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# Macrothrombocytopenia and dense granule deficiency associated with *FLI1* variants: ultrastructural and pathogenic features

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## ABSTRACT

Congenital macrothrombocytopenia is a family of rare diseases, of which a significant fraction remains to be genetically characterized. To analyze cases of unexplained thrombocytopenia, 27 individuals from a patient cohort of the Bleeding and Thrombosis Exploration Center of the University Hospital of Marseille were recruited for a high-throughput gene sequencing study. This strategy led to the identification of two novel *FLI1* variants (c.1010G>A and c.1033A>G) responsible for macrothrombocytopenia. The *FLI1* variant carriers' platelets exhibited a defect in aggregation induced by low-dose adenosine diphosphate (ADP), collagen and thrombin receptor-activating peptide (TRAP), a defect in adenosine triphosphate (ATP) secretion, a reduced mepacrine uptake and release and a reduced CD63 expression upon TRAP stimulation. Precise ultrastructural analysis of platelet content was performed using transmission electron microscopy and focused ion beam scanning electron microscopy. Remarkably, dense granules were nearly absent in the carriers' platelets, presumably due to a biogenesis defect. Additionally, 25-29% of the platelets displayed giant  $\alpha$ -granules, while a smaller proportion displayed vacuoles (7-9%) and autophagosome-like structures (0-3%). *In vitro* study of megakaryocytes derived from circulating CD34<sup>+</sup> cells of the carriers revealed a maturation defect and reduced proplatelet formation potential. The study of the *FLI1* variants revealed a significant reduction in protein nuclear accumulation and transcriptional activity properties. Intraplatelet flow cytometry efficiently detected the biomarker MYH10 in *FLI1* variant carriers. Overall, this study provides new insights into the phenotype, pathophysiology and diagnosis of *FLI1* variant-associated thrombocytopenia.

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## Introduction

Congenital reduced platelet count or impaired platelet function lead to a family of diseases, which are increasingly recognized as a significant cause of bleeding in children and adults.<sup>1</sup> Congenital macrothrombocytopenias (CMTPs) constitute a large subgroup of inherited thrombocytopenia characterized by enlarged platelets.

Several genetic variants have been associated with CMTP, of which the most frequent alter genes that encode platelet surface proteins and megakaryocyte (MK) cytoskeletal proteins.<sup>2</sup>

However, mounting evidence has shown that several forms of CMTP are caused by variants in genes encoding hematopoietic transcription factors, which lead to altered downstream expression of genes that regulate platelet formation and function.<sup>3-10</sup> In 2013, Stockley *et al.* used next-generation sequencing (NGS) to analyze 13 unrelated patients suspected of having an inherited qualitative platelet defect in the UK Genotyping and Phenotyping of Platelets (UK-GAPP) study.<sup>7</sup> The group reported two substitutions and a 4-bp frameshift deletion in the transcription factor gene *FLI1*. Two of the index cases also had mild thrombocytopenia, and one of them exhibited enlarged platelets. These findings highlight the role of *FLI1* in megakaryopoiesis and platelet function.

To analyze cases of unexplained thrombocytopenia, 27 individuals from the patient cohort of the Bleeding and Thrombosis Exploration Center of the University Hospital of Marseille were recruited for a high-throughput gene sequencing investigation. In the study herein, we report the discovery of two novel *FLI1* dominant variants linked to CMTP in two unrelated pedigrees. We investigated the pathogenesis of *FLI1*-associated thrombocytopenia by analyzing subcellular *FLI1* localization and MK differentiation from hematopoietic progenitors derived from patients. We characterized the genotype-phenotype relationship and described the intraplatelet flow cytometry-based quantification of MYH10, a biomarker of *FLI1* and *RUNX1* alterations. Focused ion beam scanning electron microscopy (FIB-SEM) is an imaging approach based on the serial sectioning of entire platelets, which allows a 3D representation of the granules with respect to their shape, content and number. Using this method, we showed a severe dense granule deficiency in platelets of the *FLI1* variant carriers.

**Methods**

Methods concerning high-throughput gene sequencing, structural model of *FLI1*-DNA interactions, platelet phenotyping,

platelet-rich plasma (PRP) serotonin level, ATP secretion, mepacrine uptake and release, flow cytometric quantification of MYH10 expression (whole blood-based assay), western blot assay, site-directed mutagenesis, luciferase reporter assay, epifluorescence microscopy and *in vitro* MK differentiation and pro-platelet formation are described in the *Online Supplementary Material*.

**Recruitment of patients and genetic analysis strategy**

Patients were recruited from the cohort of the Bleeding and Thrombosis Exploration Center (University Hospital of Marseille) after informed written consent was obtained, in accordance with protocols approved by the local Institutional Review Board and the Declaration of Helsinki. Two high-throughput sequencing methods were used to identify candidate variants: (A) Whole-exome sequencing performed at the Centre National de Génotypage (Evry, France), or (B) sequencing of a 308 gene panel (selected genes associated with platelet diseases; available on request).

**Flow-cytometric quantification of MYH10 expression (PRP-based assay)**

PRP was fixed in 1% paraformaldehyde phosphate buffered saline (PBS). Next, the fixed PRP was centrifuged at 1000 G for five min. The platelet pellet was suspended in permeabilization buffer (0.5% Triton X-100, 2mM Ethylenediaminetetraacetic acid, 0.5% bovine serum albumin (BSA) PBS) and labeled with rabbit anti-MYH10 antibody (Cell Signaling; #3404) for one hour. After a washing step, the platelets were incubated with goat anti-rabbit Alexa-488-labeled secondary antibody (Abcam; ab150077) for 30 min in permeabilization buffer. After another washing step, the data were acquired using Navios Cytometer (Beckman Coulter) and analyzed with FlowJo software (Tree Star, Inc.).

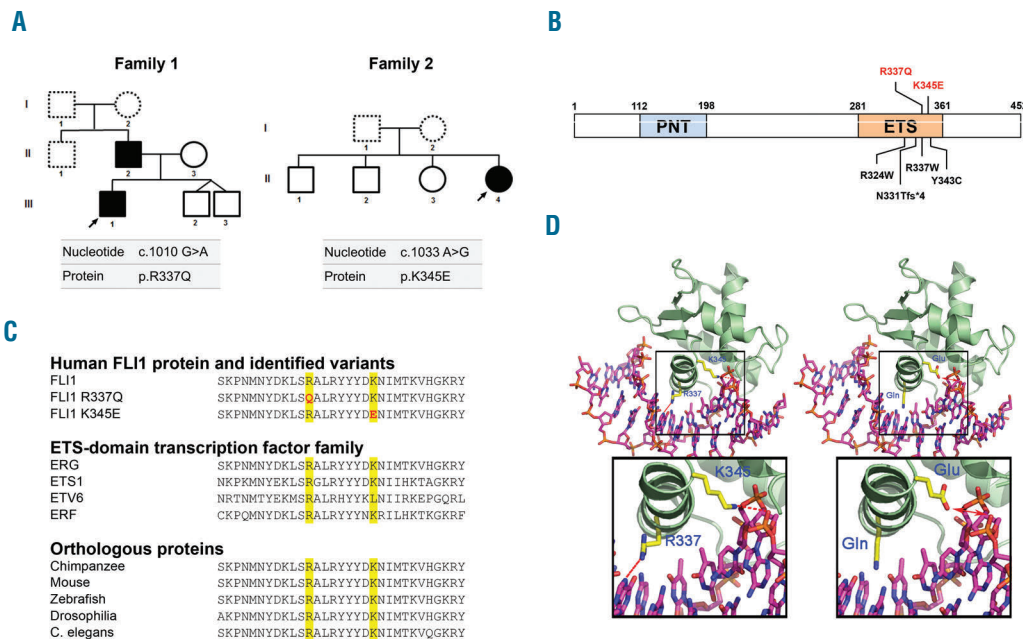
**Electron microscopy**

For ultrastructural analysis, platelets in citrated PRP were diluted and fixed in PBS pH 7.2 containing 1.25% glutaraldehyde for one hour. After centrifugation and two PBS washings, they were post-fixed in 150 mM cacodylate-hydrochloric acid (HCl) buffer containing 1% osmium tetroxide pH 7.4 for 30 min at 4°C. After dehydration in a graded alcohol, embedding in EPON was performed by polymerization at 60°C for 72 hours. Ultrathin sections (~70 nm thick) were mounted on 200 mesh copper grids, contrast-

**Table 1. Platelet phenotyping in patients carrying the *FLI1* variants.**

Patient ID	<i>FLI1</i> variant	Age <sup>†</sup> (years)	Platelet counts (x10 <sup>9</sup> /l)	MPV <sup>‡</sup> (fl)	Serum PAI1 levels (ng/ml)	PRP serotonin levels (µg/10 <sup>9</sup> plt)	Platelet aggregation maximal intensity (%)			Platelet glycoprotein MFI <sup>§</sup> (a.u.)			
							ADP	Coll* <sup>¶</sup>	AA* <sup>¶</sup>	αIIbβ3	GP1bα	CD63	P-selectin
Reference ranges <sup>§</sup>			152-402	7.1-9.6	92-283	0.30-1.2	2.5 µM:75-95	78-90	79-94	Base: 20-47	Base: 1.3-5.1	Base: 0.4-1.3	Base: 0.2-0.8
							5.0 µM:78-91			TRAP: 28-60	TRAP: 0.7-2.2	TRAP: 1.3-4.2	TRAP: 1.9-7.4
F1-II2	c.1010 G>A (p.R337Q)	46	154	10.7	430	0.26	NA	NA	NA	Base: 21	Base: 1.2	Base: 0.6	Base: 0.3
										TRAP: 30	TRAP: 0.9	TRAP: 0.8	TRAP: 1.7
F1-III1	c.1010 G>A (p.R337Q)	19	131	13.0	443	NA	2.5 µM:31	83	85	Base: 34	Base: NA	Base: 0.7	Base: NA
							5.0 µM:81			TRAP: 42	TRAP: NA	TRAP: 1	TRAP: NA
F2-II4	c.1033 A>G (p.K345E)	52	140	11.1	273	0.22	2.5 µM:58	81	83	Base: 22	Base: 2.9	Base: 0.9	Base: 0.6
							5.0 µM:76			TRAP: 30	TRAP: 2.9	TRAP: 1.3	TRAP: 5

<sup>†</sup>Age at last evaluation. <sup>‡</sup>MPV: mean platelet volume (measured by optical method using ADVIA 120, Siemens). <sup>§</sup>The reference ranges were defined as the [minimum-maximum] interval of values obtained from healthy individuals in our laboratory. <sup>¶</sup>The collagen and arachidonic acid aggregation assays were performed using agonist concentrations of 3.3 µg/mL and 0.5 mg/mL, respectively. <sup>¶</sup>Platelet glycoprotein expression was measured via flow cytometry at basal conditions and after stimulation using TRAP-14 (50 µM). PAI1: plasminogen activator inhibitor-1; PRP: platelet-rich plasma; plt: platelet; ADP: adenosine diphosphate; Coll: collagen; AA: arachidonic acid; a.u.: arbitrary unit; NA: not available; TRAP: thrombin receptor-activating peptide; MFI: mean fluorescence intensity.



**Figure 1. Identification of two novel *FLI1* variants.** (A) Pedigrees for the affected families. Squares denote males and circles denote females. Black filled symbols represent family members carrying heterozygous c.1010 G>A and c.1033 A>G *FLI1* variants. Dotted line symbols represent non-tested members. Arrows indicate the probands. (B) Schematic diagram of the FLI1 protein. The functional N-terminal Pointed domain (PNT), and C-terminal ETS DNA-binding domain (ETS) are depicted. The positions of the alterations in FLI1 are indicated in red (alterations reported in this study) or black (previously reported alterations).<sup>7,8</sup> (C) Sequence alignment of the FLI1 protein (variant NM\_002017.3). The variants reported in this study are indicated in red (top). Alignments of various members of the ETS-domain transcription factor family (middle) and different species (bottom) are provided. (D) Diagram of the simulated interactions between FLI1 and double-stranded DNA (dsDNA). Left: Interaction of native FLI1 with dsDNA. The FLI1 dsDNA-interacting domain is represented as a green ribbon. The dsDNA is represented as a stick model (carbons: pink; nitrogen: blue; oxygen: red; phosphate: orange). The interactions of R337 with Guanine 5 atoms N7 and O6 as well as K345 with Adenine 15 atom OP2 (side chain carbons: yellow; nitrogen: blue; oxygen: red) are indicated in red. Right: model of FLI1 structure with mutated residues p.R337Q and p.K345E. An expanded view of the interaction between altered residues and DNA is shown at the bottom.

ed with uranyl acetate and lead citrate and examined using a JEOL JEM1400 transmission electron microscope equipped with a Gatan Orius 600 camera and Digital Micrograph software (Lyon Bio Image, Centre d'Imagerie Quantitative de Lyon Est, France). Morphometric measurements were done using ImageJ software (National Institutes of Health, USA). Preparations of platelet whole mounts were obtained after brief contact of formvar coated grids on small drops of PRP. Then, grids were rinsed rapidly with distilled water, dried with the edge of a filter paper, waved in the air and examined under the transmission electron microscope. Dense granules were identified and counted as previously described.<sup>11</sup> FIB-SEM samples were processed as for transmission electron microscopy (TEM), and examined using a Helios NanoLab microscope (FEI). The 3D models were computed using Amira software.

### Statistical analyses

Statistical significance was determined *via* a 2-tailed Mann-Whitney test unless otherwise specified.  $P < 0.05$  was considered statistically significant. Quantitative variables were presented as mean  $\pm$  standard error of the mean. Analyses were performed using GraphPad Prism software.

## Results

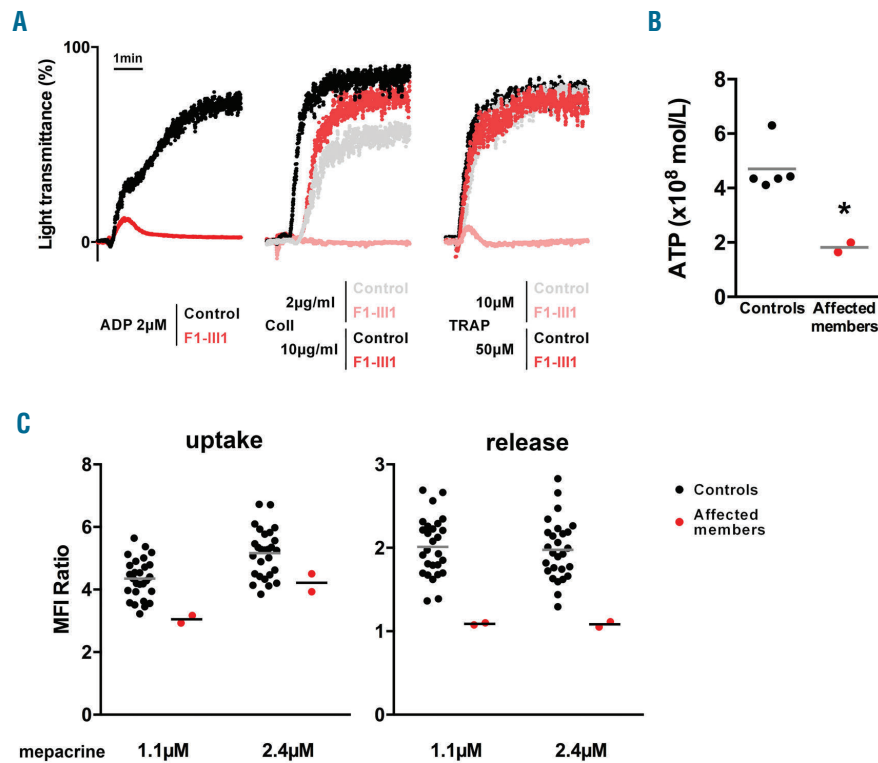
### Identification of two novel *FLI1* variants

Two high-throughput sequencing methods were used to identify candidate variants in 27 patients with suspected congenital thrombocytopenia. DNA samples from 12 individuals (three members of four unrelated families)

with unexplained autosomal dominant macrothrombocytopenia were analyzed *via* whole-exome sequencing. In one family (Family F1), this analysis revealed a heterozygous single nucleotide change in *FLI1*. This missense variant, c.1010G>A, encodes an arginine to glutamine change (p.R337Q) in the highly conserved E twenty-six (ETS) DNA-binding domain of FLI1 (Figure 1A,B). The presence of the variant was confirmed using capillary sequencing. There was a segregation between the *FLI1* variant and a history of thrombocytopenia in all cases for which DNA samples were available. In the three other families, no candidate variant was detected. Next, 15 additional individuals from 14 unrelated families were screened using a 308 gene panel NGS strategy. This screening led to the genetic diagnosis of four patients with macrothrombocytopenia. Aside from the identification of *MYH9*, *ACTN1* and *GP1BA* variants in three unrelated individuals, one patient carried c.1033A>G heterozygous *FLI1* variant, yielding p.K345E substitution (Family F2). In this family, the clinical and genetic status of the proband's father (F2-I1) was unknown and the proband's mother (F2-I2) displayed a normal platelet count, but the *FLI1* gene was not sequenced. This second *FLI1* variant also affected the ETS domain (Figure 1A,B). Sequence alignment of FLI1 orthologs and paralogs demonstrated that both variants affected amino acid residues in a highly conserved sequence (Figure 1C).

As these variants were located in the ETS domain, a simulation of the protein-DNA interactions was performed. Regarding the p.R337Q variant, glutamine 337





**Figure 2. Platelet function analysis.** (A) Light transmission aggregometry upon ADP (2 µM), collagen (2 and 10 µg/ml) and TRAP-6 (10 and 50 µM) stimulation in a *FLI1* variant carrier (F1-III1) and in a control who was representative of 20 controls investigated at the same period of time. The PRP platelet count was 268 x10<sup>9</sup>/l for the patient and 331 x10<sup>9</sup>/l for the control. (B) Luminometry-based ATP secretion assay. ATP secretion was measured in two *FLI1* variant carriers (F1-II2 and F1-III1) and unrelated controls after 100 µM TRAP-6 stimulation in 100 µl of diluted PRP (10<sup>7</sup> platelet / ml). \*P<0.05 vs. controls (Mann-Whitney test). (C) Flow cytometric mepacrine uptake and release assay in two *FLI1* variant carriers (F1-II2 and F2-II4) and unrelated controls. The platelets were incubated with 1.1 or 2.4 µM mepacrine and stimulated with 40 µM TRAP-14 to evaluate the mepacrine release. The mepacrine uptake was defined as the MFI ratio of platelets incubated with mepacrine to platelets incubated without mepacrine, and the mepacrine release was defined as the MFI ratio of resting platelets to stimulated platelets. ADP: adenosine diphosphate; ATP: adenosine triphosphate; TRAP: thrombin receptor-activating peptide; Coll: collagen; MFI: mean fluorescence intensity.

retained the ability to interact with DNA; however, the putative hydrogen (H)-bonds should be weaker than those established with the positively charged arginine guanidinium moiety. In the p.K345E variant, glutamic acid 345 is negatively charged and likely has a repelling effect on DNA binding (Figure 1D).

**Description of the families**

The affected members of the first family (F1-II2 and F1-III1) were both males, aged 46 and 19, respectively. They had not experienced significant bleeding (International Society on Thrombosis and Haemostasis Bleeding Assessment Tool (ISTH BAT) score: 1 for both), whereas the female proband in the second family (F2-II4), who was 52 years of age, had displayed excessive bleeding (ISTH BAT score: 16) with predominantly gynecological, obstetrical and oral cavity hemorrhaging. The three affected members did not display any extra hematologic abnormality. The first-line laboratory tests are reported in Table 1. All affected members presented with mild thrombocytopenia (mean value: 142 ± 7 x 10<sup>9</sup>/l). The mean platelet volume (MPV) was elevated (11.6 ± 1.2 fl; reference range 7.1 - 9.6 fl). In these families, coagulation, fibrinolysis and von Willebrand factor levels were normal (*data not shown*).

**Platelet receptor expression and platelet function evaluation**

The CD63 expression did not increase significantly upon TRAP stimulation (Table 1). The basal expression of αIIbβ3 was in the normal range among *FLI1* variant carriers (Table 1). The platelet light transmission aggregometry (Figure 2A and Table 1) revealed a significant aggregation defect upon stimulation with different agonists, when used at a low concentration (2 and 2.5 µM ADP, 2 µg/ml

collagen, 10 µM TRAP-6). The use of higher concentrations of agonists (5 µM ADP, 10 µg/ml collagen, 50 µM TRAP-6) induced normal platelet aggregation. The ATP secretion upon TRAP-6 stimulation was 2.5-fold decreased in patients F1-II2 and F1-III1 as compared to controls (Figure 2B). The mepacrine uptake and release upon TRAP-14 stimulation was reduced (Figure 2C). Among *FLI1* variant carriers, the platelet serotonin level was low but the serum plasminogen activator inhibitor-1 (PAI-1) antigen level (an α-granule biomarker) was normal or high (Table 1).

**Increased MYH10 expression in the platelets of affected members**

The expression of MYH10 in platelets has been recently proposed as a biomarker for *RUNX1* and *FLI1* alterations.<sup>7,8,12</sup> Intraplatelet flow-cytometry showed an almost 4-fold overexpression of MYH10 among the *FLI1* variant carriers as compared to controls, using a PRP-based and a whole blood-based experimental procedure (Figure 3A and *Online Supplementary Figure S1*, respectively). The MYH10 mean fluorescence intensity (MFI) was expressed as a function of MPV in 13 control individuals (MPV range 7.2 - 11.4 fl) and two *FLI1* patients (MPV 10.7 and 13 fl), demonstrating that the increased MPV could not explain the specific increase in MYH10 MFI observed in the *FLI1* patients (*Online Supplementary Figure S1*). Western blot analysis confirmed the increased expression of MYH10 in the platelet lysates of the *FLI1* variant carriers (Figure 3B).

**Functional assessment of the *FLI1* variants**

Western blot analysis did not reveal an alteration in *FLI1* protein expression in the platelets of affected members compared with healthy controls (Figure 4A). Accordingly, protein expression of wild-type (WT) and both *FLI1* vari-

ants in transfected cells were equivalent, thereby suggesting that these variants do not influence protein expression or stability (Figure 4B). To investigate the effect of the p.R337Q and p.K345E variants on FLI1 transcriptional activity, we investigated the ability of the recombinant FLI1 variants to regulate transcriptional activity using a dual-luciferase reporter assay. Co-transfection of the reporter plasmid containing the ETS-binding site and a plasmid encoding WT FLI1 resulted in almost 70% inhibition of luciferase activity. There was a loss in the ability of the p.R337Q and p.K345E variants to repress luciferase activity compared with that of WT FLI1 ( $P < 0.0001$ , Figure 4C). Reduced transcriptional activity was also observed in this model for the FLI1 p.R324Q variant previously reported by Stevenson *et al.*<sup>8</sup> The co-transfection of WT and variant *FLI1* led to normal transcriptional activity (Figure 4C).

The R337 and K345 residues were predicted to be part of a nuclear localization signal sequence using several prediction tools (cNLS mapper and NLStradamus; *data not*

*shown*). To examine whether the p.R337Q and p.K345E variants alter the subcellular localization of FLI1 protein, we performed cell fractionation assay and immunofluorescence staining of cells overexpressing WT or variant FLI1. Western blot analysis of subcellular fractions showed increased FLI1 protein level in the cytoplasmic fraction and decreased protein level in the nuclear fraction in cells expressing the p.R337Q and p.K345E FLI1 variants compared with cells expressing the WT protein (Figure 4D). Concordant with the western blot data, fluorescence microscopy showed that WT FLI1 concentrated primarily in cell nuclei. In contrast, both *FLI1* variants exhibited predominant cytoplasmic localization (Figure 4E).

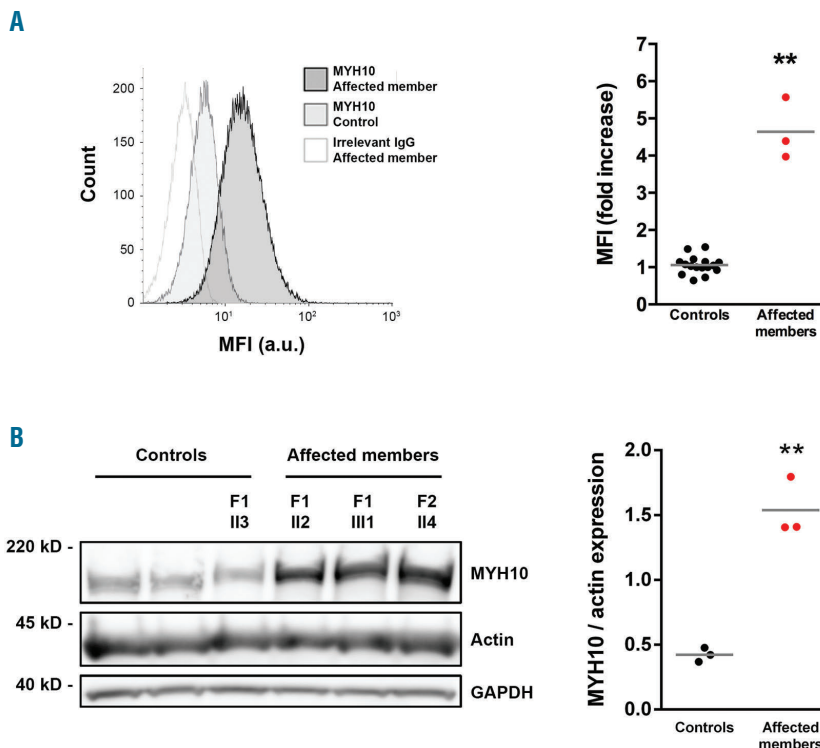
**FLI1 variants are associated with reduced CD34<sup>+</sup>-derived megakaryocyte differentiation and proplatelet formation *in vitro***

We generated MKs from peripheral blood CD34<sup>+</sup> cells in the presence of thrombopoietin (TPO) and stem cell factor

**Table 2. FIB-SEM analysis of platelet ultrastructure.**

Patient ID	FLI1 variants	Platelet volume (fl)	$\alpha$ -granule volume (fl)	Giant $\alpha$ -granule volume (fl)	Proportion of platelets displaying			
					Giant $\alpha$ -granules (%)	Autophagosome-like structures (%)	Glycogen accumulation (%)	Vacuoles (%)
Control		7.6±0.5	0.018±0.003	None detected	None detected	None detected	8	None detected
F1-II2	c.1010 G>A (p.R337Q)	12.6±0.1	0.014±0.005	0.22±0.02	25	3	86	9
F2-II4	c.1033 A>G (p.K345E)	12.8±2.0	0.017±0.005	0.08±0.02	29	None detected	51	7

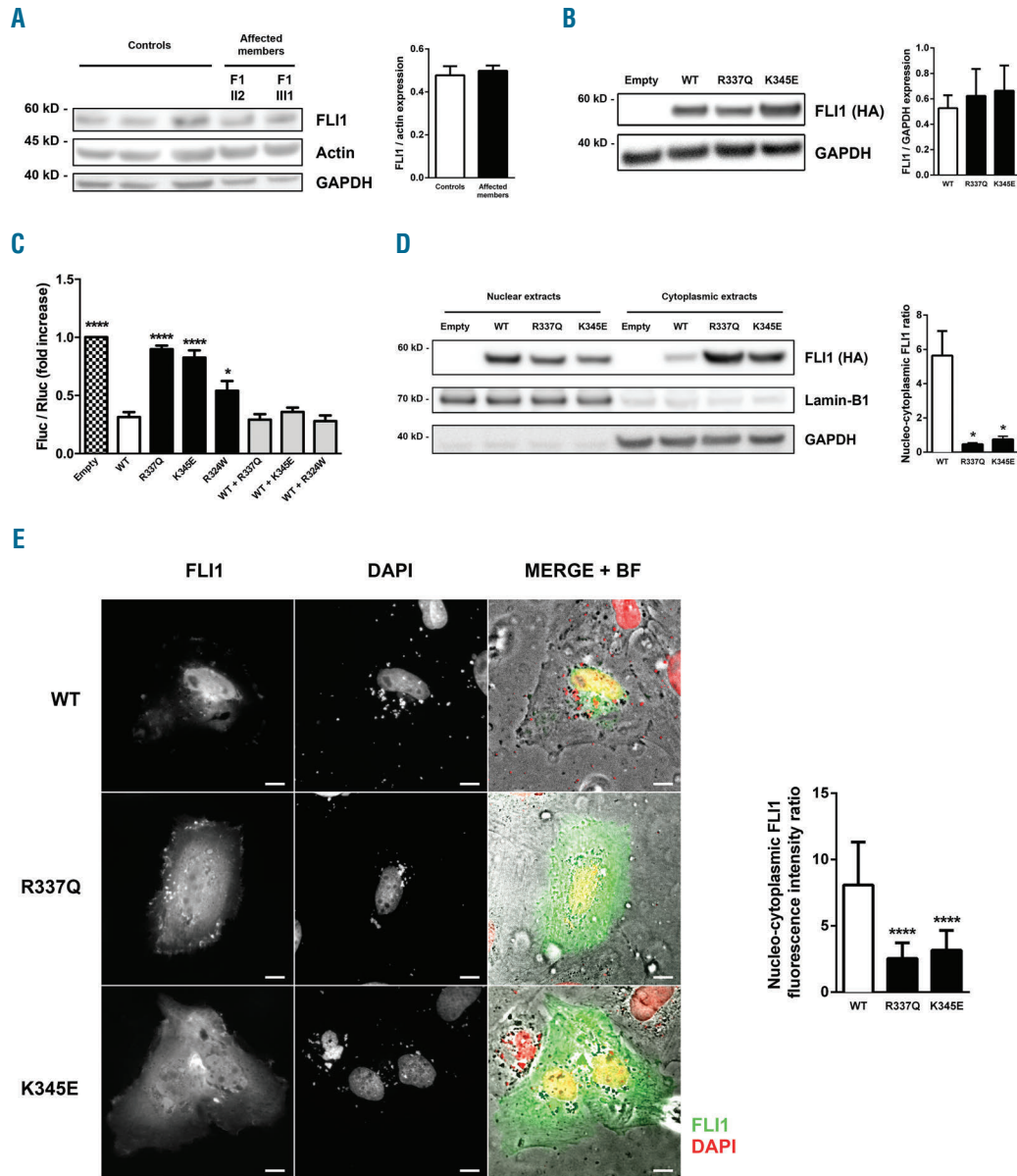
Giant  $\alpha$ -granules were defined by a diameter of >400 nm. Values are shown as the mean ± SEM as quantified for n=38, 52 or 45 randomly selected platelets for control, F1-II2 and F2-II4, respectively.



**Figure 3. Quantification of platelet MYH10 expression.** (A) Flow cytometry-based quantitative detection of intraplatelet MYH10 expression (PRP-based assay). Left: representative overlay of histograms of intraplatelet MYH10 expression in an affected member (F1-II2) and a healthy control. The non-specific staining (irrelevant IgG) is only presented for the affected member, which can be superimposed for that of the control. Right: MYH10 mean fluorescence intensity (MYH10 MFI - irrelevant IgG MFI) in affected members (n=3) or unrelated controls (n=8) from three independent experiments. The results are expressed as fold change relative to corresponding controls; \*\* $P < 0.01$  vs. controls (Mann-Whitney test). (B) Left: representative western blot analysis of MYH10 expression in platelets from the affected members (F1-II2, F1-III1, F2-II4), one unaffected member (F1-II3) and two unrelated controls. GAPDH and actin were used as a protein loading control. Right: the results of densitometric analysis were normalized to actin and expressed as mean ± SEM; \*\* $P < 0.01$  vs. controls (Mann-Whitney test). Three independent experiments were performed. GAPDH: glyceraldehyde 3-phosphate dehydrogenase; IgG: immunoglobulin G; a.u.: arbitrary units.

(SCF) in liquid culture. At day 11, the percentage of mature CD41<sup>hi</sup>CD42a<sup>hi</sup> MKs was strikingly reduced, while the percentage of CD41<sup>low</sup>CD42<sup>-</sup> and CD41<sup>-</sup>CD42<sup>-</sup> cells was increased in F1-II2 compared with the control (Figure 5A). The percentage of high-ploidy cells ( $\geq 8n$ ) was

reduced among *FLI1* variant carriers at day 12 (11.9, 8.2 and 5.8% for the control, the F1-II2 and the F1-III1 affected members, respectively) and day 14 of maturation (11.3, 6.2 and 4.5% for the control, the F1-II2 and the F1-III1 affected members, respectively) (Figure 5B). At days 12-



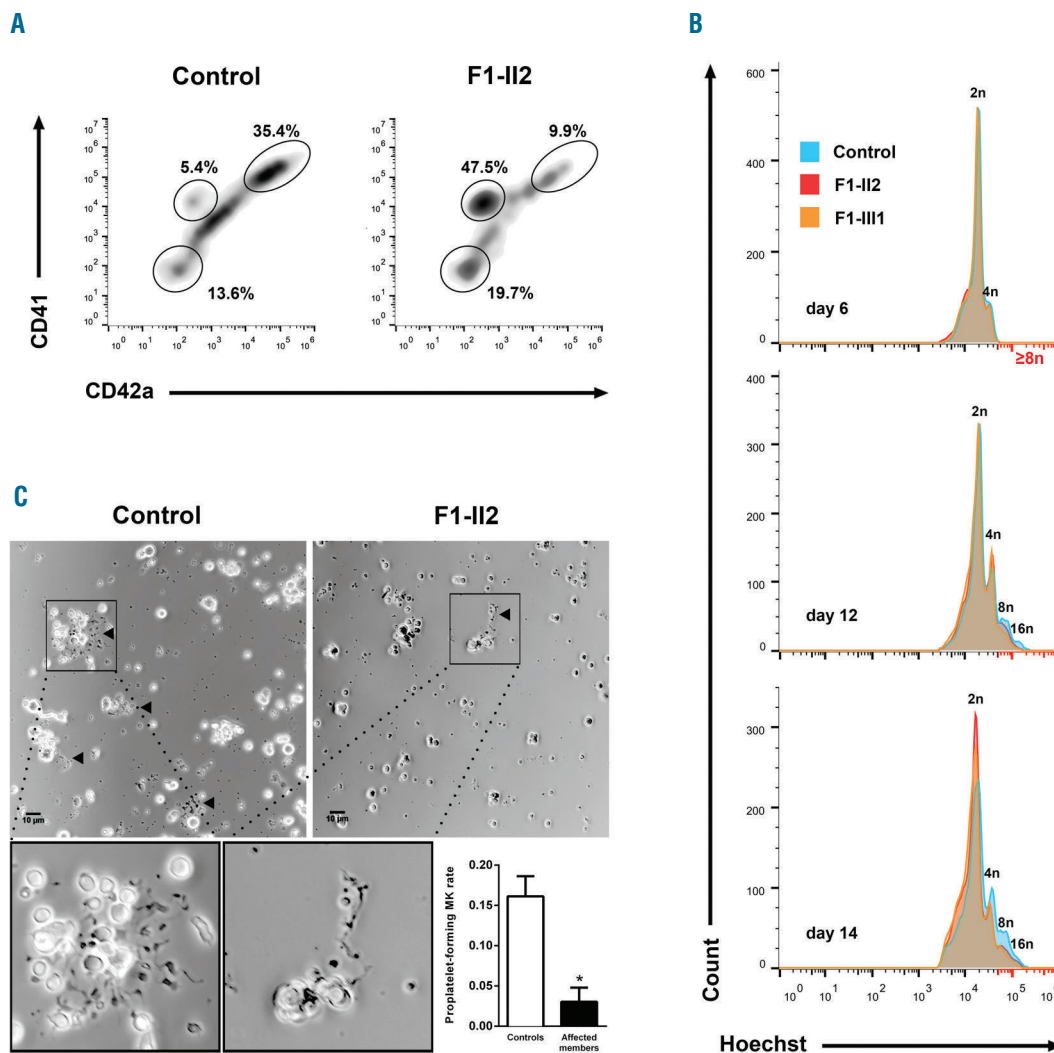
**Figure 4. Functional characterization of the *FLI1* variants.** (A) Left: representative western blot analysis of FLI1 expression in platelets from the affected members (F1-II2, F1-III1) and three control individuals. GAPDH and actin were used as a protein loading control. Right: the results of the densitometric analysis were normalized to actin and expressed as mean  $\pm$  SEM from three independent experiments. (B) Left: representative western blot analysis of FLI1 expression in GripTite 293 MSR cells transfected with an empty vector, wild-type (WT) or variant *FLI1* constructs using an anti-HA antibody. GAPDH was used as a protein loading control. Right: the results of the densitometric analysis are expressed as mean  $\pm$  SEM. Three independent experiments were performed. (C) GripTite 293 MSR cells were co-transfected with an empty vector, WT or variant *FLI1* constructs including the c.970C.T *FLI1* variant previously reported by Stevenson *et al.*<sup>9</sup> (p.R324W) along with the luciferase reporter plasmid containing three tandem copies of the ETS-binding site upstream of the HSV tk promoter (E743tk80Luc) and pGL4.73 Renilla luciferase control vector. Firefly to renilla luminescence ratios (Fluc/Rluc) were calculated to compensate for transfection efficiency and expressed as fold change relative to empty vector. The data represent the mean  $\pm$  SEM of three independent experiments; \* $P < 0.05$ , \*\*\*\* $P < 0.0001$  vs. WT (one-way ANOVA with Dunnett's *post hoc* test). (D) Western blot analysis of WT and variant FLI1 subcellular localization. GripTite 293 MSR cells were transfected with an empty vector, WT or variant *FLI1* constructs. The lamin-B1 and GAPDH expression were used as nuclear and cytoplasmic markers, respectively. The data are the mean  $\pm$  SEM of four independent experiments; \* $P < 0.05$  vs. WT (Mann-Whitney test). (E) Left: representative immunofluorescence microscopy images of H9C2 cells transfected with WT or variant *FLI1* constructs visualized using bright field illumination and immunofluorescence after FLI1 and DAPI staining; scale bar, 5  $\mu$ m. Right: quantification of the nuclear and cytoplasmic integrated density of fluorescence. The data are expressed as mean  $\pm$  SEM of the nucleocytoplasmic ratio of fluorescence integrated density from two independent experiments ( $\geq 30$  total cells were assessed for each condition); \*\*\*\* $P < 0.0001$  vs. WT (Mann-Whitney test). GAPDH: glyceraldehyde 3-phosphate dehydrogenase; HA: hemagglutinin; BF: bright field.

13, the percentage of proplatelet (PPT)-forming MKs was significantly reduced in the affected members F1-II2 and F2-II4 compared with three controls ( $16 \pm 1\%$  vs.  $3 \pm 1\%$ ,  $P < 0.05$ , Figure 5C). MKs from patients were smaller and formed very few PPTs, which displayed reduced extensions and branching (Figure 5C).

**Platelets from affected members exhibit giant  $\alpha$ -granules and an absence of dense granules**

May-Grünwald-Giemsa staining of peripheral blood smears of patient F1-II2 revealed enlarged platelets with giant  $\alpha$ -granules (Figure 6A). Accordingly, TEM analysis showed the presence of large fused  $\alpha$ -granules in the carriers' platelets (Figure 6B). The mean diameter of the  $\alpha$ -

granules was significantly increased (Figure 6C). The mean  $\alpha$ -granule number per  $\mu\text{m}^2$  of platelet section was slightly reduced in the FLI1 p.R337Q variant carrier but not in the p.K345E variant carrier. We quantified the platelet surface on sections (Figure 6C), which confirmed the increased platelet size associated with *FLI1* variants (Table 1). Whole mount electron microscopy, FIB-SEM and TEM was used to quantify the dense granules, which were nearly absent in carriers' platelets (Figure 6B,C and *Online Supplementary Figure S2*), thereby indicating a dense granule storage pool deficiency. Precise platelet content analysis and 3D reconstruction of the  $\alpha$ - and dense granules within platelets was performed using the FIB-SEM technique. Among affected members, the proportion of

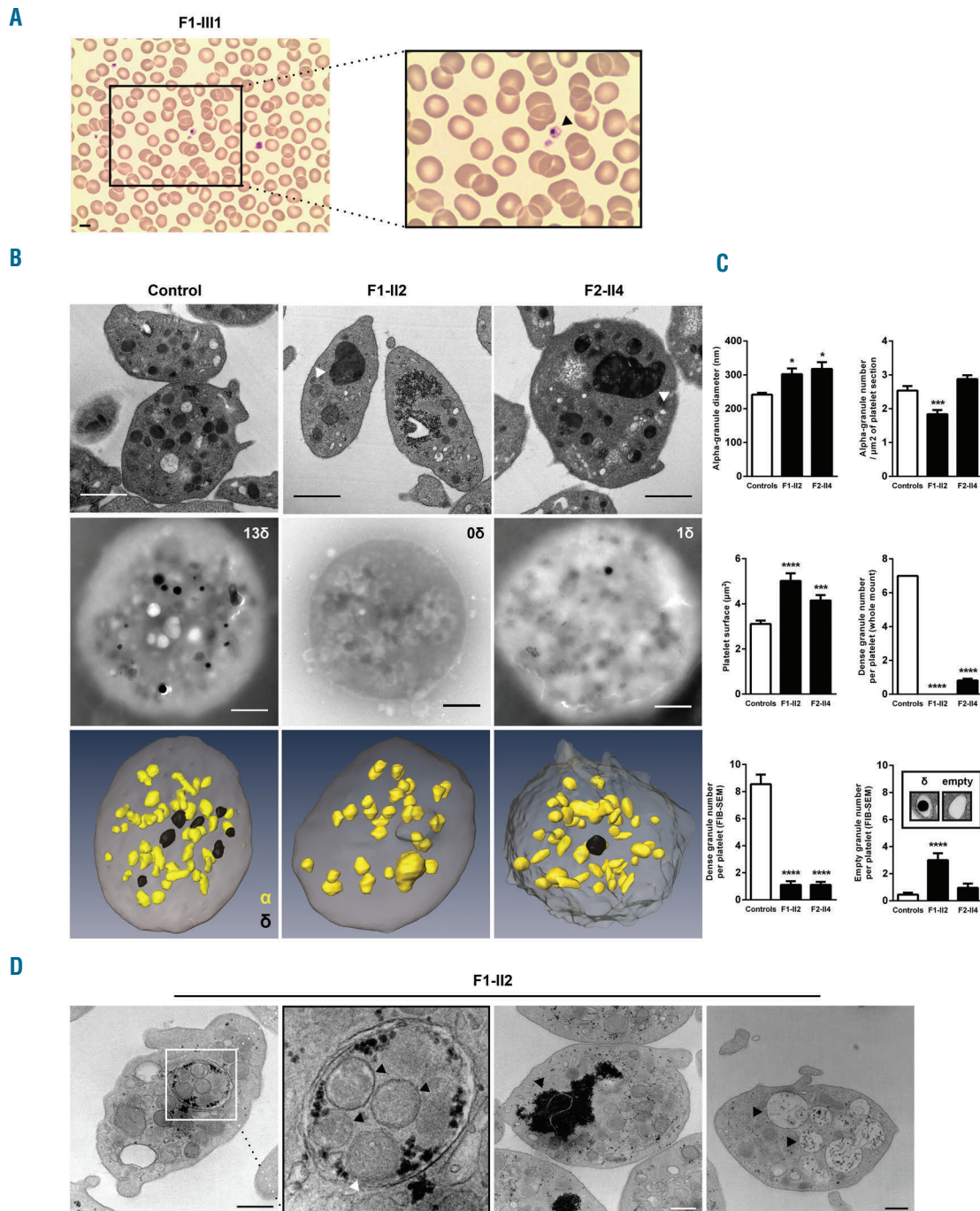


**Figure 5. Megakaryocyte differentiation and proplatelet formation.** Circulating CD34<sup>+</sup> progenitors from affected members or controls were isolated and cultured in the presence of TPO and SCF to induce megakaryocytic commitment. (A) MK differentiation was monitored using flow cytometry. The density plots represent CD41 and CD42a expression in Hoechst<sup>+</sup> cells from an affected member (F1-II2) and an unrelated control at day 11 of culture. The ellipse gates show the populations CD41<sup>+</sup>CD42a<sup>-</sup>, CD41<sup>int</sup>CD42a<sup>-</sup> cells and mature CD41<sup>+</sup>CD42a<sup>+</sup> MKs. (B) Ploidy level was monitored by flow cytometry. The histograms represent frequency distribution of Hoechst levels among the CD34<sup>+</sup>-derived cells from two affected members (F1-II2 and F1-III1) and an unrelated control at day 6, 12 and 14 of culture. (C) Representative microscopic images of proplatelet (PPT) formation after 13 days of culture. An expanded view of PPT formation is shown at the bottom. PPT formation was quantified in two affected members (F1-II2 and F2-II4) and three unrelated controls at culture days 12 and 13 from three independent experiments. The PPT-forming MK are indicated with triangle markers; scale bar, 10  $\mu\text{m}$ . The percentage of PPT-forming MKs was estimated by counting MKs harboring  $\geq 1$  cytoplasmic process with areas of constriction;  $\geq 180$  total cells were assessed for each individual. The results are expressed as mean  $\pm$  SEM; \* $P < 0.05$  (Mann-Whitney test). MK: megakaryocyte.



platelets displaying giant  $\alpha$ -granules (diameter > 400nm) was 25-29% (Table 2). The giant  $\alpha$ -granules had a mean volume of almost 8-fold that of normal  $\alpha$ -granules. While dense granules were easily distinguished through their typical dark central core and spherical geometry in control

platelets ( $8.6 \pm 0.7$  dense granules per platelet;  $n=12$ ), they were almost absent within platelets of *FLI1* variant carriers ( $1.1 \pm 0.3$ ,  $1.1 \pm 0.2$  dense granules per platelet for F1-II2 and F2-II4 affected members, respectively;  $n=12$ ) (Figure 6C). Empty dense granule membrane structures



**Figure 6. Platelet structure defects associated with *FLI1* variants.** (A) Representative May-Grünwald-Giemsa stained blood smears showing enlarged platelets with giant  $\alpha$ -granules (patient F1-III3). Scale bar, 5  $\mu$ m. (B) Platelet ultrastructural analysis from *FLI1* variant carriers (F1-II2 and F2-II4) and an unrelated control. Representative electron microscopy images of platelets transmission electron microscopy (TEM) ultrathin sections (top) and whole mount (middle). 3D reconstruction of platelet  $\alpha$ - and dense granules from focused ion beam scanning electron microscopy (FIB-SEM) images (bottom); Giant  $\alpha$ -granules are indicated with triangle markers;  $\delta$ : dense granule,  $\alpha$ : alpha-granule; Scale bar, 1  $\mu$ m. (C) Measurement of the  $\alpha$ -granules diameter and number /  $\mu$ m<sup>2</sup> of platelet section, platelet surface, number of dense granules and number of empty granules. Values are shown as the mean  $\pm$  SEM as quantified for  $\geq 100$  randomly selected platelets for TEM and whole mount and 12 randomly selected platelets for FIB-SEM; \* $P < 0.05$ , \*\*\* $P < 0.001$ , \*\*\*\* $P < 0.0001$  vs. controls (Mann-Whitney test). (D) Representative ultrastructural image of double membrane structures resembling autophagosome (far left panel with enlarged view middle left panel; black triangle marker:  $\alpha$ -granule; white triangle marker: mitochondrion), area of glycogen accumulation (middle right panel; black triangle marker: glycogen), and vacuoles (far right panel; black triangle marker: vacuole) in the platelet of the affected member F1-II2.



were rarely detected ( $3 \pm 0.5$ ,  $1 \pm 0.3$  and  $0.5 \pm 0.2$  copies for F1-II2, F2-II4 and control platelets, respectively;  $n=12$ ; Figure 6C). Double membrane structures resembling autophagosomes were detected in 3% of the platelets from the affected member F1-II2 (Table 2; Figure 6D, left panel), while these were absent in the affected member F2-II4. Accumulation of glycogen was found in the cytoplasm of many platelets from the affected members (86% and 51% in F1-II2 and F2-II4 platelets, respectively; Table 2 and Figure 6D, middle right panel) and vacuoles containing glycogen were also observed in 9% and 7% of F1-II2 and F2-II4 platelets, respectively (Table 2 and Figure 6D, far right panel). FIB-SEM image stacks in F1-II2 and a control allowed the 3D observation of the platelet morphological defects associated with *FLI1* variants (*Online Supplementary Videos S1* and *S2*).

## Discussion

In the present study, we identified two novel *FLI1* variants in patients from two unrelated families displaying CMTP with marked bleeding diathesis in one of them. The reported *FLI1* variants were associated with impaired megakaryocytic differentiation and PPT formation *in vitro* and with a defect in the protein subcellular localization, thereby providing further insights into the pathophysiology of *FLI1*-associated thrombocytopenia. Remarkably, we demonstrated that dense granules were nearly absent in the carriers' platelets. In addition, we showed that intraplatelet flow cytometry efficiently detected the biomarker MYH10 in *FLI1* variant carriers.

*FLI1* belongs to the ETS-domain transcription factor family, in which members bind a specific DNA consensus sequence to control the expression of genes that are essential in the regulation of cellular proliferation, differentiation and programmed cell death.<sup>15</sup> *FLI1* is mainly expressed in hematopoietic and vascular endothelial cells and plays a role in the regulation of essential MK genes.<sup>7,8,14,23</sup> In humans, a role for *FLI1* in megakaryopoiesis was highlighted by studies of Paris-Trousseau syndrome (PTS)<sup>24,26</sup> in which the *FLI1* locus is hemizygotously deleted, and more recently by two articles reporting *FLI1* variants associated with congenital thrombocytopenia.<sup>7,8</sup>

The affected members carried heterozygous variants, with a dominant mode of inheritance in the F1 family. The thrombocytopenia associated with the *FLI1* variants reported by Stockley *et al.*<sup>7</sup> and the PTS<sup>24,25</sup> were also shown to be transmitted in a dominant manner. Inversely, Stevenson *et al.* have described a family displaying inherited thrombocytopenia caused by a recessive *FLI1* variant.<sup>8</sup> We hypothesize that the mode of inheritance could rely on the degree of the protein functional impairment, as suggested by the moderately reduced transcriptional activity of the p.R324W variant, associated with the recessive inheritance pattern. Intriguingly, in mouse models of a targeted null *Fli1* variant, the mice carrying a heterozygous deletion of *Fli1* are phenotypically indistinguishable from the WT mice. We hypothesize that these mouse models do not undergo silencing of one *FLI1* allele during the early stages of megakaryopoiesis, contrary to that observed in humans.<sup>26</sup>

The bleeding phenotype of F2-II4 was much more severe than that of the affected members of the family F1 (ISTH BAT score 16 vs. 1). An additional bleeding disorder

(affecting coagulation, von Willebrand factor or fibrinolysis) was ruled out. A combination of genetic, environmental and lifestyle factors may interfere in the genotype-phenotype relationship. This difference is not likely to be directly related to platelet count or patient age; however, this could be explained in part by the sex of the patient. Indeed, the individual with a high bleeding score (F2-II4) predominantly suffered from gynecological and obstetrical bleeding. This may reflect the limitation of the ISTH BAT score to evaluate the bleeding risk in males who have not been challenged enough as compared to females.

In PTS, the platelet count seems to be as low as  $20 \times 10^9/l$  early in life and subsequently increases to near or low normal levels.<sup>25,27</sup> Similar to that shown in a previous report of *FLI1* variant carriers,<sup>7</sup> the older patient in the F1 family exhibited a higher platelet count than the younger one. The platelet counts and the increased MPV of the affected members were consistent with that observed in most PTS cases and *FLI1* variant-associated thrombocytopenia patients.<sup>7,25</sup> As noted in some PTS patients, the bleeding phenotype of patient F2-II4 was more severe than that expected with a mild thrombocytopenia, thereby suggesting associated abnormal platelet function.<sup>25,27</sup> In the present study, the affected members exhibited an aggregation defect upon low dose ADP, collagen, and TRAP-6 stimulation, which is a hallmark of dense granule storage pool deficiency. Data concerning aggregation assays has varied widely in the literature. Breton-Gorius *et al.* have reported normal aggregation curves in PTS patients.<sup>24</sup> Regarding *FLI1* variant-associated thrombocytopenia, Stevenson *et al.* have reported reduced aggregation to collagen and ADP.<sup>8</sup> Finally, in a murine model expressing a C-terminal truncated *Fli1* protein, aggregation tests revealed reduced ADP- and thrombin-induced aggregation.<sup>28</sup> These discrepancies could be in part due to differences regarding agonist concentrations or sources. Furthermore, storage pool diseases are known to be associated with variable aggregation profiles.<sup>29</sup>

In the study herein, NGS-based techniques led to the identification of the *FLI1* variants. These strategies require careful interpretation as they lead to the identification of a high number of non-pathogenic variants. However, clinical phenotype and screening laboratory tests are not specific and often fail to yield a specific diagnosis. Identifying biomarkers could facilitate the diagnostic process of inherited platelet disorders. In this regard, platelet-specific expression of MYH10, whose silencing is required for the switch from mitosis to endomitosis during MK maturation,<sup>30</sup> has been shown to constitute a biomarker for the inherited platelet disorders associated with *FLI1* and *RUNX1* alterations.<sup>7,8,12</sup> Platelet MYH10 expression is usually detected *via* western blot. This assay is complex and time-consuming, and the results must be carefully interpreted as control platelets exhibit low expression of MYH10. In this study, we developed a quantitative assay to detect MYH10 overexpression in *FLI1* variant carriers' platelets using intracellular flow cytometry. This test enabled rapid and accurate discrimination between the affected members and controls using only a small volume of PRP or whole blood.

$\alpha$ - and dense granule defects were identified as the main platelet defect in the affected members. It is important to study the mechanisms of this functional impairment, as bleeding diathesis may still occur in *FLI1*-associated thrombocytopenia patients with subnormal or normal

platelet counts.<sup>27</sup> We report that up to 29% of the patients' platelets displayed giant  $\alpha$ -granules, which is a characteristic feature of PTS<sup>24,25</sup> and have been previously reported in *FLI1* variant-associated thrombocytopenia,<sup>8</sup> albeit in a much smaller proportion of platelets (4-15%). This discrepancy can be attributed to differences in the analysis methods. Contrary to classic TEM, FIB-SEM enables evaluation of the whole granule content of a platelet. In the present study, normal or even increased serum PAI-1 levels suggest normal  $\alpha$ -granule content and release.<sup>31</sup> Taken together, these results indicate that the vesicle trafficking or the  $\alpha$ -granule proteins packaging but not the storage mechanisms may be impaired. Regarding dense granules, we demonstrated for the first time using electron microscopy (notably FIB-SEM), ATP secretion, mepacrine uptake and release, CD63 expression upon stimulation and serotonin measurement, that dense granules were nearly absent in the carriers' platelets and that empty dense granule membrane structures were rarely detected. This suggests that the platelet dense granule deficiency is due to an impaired biogenesis. A previous study on Jacobsen syndrome, a variant of PTS, has reported a dense granule storage pool deficiency in six patients<sup>32</sup> using whole mount electron microscopy, which only allows the evaluation of the electron opaque dense granules content. More recently, Stockley *et al.* showed that the platelet of *FLI1* variants carriers exhibited an ATP secretion defect<sup>7</sup> but have not evaluated the dense granule content. Interestingly, in a recent study reporting ChIP-seq data in murine MK,<sup>35</sup> FLI1 was shown to interact with *HPS4* and *RAB27B* genes, which are known regulators of dense granules biogenesis.<sup>34,35</sup> Autophagosome-like structures were noted in 3% of platelets from one affected member. Autophagy is known to be important in both megakaryopoiesis<sup>36</sup> and platelet function.<sup>37</sup> This observation thus suggests that autophagy may play a role in the pathophysiology of *FLI1*-associated thrombocytopenia. Glycogen accumulations were noted in the platelets of most of the affected members. The significance of this abnormal feature is unclear, although it may suggest that FLI1 plays a role in the regulation of platelet intracellular metabolic processes.

The p.R337Q and p.K345E variants are located within the ETS-domain, similar to the previously reported variants.<sup>7,8</sup> The pathogenic effect of these novel *FLI1* variants was confirmed by the significant reduction in transcriptional activity, as with all other *FLI1* variants reported in the literature.<sup>7,8</sup> Furthermore, we show that both p.R337Q and p.K345E variants displayed significantly altered cellular localization with a reduced nuclear accumulation. In a previous study based on directed mutagenesis, these residues were found to be important for the DNA-bound conformation of FLI1 and its nuclear accumulation.<sup>38</sup> Additionally, the p.R337A variant abolished ETS DNA-

binding activity.<sup>38</sup> Overall, these results suggest that both nuclear accumulation and DNA-binding capacity could interfere with FLI1 regulation of MK-specific genes. Other mechanisms may also cause this impairment, such as abnormal recruitment of co-regulators or altered activation of histone deacetylases or acetylases.<sup>39</sup>

*In vitro*, we noted a marked diminution of mature CD41<sup>hi</sup>CD42a<sup>hi</sup> MKs, along with an increase in CD41<sup>low</sup>CD42a<sup>-</sup> and CD41<sup>-</sup>CD42<sup>-</sup> cells among *FLI1* variant carriers. Even if we cannot formally exclude an impaired expression of CD41 and CD42a by FLI1 variants, the reduced percentage of high-ploidy cells ( $\geq 8n$ ) confirmed the MK differentiation defect. Furthermore, the platelet from the *FLI1* variant carriers displayed normal expression of CD42a (*data not shown*) and CD41. Consistent with our findings, data from PTS patients has shown that the MK transition from CD42a<sup>-</sup> to CD42a<sup>+</sup> was especially sensitive to a reduction in FLI1 dosage.<sup>26</sup> In a murine model with complete *Fli1* deficiency, the authors reported that megakaryocytic progenitors displayed an early blockage in MK differentiation.<sup>40</sup> Additionally, in a murine model expressing a truncated *Fli1* protein lacking the C-terminal transcriptional activation domain, bone marrow cell analysis revealed a 2-fold increase in CD34<sup>+</sup>CD41<sup>+</sup>CD42<sup>-</sup> cells.<sup>28</sup>

The study herein provides new insights into the mechanisms that drive *FLI1*-associated thrombocytopenia and highlights its function in normal and pathologic megakaryopoiesis and in platelet granule biogenesis. TEM and FIB-SEM enabled a more detailed description of the ultrastructural features associated with this disease, demonstrating that dense granules were nearly absent in the carriers' platelets. This work also shows that the increased expression of MYH10 in the platelets of *FLI1* variant carriers was efficiently detected by flow cytometry.

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