained in the ER, we performed a co-staining of VWF and ER marker PDI. Indeed, a clear overlap of VWF and PDI staining is observed in ECFC 2A transfected with siNEG, indicating ER retention (Fig. 4A; ECFC 2A + siNEG, second column). This was not observed in a control ECFC transfected with siNEG (Fig. 4A; control + siNEG, first column). No colocalization of VWF with the Golgi was observed, indicating pure ER retention for this mutation (Supplementary Fig. 3, available in the online version). When the mutant allele was inhibited by si1451A, retention of VWF in the ER was clearly decreased (best observed in the VWF grayscale image) and many cells showed no ER retention at all (Fig. 4A; ECFC 2A + si1451A, third column). Inhibition of wild-type VWF by si1451G resulted in a rather severe cellular phenotype, with increased ER retention (Fig. 4A; ECFC 2A + si1451G, fourth column).

VWF is produced as proVWF that is dimerized in the ER. After dimerization of proVWF in the ER, VWF is translocated to the Golgi where the propeptide is cleaved from VWF by furin.23 Increased ER retention of VWF results in an increase in unprocessed proVWF in the cells. This can be quantified by an ELISA that measures the amount of propeptide that is still bound to VWF. Protein lysates of ECFC 2A showed a 4.9-fold increase in unprocessed VWF compared with protein lysates of control ECFCs. This was a 1.9-fold decreased after inhibition
Fig. 3 Correction of the VWF multimerization defect in ECFC 2A. (A) Phasing of the heterozygous SNP, VWF c.1451A|G, with the dominant-negative mutation VWF p.Cys1190Tyr (c.3569A). PacBio sequencing revealed that VWF c.1451A is located on the same allele as VWF c.3569A. (B) Relative VWF RNA expression of ECFC 2A transfected with 20 nM si1451A or si1451G. Shown are the RNA expression levels of VWF alleles c.1451A and c.1451G as well as the total VWF expression. Expression levels were determined in RNA lysates of ECFC 2A transfected with si1451A or si1451G, normalized to the expression level of the same allele measured in ECFCs transfected with siNEG. Shown are the mean ± 1 SD of three independent experiments performed in triplicate. Mann–Whitney (c.1451A vs. c.1451G expression), ****p < 0.0001. VWF:CB/VWF:Ag determined in (C) protein lysates and (D) conditioned medium of ECFC C1, C2, and C3, transfected with siNEG and ECFC 2A transfected with siNEG, si1451A, and si1451G. Shown are the mean ± 1 SD of three independent experiments performed in triplicate for ECFC 2A and the mean ± 1 SD of three independent experiments performed in duplicate for ECFC C1, C2, and C3. One-way ANOVA, *p < 0.05, **p < 0.001, ***p < 0.0001.

Normal reference line of VWF:CB/VWF:Ag plotted against the VWF:Ag levels measured in (E) protein lysates and (F) conditioned medium of ECFC C1, C2, and C3 transfected with siNEG, siVWF, and various allele-specific siRNAs. Every white dot represents the average of three independent experiments performed in duplicate of an ECFC line transfected with a specific siRNA. Included in the graphs are the average of VWF:CB/VWF:Ag of three experiments performed in triplicate in which ECFC 2A was transfected with si1451A (green), si1451G (purple), or siNEG (red). VWF multimerization analysis of (G) protein lysates, (H) plasma, and (I) conditioned medium. Protein lysates and conditioned medium were harvested from a healthy control ECFC line transfected with siNEG and ECFC 2A transfected with si1451A, si1451G, and siNEG. ECFC, endothelial colony-forming cell; qPCR, quantitative PCR; siNEG, negative control siRNA; si1451G, indicates “small interfering RNA against VWF c.1451G,” all siRNAs are indicated according to this principle; VWF, von Willebrand factor; VWF:Ag, VWF antigen; VWF:CB, VWF collagen binding.
of production of the mutant allele by si1451A (►Fig. 4B).

Compared with control ECFCs, sixfold more unprocessed VWF was present in ECFC 2A in which the wild-type allele was inhibited by si1451G.

A western blot in which the samples are run under reducing conditions can be used as an alternative method to identify defects in the processing of VWF. In this western blot, an increase in the intensity of the unprocessed proVWF band was detected in protein lysates of ECFC 2A compared with control ECFCs (►Fig. 4C). The intensity of the unprocessed proVWF band decreased after inhibition of the mutant allele by si1451A. An increase in the intensity of the unprocessed proVWF band was observed when the wild-type allele was inhibited by si1451G. Quantification of the western blot confirms the defective processing of VWF and improvement of the phenotype after transfection of si1451A that inhibited expression of the mutant allele (►Fig. 4D).

**Discussion**

Allele-specific siRNAs can be used to selectively inhibit mutant alleles to improve disease phenotypes. Previously, we have shown that allele-specific siRNAs that distinguish two VWF alleles based on one nucleotide mismatch can specifically inhibit VWF alleles in HEK293 cells, and that these siRNAs can improve a VWD phenotype. Here, we aimed to extend the proof of concept of allele-specific VWF inhibition to an ex vivo setting by the use of patient-derived and healthy control ECFCs. In this study, we prove that allele-specific siRNAs can inhibit single VWF alleles based on one nucleotide mismatch in healthy control ECFCs. This was shown both on protein and RNA levels. Also, inhibition of the mutant allele in ECFCs isolated from a VWD type 2A patient resulted in clear improvements in the patient's cellular phenotype (►Fig. 5).
SNPs in VWF with a high minor allele frequency have been used in this study as a target to inhibit VWF alleles. This approach was chosen since with only a small number of SNPs, a large percentage of the patient population can be reached. From this study it became evident that especially si1451A and si1451G have high potency as allele-specific siRNAs, and both siRNAs proved successful to, respectively, correct or deteriorate a VWD type 2A phenotype ex vivo. Whether the other siRNAs have the same ability to correct for VWD phenotypes remains to be unanswered. It is however unlikely that si2365A or si2365G have similar effects as si1451A or si1451G. Fortunately, exclusion of VWF c.2365A|G does not reduce the fraction of the targeted patient population, since VWF c.2365A|G is in almost complete linkage disequilibrium with VWF c.2385C|T. Furthermore, the efficacy of the allele-specific siRNAs in this study may be improved by alternating the chemical modification of the siRNAs. Also, more SNP targets could be tested to increase the percentage of the targeted patient population.

The ability of the allele-specific siRNAs to correct a VWD phenotype was tested in ECFCs isolated from a VWD type 2A patient with the VWF p.Cys1190Tyr mutation. This mutation is characterized by a clearly defined laboratory phenotype with reduced HMW multimers and reduced VWF:CB. Inhibition of mutant VWF by si1451A resulted in clear improvements in VWF processing and the VWF:CB/VWF:Ag in protein lysates. Remarkably, no increase of VWF:CB/VWF:Ag was observed in the conditioned medium of ECFC 2A in which expression of the mutant allele was inhibited (►Fig. 3D). It appeared, however, that also a reduced VWF:CB/VWF:Ag was observed in healthy control ECFCs transfected with siRNAs that reduced the overall expression of VWF (►Fig. 3F). This decrease in VWF:CB/VWF:Ag might be a result of the formation of shorter Weibel–Palade bodies after inhibition of VWF, as was observed by Ferraro et al. When the VWF:CB/VWF:Ag of ECFC 2A transfected with si1451A was plotted against the normal reference line, VWF:CB/VWF:Ag was almost normalized. Also, the multimerization pattern of VWF in the conditioned medium showed an increase in HMW VWF in ECFCs transfected with si1451A compared with the siNEG-transfected cells. The reason for discrepancy between the VWF:CB assay and the VWF multimer analysis remains uncertain. It is possible that the VWF:CB assay is sensitive for altered ratios between HMW and low-molecular-weight VWF. Indeed, the decrease in VWF:CB after inhibition is undesirable. Further in vivo studies should reveal whether reduced VWF:CB after downregulation of VWF is also apparent in vivo and what the potential consequences of this effect are.

ECFCs are used in this study as an ex vivo cell model for VWD. ECFCs are the only source of proliferative endothelial cells that can be isolated from patients by a peripheral blood venepuncture. The use of ECFCs has the advantage that the cells are proliferative, have generally a high VWF production, and are easy to transfect. However, also clear variations between various ECFC cell lines emerge. Most importantly, clear variations in VWF expression exist between ECFC cell lines and it was therefore suggested to be cautious when comparing different ECFC lines. When allele-specific siRNAs are tested in (patient-derived) ECFCs, the effects are determined in ECFCs transfected with either an allele-specific siRNA or a negative control siRNA. Any effects of the allele-specific siRNA compared with the negative control siRNA are an effect of the siRNA and not due to potential variability of the ECFC, as the cells are used as their own control. Therefore, we are confident to draw conclusions on the effects of allele-specific siRNAs on ECFCs from a single patient. To assess the efficiency and specificity of the allele-specific siRNAs on a protein level, we had no
choice but to compare different healthy control ECFC lines (►Fig. 2A). Although the ECFC lines in this study were carefully selected based on their proliferative state and VWF production, differences between ECFC lines could not be avoided. This was also reflected in variations in the efficiency of siVWF to inhibit VWF production in ECFCs. For example, we reproducibly observed a stronger relative VWF inhibition by siVWF in ECFC C1 compared with ECFC C2 (data not shown). No firm conclusions can therefore be drawn on the efficiency and specificity of the allele-specific siRNAs on a protein level that have been tested in the different healthy control ECFC lines (►Fig. 2A).

The use of siRNAs as a therapeutic application requires persistence of the siRNA activity over a longer period of time. Clinical trials with siRNAs that target liver-expressed genes showed persistent siRNA-mediated downregulation of several genes for more than a month.30–32 In ECFCs, downregulation of VWF was observed for up to 28 days. Furthermore, siRNA-mediated inhibition of Icam2 and Tie2 in mouse endothelium by siRNAs that were complexed in polymeric nanoparticles or a cationic lipoplex delivery system, respectively, resulted in persistent, and around 80%, Icam2 and Tie2 inhibition for more than 21 days after a single dose.33,34 These results of long-term gene inhibition are promising for further developments of siRNA therapeutics for endothelial genes, like VWF.

To conclude, we show that allele-specific siRNAs are effective in the inhibition of single VWF alleles in ECFCs. Inhibition of expression of the mutant allele in ECFCs isolated from a VWD type 2A patient with the VWF p.Cys1190Tyr mutation resulted in clear improvements of the cellular phenotype. Further studies in VWD mouse models are needed to translate the positive ex vivo results to an in vivo model, and to show whether the allele-specific siRNAs are able to correct a bleeding phenotype.

What is known about this topic?

• Continued circulation of mutant VWF negatively affects hemostasis in VWD patients.
• Most VWDs are caused by dominant-negative mutations.
• siRNAs can discriminate between two alleles with only one nucleotide mismatch and can therefore inhibit mutant alleles.

What does this paper add?

• Inhibition of VWF in ECFCs by siRNAs results in reduced VWF levels up to 21 days in cultured endothelial cells.
• siRNAs allow allele-specific inhibition of VWF alleles in ECFCs.
• siRNA-mediated inhibition of mutant VWF in patient-derived ECFCs results in clear improvements in the cellular phenotype.

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Authors’ Contributions

A.D.J. designed the study, performed the experiments and data analyses, and wrote the manuscript. R.J.D. performed the experiments. J.B., F.A., and F.W.G.L. were involved in the control and patient inclusions. S.Y.A. contributed to the analysis of PacBio sequencing data. B.J.M.v.V. contributed to discussions and reviewing of the manuscript. J.E. designed the study, interpreted the data, and contributed to writing of the manuscript. All authors revised and approved the final version of the manuscript.

Conflict of Interest

J.B. received research funding from CSL Behring and is an employee of Sobi. F.A. received research funding from CSL Behring and a travel grant from Sobi. F.W.G.L received research funding from CSL Behring, Takeda/Shire, and uniQure. He is member of a DSMB for Roche. He is consultant for Takeda, Biomarin, and uniQure of which fees go to the institution. J.E. received research funding from CSL Behring. J.E. reports grants from Landsteiner Foundation for Blood Transfusion Research, during the conduct of the study. Other authors declare no conflicts of interest.

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