Regular Article

Targeting anticoagulant protein S to improve hemostasis in hemophilia

Raja Prince,[1,2,]* Luca Bologna,[1,2,]* Mirko Manetti,[3] Daniela Melchiorre,[4] Irene Rosa,[3] Natacha Dewarrat,[1,2] Silvia Suardi,[5] Poorya Amini,[6] José A. Fernández,[7] Laurent Burnier,[7] Claudia Quarroz,[1,2] Maria Desiré Reina Caro,[1,2] Yasuhiro Matsumura,[8] Johanna A. Kremer Hovinga,[1,2] John H. Griffin,[7] Hans-Uwe Simon,[6] Lidia Ibba-Manneschi,[3] François Saller,[9] Sara Calzavarini,[1,2,]* and Anne Angelillo-Scherrer[1,2]

[1]Department of Hematology and Central Hematology Laboratory, Inselspital, Bern University Hospital, and [2]Department of Clinical Research, University of Bern, Bern, Switzerland; [3]Department of Experimental and Clinical Medicine, Section of Anatomy and Histology, and [4]Department of Experimental and Clinical Medicine, Section of Internal Medicine, Rheumatology Unit, Careggi University Hospital, University of Florence, Florence, Italy; [5]Vetsuisse, Institute of Animal Pathology, Comparative Pathology Platform, and [6]Institute of Pharmacology, University of Bern, Bern, Switzerland; [7]Department of Molecular and Experimental Medicine, Scripps Research Institute, La Jolla, CA; [8]Division of Developmental Therapeutics, Research Centre for Innovative Oncology, National Cancer Centre Hospital East, Chiba, Japan; and [9]INSERM UMR-S 1176, Université Paris-Sud, Université Paris-Saclay, Le Kremlin-Bicêtre, France

KEY POINTS

- **Targeting anticoagulant protein S rebalances coagulation in hemophilia.**

- **Protein S in joints is a novel pathophysiological contributor to hemarthrosis and a potential therapeutic target in hemophilia.**

Improved treatments are needed for hemophilia A and B, bleeding disorders affecting 400 000 people worldwide. We investigated whether targeting protein S could promote hemostasis in hemophilia by rebalancing coagulation. Protein S (PS) is an anticoagulant acting as cofactor for activated protein C and tissue factor pathway inhibitor (TFPI). This dual role makes PS a key regulator of thrombin generation. Here, we report that targeting PS rebalances coagulation in hemophilia. PS gene targeting in hemophilic mice protected them against bleeding, especially when intra-articular. Mechanistically, these mice displayed increased thrombin generation, resistance to activated protein C and TFPI, and improved fibrin network. Blocking PS in plasma of hemophilia patients normalized in vitro thrombin generation. Both PS and TFPIα were detected in hemophilic mice joints. PS and TFPI expression was stronger in the joints of hemophilia A patients than in those of hemophilia B patients when receiving on-demand therapy, for example, during a bleeding episode. In contrast, PS and TFPI expression was decreased in hemophilia A patients receiving prophylaxis with coagulation factor concentrates, comparable to osteoarthritis patients. These results establish PS inhibition as both controller of coagulation and potential therapeutic target in hemophilia. The murine PS silencing RNA approach that we successfully used in hemophilic mice might constitute a new therapeutic concept for hemophilic patients. (*Blood*. 2018;131(12):1360-1371)

Introduction

Hemophilia A (HA) and B (HB) are hereditary X-linked disorders.[1-3] They are caused by mutations in the factor VIII (FVIII) gene (*F8*) or the factor IX (FIX) gene (*F9*), respectively, leading to the deficiency of the encoded protein that is an essential component of the intrinsic pathway of coagulation (Figure 1A).

Patients with severe hemophilia often suffer from spontaneous bleeding within the musculoskeletal system, such as hemarthrosis. This can result in disability at a young age if left untreated.[4]

Current hemophilia treatment involves factor replacement therapy.[1,4] This therapy improves quality of life (QoL) but some drawbacks remain. Factors are administered intravenously, and because of their short half-life, they must be repetitively infused, a practice that carries with it major discomfort for the patient and a risk of infection and venous damage. More important, patients under factor replacement therapy can develop inhibitory alloantibodies. Inhibitors render replacement therapy ineffective,

limit patient access to a safe and effective standard of care, and predispose them to an increased morbidity and mortality risk.[1,4,5]

New therapies focus on the development of products capable of decreasing the frequency of prophylactic infusions, thus potentially improving both compliance to therapy and QoL. Besides long-lasting FVIII and FIX, novel approaches comprise the replacement of the gene necessary for production of the endogenous coagulation factor, the bispecific antibody technology to mimic the coagulation function of the missing factor, and the targeting of coagulation inhibitors such as tissue factor pathway inhibitor (TFPI) or antithrombin as a strategy to rebalance coagulation in patients with hemophilia.[5] Recently, it was shown that an activated protein C (APC)–specific serpin rescues thrombin generation in vitro and restores hemostasis in hemophilia mouse models.[6]

Here, we investigated whether targeting protein S (PS)[7] could promote hemostasis in hemophilia by rebalancing coagulation (Figure 1B). PS, encoded by the *PROS1* gene, acts as a cofactor

for APC in the inactivation of factor Va (FVa) and activated FVIII (FVIIIa)[8] and for TFPI in the inhibition of FXa.[9,10] This dual role makes PS a key regulator of thrombin generation. The importance of PS as an anticoagulant is illustrated by the dramatic clinical manifestations observed in patients with severe PS deficiency.[11] Homozygous PS deficiency leads to purpura fulminans and disseminated intravascular coagulation (DIC), which are fatal if untreated. Heterozygous PS deficiency has variable penetrance but can be associated with an increased risk of thromboembolic events.[12,13]

Previous studies showed that $Pros1^{-/-}$ mice die in utero in late gestation with a phenotype similar to the one observed in humans.[14,15] We therefore consider $Pros1^{-/-}$ mice to be a suitable model of the human disease.

Materials and methods

Mice

$F8^{-/-}$ mice (B6;129S4-$F8^{tm1Kaz}$/J) and $F9^{-/-}$ mice (B6.129P2-$F9^{tm1Dws}$/J) with C57BL/6J background were obtained from the Jackson Laboratory. $Pros1^{+/-}$ mice were progeny of the original colony.[15] The Swiss Federal Veterinary Office approved the experiments. Mice were genotyped, as has been described previously.[15-17]

TF-induced pulmonary embolism

A model of venous thromboembolism was adapted from Weiss et al[18] with minor modifications.[15] Anesthetized mice, ages 6 to 9 weeks old, received human recombinant tissue factor (TF) (Dade Innovin, Siemens) intravenously (2 μL/g) at 4.25 nM (1:2 dilution) or 2.1 nM (1:4 dilution). Two minutes after the onset of respiratory arrest or at the completion of the 20-min observation period, lungs were harvested and fixed in 4% paraformaldehyde (PFA). Lung sections were stained with hematoxylin and eosin and for fibrin. The extent of fibrin clots in the lungs was assessed as the number of intravascular thrombi in 10 randomly chosen nonoverlapping fields (×10 magnification).

Tail clipping model in HA mice

We assessed 2 different tail clipping models to evaluate bleeding phenotype, as has been described.[6] Briefly, the distal tail of 8- to 10-week-old mice was transected at 2 mm (mild injury), and the bleeding was venous, or at 4 mm (severe injury), and the bleeding was arterial and venous.[19] Bleeding was quantified as blood lost after 30 min or 10 min, respectively. In the severe injury model, some $F8^{-/-}Pros1^{+/-}$ mice received a rabbit anti-human PS immunoglobulin G (IgG) (Dako) or rabbit isotype IgG (R&D Systems) intravenously at a dose of 2.1 mg/kg 2 min before tail transection.

Acute hemarthrosis model

Induction of joint bleeding in anesthetized 9- to 12-week-old mice, knee diameter measurements, and analgesic coverage were performed according to Øvlisen et al.[20] Joint diameters were measured at 0 hour and 72 hours with a digital caliper (Mitutoyo 547-301; Kanagawa). At 72 hours, mice were sacrificed, and knees were isolated, fixed in 4% PFA, decalcified, and embedded in paraffin. The intra-articular bleeding score (IBS) was assessed, as has been described.[21]

In vivo PS inhibition

Ten-week-old mice received a continuous infusion of rabbit anti-human PS-IgG (Dako Basel, Switzerland) or rabbit isotype IgG (R&D Systems) at 1 mg/kg/d through subcutaneous osmotic minipumps (model 2001; Alzet).

Alternatively, 10-week-old mice were treated with a single dose of mouse-specific small interfering RNA (siRNA) (s72206; Life Technologies) or control siRNA (4459405, in vivo negative control 1 Ambion; Life Technologies) at 1 mg/kg by using a transfection agent (Invivofectamine 3.0; Invitrogen, Life Technologies), following the manufacturer's instructions. An acute hemarthrosis model was applied 2.5 days after PS inhibition.

Statistical methods

Values were expressed as means ± SEMs. The chi-square for nonlinked genetic loci was used to assess the Mendelian allele segregation. Survival data in the TF-induced venous thromboembolism model were plotted using the Kaplan-Meier method. A log-rank test was used to statistically compare the curves (Prism 6.0d; GraphPad). The other data were analyzed by t test and 1-way and 2-way analyses of variance with GraphPad Prism 6.0d. A P value of less than .05 was considered statistically significant.

Additional methods are described in the supplemental Materials and methods (available on the *Blood* Web site).

Results

Loss of X-ase activity rescues $Pros1^{-/-}$ mice

$Pros1^{+/-}$ females crossed with $F8^{-/-}$ males produced 25% $F8^{+/-}Pros1^{+/-}$ progeny. $F8^{+/-}Pros1^{+/-}$ females bred with $F8^{-/-}$ males resulted in 25% $F8^{-/-}Pros1^{+/-}$ progeny (supplemental Figure 1A-C). Similar observations were made with $F9^{-/-}Pros1^{+/-}$ mice (supplemental Figure 1D-F). As was expected, $F8^{-/-}Pros1^{-/-}$ and $F9^{-/-}Pros1^{-/-}$ mice did not display FVIII and FIX plasma activity, respectively, and PS was not detected in $F8^{-/-}Pros1^{-/-}$ and $F9^{-/-}Pros1^{-/-}$ mice plasma (Figure 1C-D). PS levels in $F8^{-/-}Pros1^{+/-}$ and $F9^{-/-}Pros1^{+/-}$ mice were ~50% to 60% less than in $F8^{-/-}Pros1^{+/+}$ and $F9^{-/-}Pros1^{+/+}$ mice (Figure 1C-D), as reported.[14,15]

Of 295 pups from $F8^{-/-}Pros1^{+/-}$ breeding pairs, 72 (24%) were $F8^{-/-}Pros1^{+/+}$, 164 (56%) were $F8^{-/-}Pros1^{+/-}$, and 59 (20%) were $F8^{-/-}Pros1^{-/-}$ ($\chi^2 = 4.8$, $P = .09$). Thus, $F8^{-/-}Pros1^{-/-}$ mice were present at the expected Mendelian ratio. In contrast, of 219 pups from $F9^{-/-}Pros1^{+/-}$ breeding pairs, 56 (26%) were $F9^{-/-}Pros1^{+/+}$, 132 (60%) were $F9^{-/-}Pros1^{+/-}$, and 31 (14%) were $F9^{-/-}Pros1^{-/-}$ ($\chi^2 = 14.95$, $P = .001$). This is compatible with a transmission ratio distortion for $F9^{-/-}Pros1^{-/-}$ mice consistent with the decreased litter sizes compared with those of matings from $F9^{+/+}Pros1^{+/+}$ mice (5.2 ± 0.7 vs 9.8 ± 1.8, $n = 4$ matings, over 3 generations, $P = .046$). We then assessed the viability of 101 E12 to E16 embryos from $F9^{-/-}Pros1^{+/-}$ breeding pairs. Twelve embryos (=12%) were found dead. A comparable mortality rate was found in $F9^{+/+}Pros1^{+/+}$ embryos. The genetic scoring of the 89 live E12 to E16 embryos revealed that 27 (27% of the total number of embryos, 101) were $F9^{-/-}Pros1^{+/+}$, 46 (46%) were $F9^{-/-}Pros1^{+/-}$, and 16 (16%) were $F9^{-/-}Pros1^{-/-}$ ($\chi^2 = 2.8$, $P = .24$). Therefore, $F9^{-/-}Pros1^{-/-}$ embryos were present at the expected Mendelian ratio, pointing to an increased mortality of $F9^{-/-}Pros1^{-/-}$ embryos during late gestation (>E16) and delivery. Similar observations were made in $Pros1^{-/-}$ embryos.[15]
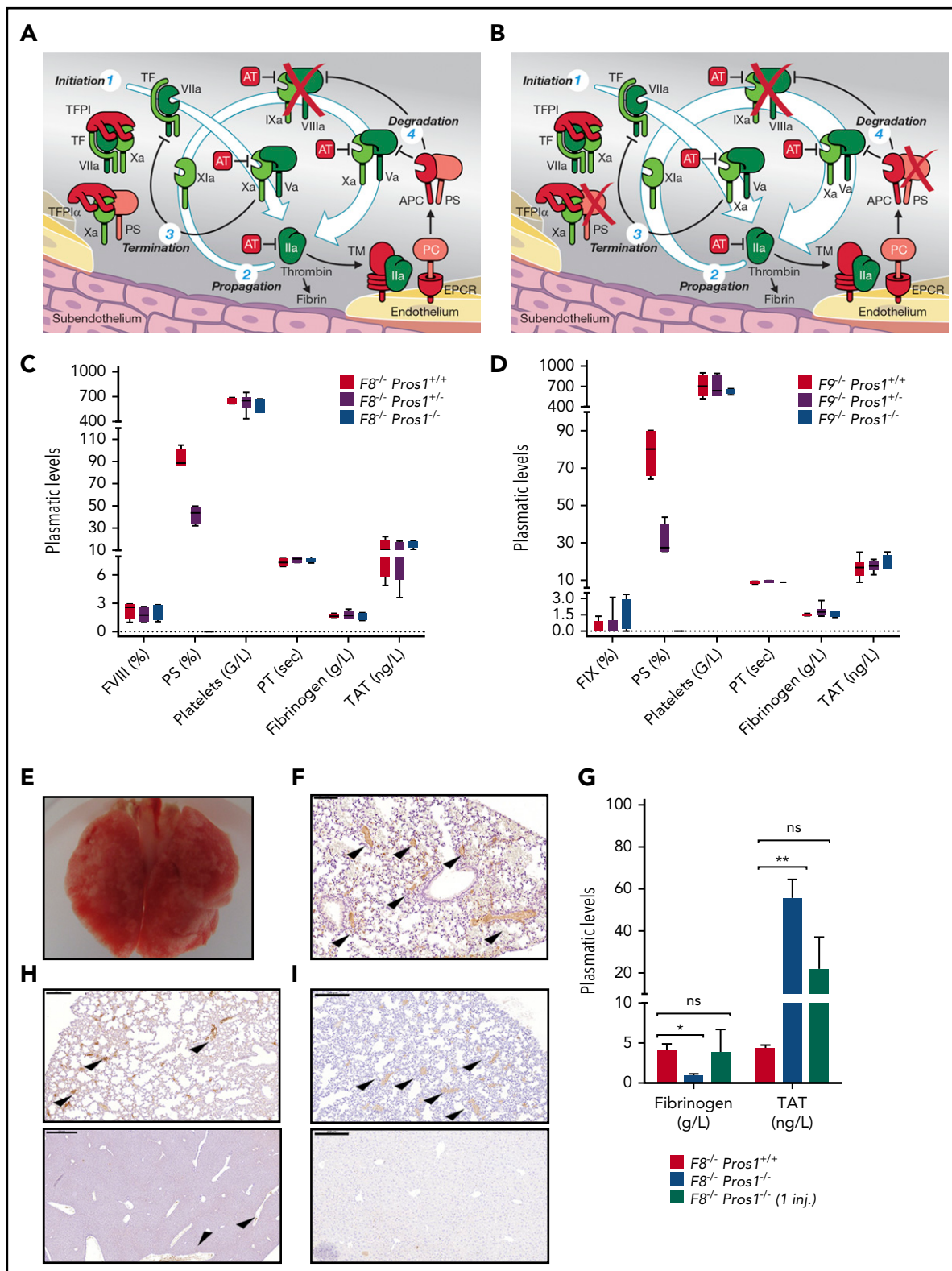
Figure 1. Loss of X-ase activity rescues *Pros1*−/− mice. (A) Schematic model of thrombin generation in hemophilic condition. One of the major coagulation complexes is the intrinsic tenase (X-ase) complex.[45] X-ase comprises activated FIX (FIXa) as the protease, activated FVIII as the cofactor, and factor X (FX) as the substrate. Although the generation or exposure of TF at the site of injury is the primary event in initiating coagulation via the extrinsic pathway, the intrinsic pathway X-ase is important because of the limited amount of available active TF in vivo and the presence of TFPI, which when complexed with activated FX (FXa), inhibits the TF/activated factor VII (FVIIa) complex[46] (Figure 1A). Thus, sustained thrombin generation depends upon the activation of both FIX and FVIII[47] (Figure 1A). This process is amplified because FVIII is activated by both FXa and thrombin, and FIX is activated by both FVIIa and activated factor XI (FXIa), the latter factor being previously activated by thrombin. Consequently, a progressive increase in FVIII and FIX activation occurs as FXa and thrombin are formed. (B) The experimental approach to enhancing thrombin generation in severe hemophilia A and B by targeting *Pros1*. (C-D) Murine model validation and evaluation of DIC hematologic parameters in hemophilic adult mice with and without *Pros1* deficiency: PS (antigenic), FVIII (coagulant activity), or

$F8^{-/-}Pros1^{-/-}$ and $F9^{-/-}Pros1^{-/-}$ mice appeared completely normal. Their viability was monitored up to 20 months ($n = 4$) and 16 months ($n = 2$), respectively, without showing any difference from $F8^{-/-}Pros1^{+/+}$ and $F9^{-/-}Pros1^{+/+}$ mice, respectively.

Because a complete $Pros1$ deficiency in mice leads to consumptive coagulopathy,[15] we assessed whether $F8^{-/-}Pros1^{-/-}$ and $F9^{-/-}Pros1^{-/-}$ mice developed DIC. DIC parameters were comparable in $F8^{-/-}Pros1^{+/+}$, $F8^{-/-}Pros1^{+/-}$, and $F8^{-/-}Pros1^{-/-}$ mice (Figure 1C) and in $F9^{-/-}Pros1^{+/+}$, $F9^{-/-}Pros1^{+/-}$, and $F9^{-/-}Pros1^{-/-}$ mice (Figure 1D). Activated partial thromboplastin time was equally prolonged in $F8^{-/-}Pros1^{+/+}$ (69 ± 2 s), $F8^{-/-}Pros1^{+/-}$ (68 ± 3 s), and $F8^{-/-}Pros1^{-/-}$ (63 ± 3 s) mice (mean $\pm$ SEM, $n = 6$ per group, $P = .3$) because of the absence of FVIII. Comparable data were obtained with $F9^{-/-}Pros1^{+/+}$, $F9^{-/-}Pros1^{+/-}$, and $F9^{-/-}Pros1^{-/-}$ mice. Moreover, no thrombosis or fibrin deposition was found in brain, lungs, liver, or kidney of $F8^{-/-}Pros1^{-/-}$, and $F9^{-/-}Pros1^{-/-}$ mice (supplemental Figure 2).

To explore whether restoring intrinsic X-ase activity by FVIII infusion induces DIC, thrombosis, and purpura fulminans in $F8^{-/-}Pros1^{-/-}$ mice, we administered recombinant FVIII (rFVIII) intravenously. No mouse died following rFVIII injection. Thrombi in numerous blood vessels and bleeding in the lungs were found in $F8^{-/-}Pros1^{-/-}$ mice 24 hours after a single injection of an overdose of rFVIII (Figure 1E-F). Twenty-four hours after repeated administration of a normal dose of rFVIII, coagulation analyses showed incoagulable prothrombin time (PT) (not shown), low fibrinogen, and high thrombin–antithrombin (TAT) levels, compatible with an overt DIC (Figure 1G). Finally, after a single injection of a normal dose of rFVIII in $F8^{-/-}Pros1^{-/-}$ mice, fibrinogen and TAT levels were comparable to those of untreated $F8^{-/-}Pros1^{-/-}$ mice (Figure 1G). Although numerous thrombi were visible in lungs and liver (Figure 1H-I), none of these mice developed purpura fulminans.

Twenty-four hours after 2 injections of an overdose of recombinant FIX (rFIX) administered 12 hours apart in $F9^{-/-}Pros1^{-/-}$ mice (FIX plasma level was 200% in $F9^{-/-}Pros1^{-/-}$ at 24 hours), there were thrombi visible in lungs (supplemental Figure 3) but not in other organs (data not shown). However, none of these mice developed purpura fulminans or died.

Therefore, loss of X-ase activity rescues the embryonic lethality of complete $Pros1$ deficiency. However, the rescue was only partial with the loss of FIX activity, and in reconstitution experiments, the effect was less prominent with rFIX in $F9^{-/-}Pros1^{-/-}$ mice than with rFVIII in $F8^{-/-}Pros1^{-/-}$ mice. A possible explanation is that severe HB appears to be a less serious bleeding condition than is severe HA.[22-26] In the current experimental context, this might also be due to the fact that activated FIX (FIXa), in the presence of its cofactor FVIIIa, is more effectively inhibited by PS.[27]

Loss of X-ase activity does not prevent lethality caused by TF-induced thromboembolism in $Pros1^{-/-}$ mice

We demonstrated previously that although 88% of $Pros1^{+/+}$ mice survived to a TF-induced thromboembolism model,[18] only 25% of $Pros1^{+/-}$ mice were still alive 20 min after a low TF dose injection ($\sim$1.1 nM).[15] When using a higher TF dosage ($\sim$4.3 nM), both $Pros1^{+/+}$ and $Pros1^{+/-}$ mice died within 20 min. However, $Pros1^{+/-}$ mice died earlier than did $Pros1^{+/+}$ mice. HA and wild-type (WT) mice were equally sensitive to this high TF dose, with more than 85% of them succumbing within 15 min (Figure 2A). In contrast, >75% WT mice under thromboprophylaxis with a low molecular weight heparin (LMWH) survived (Figure 2A). Thus, in contrast with LMWH, HA does not protect mice against TF-induced thromboembolism. We then investigated $F8^{-/-}Pros1^{+/+}$, $F8^{-/-}Pros1^{+/-}$, and $F8^{-/-}Pros1^{-/-}$ mice in the same model. After the infusion of TF ($\sim$2.1 nM), 40% to 60% of the mice died ($P > .05$), independently of their $Pros1$ genotype (Figure 2B). However, there was a trend for $F8^{-/-}Pros1^{-/-}$ and $F8^{-/-}Pros1^{+/-}$ mice succumbing earlier than $F8^{-/-}Pros1^{+/+}$ mice and for $F8^{-/-}Pros1^{+/-}$ dying earlier than $F8^{-/-}Pros1^{+/+}$ mice (mean time to death: 12 ± 4 min for $F8^{-/-}Pros1^{+/+}$, 7 ± 2 min for $F8^{-/-}Pros1^{+/-}$, 8 ± 3 min for $F8^{-/-}Pros1^{-/-}$ mice; $n = 4$-6 per group; $P = .43$). Similar data were obtained with $F9^{-/-}Pros1^{+/+}$, $F9^{-/-}Pros1^{+/-}$, and $F9^{-/-}Pros1^{-/-}$ mice (data not shown).

Fibrin clots were detected in the lung arteries of $F8^{-/-}Pros1^{+/+}$ and $F8^{-/-}Pros1^{-/-}$ mice, which died during the TF-induced thromboembolic challenge (Figure 2C). Importantly, there were more thrombi in lungs from $F8^{-/-}Pros1^{-/-}$ than in lungs from $F8^{-/-}Pros1^{+/+}$ mice ($n = 48$ vs 26, respectively). Moreover, most arteries in $F8^{-/-}Pros1^{-/-}$ lungs were completely occluded, whereas they were only partially occluded in $F8^{-/-}Pros1^{+/+}$ lungs.

None of the $F8^{-/-}Pros1^{-/-}$ mice that succumbed during the TF-induced thromboembolic challenge developed purpura fulminans. Similar data were obtained with $F9^{-/-}Pros1^{+/+}$, $F9^{-/-}Pros1^{+/-}$, and $F9^{-/-}Pros1^{-/-}$ mice (not shown).

Loss of FVIII partially protects $Pros1^{-/-}$ mice against thrombosis in mesenteric arterioles

We then recorded thrombus formation in mesenteric arterioles, a model sensitive to defects in the intrinsic pathway of coagulation.[28] In $F8^{+/+}Pros1^{+/+}$ mice, thrombi grew to occlusive size in 20 min, and all injured arterioles were occluded (Figure 2D). As was expected, none of the arterioles of $F8^{-/-}Pros1^{+/+}$ displayed thrombosis, whereas $F8^{-/-}Pros1^{-/-}$ mice showed partial thrombi (Figure 2D).

Emboli were generated during thrombus formation in $F8^{+/+}Pros1^{+/+}$ mice but not in $F8^{-/-}Pros1^{+/+}$ mice. In $F8^{-/-}Pros1^{-/-}$ mice, multiple microemboli detached during partial thrombus growth, preventing the formation of occlusive thrombi. This

Figure 1 (continued) FIX (coagulant activity) plasma levels in $F8^{-/-}Pros1^{+/+}$, $F8^{-/-}Pros1^{+/-}$, and $F8^{-/-}Pros1^{-/-}$ adult mice (C) and in $F9^{-/-}Pros1^{+/+}$, $F9^{-/-}Pros1^{+/-}$ and $F9^{-/-}Pros1^{-/-}$ adult mice (D) ($n = 5$ per group); platelets ($n = 7$ per group), fibrinogen ($n = 8$ per group), PT ($n = 6$ per group), and TAT ($n = 6$ per group) in hemophilia A group (C) and platelets ($n = 5$ per group), fibrinogen ($n = 4$ per group), PT ($n = 4$ per group), and TAT ($n = 4$ per group) in hemophilia B group (D). (E-F) Macroscopic image of lungs from $F8^{-/-}Pros1^{-/-}$ mice 24 hours after a single intravenous injection of 2 U/g recombinant FVIII (Advate) infusion (E) and corresponding microscopic evaluation of fibrin clots (indicated by arrowheads) in lung section (F). (G-I) Recombinant FVIII (Advate) administration in $F8^{-/-}Pros1^{+/+}$ and $F8^{-/-}Pros1^{-/-}$: plasma levels of fibrinogen and TAT at 24 hours after 5 injections of 0.3 U/g Advate IV (injection time points: 1 hour before catheter insertion and 1 hour, 4 hours, 8 hours, and 16 hours after catheter insertion) ($n = 3$) (G, open and solid columns) and 24 hours after a single IV injection in $F8^{-/-}Pros1^{-/-}$ ($n = 3$) (G, dashed column), and representative immunohistochemistry allowing the detection of fibrin clots (indicated by arrowheads) in lungs and liver sections in $F8^{-/-}Pros1^{-/-}$ 24 hours after 0.3 U/g repeated IV injections of Advate (H) and after a single IV injection of 0.3 U/g Advate IV (I). All data are expressed as means $\pm$ SEM. *$P < .05$; **$P < .005$. AT, antithrombin; EPCR, endothelial protein C receptor; inj., injection; ns, not significant. Scale bars: 200 μm.
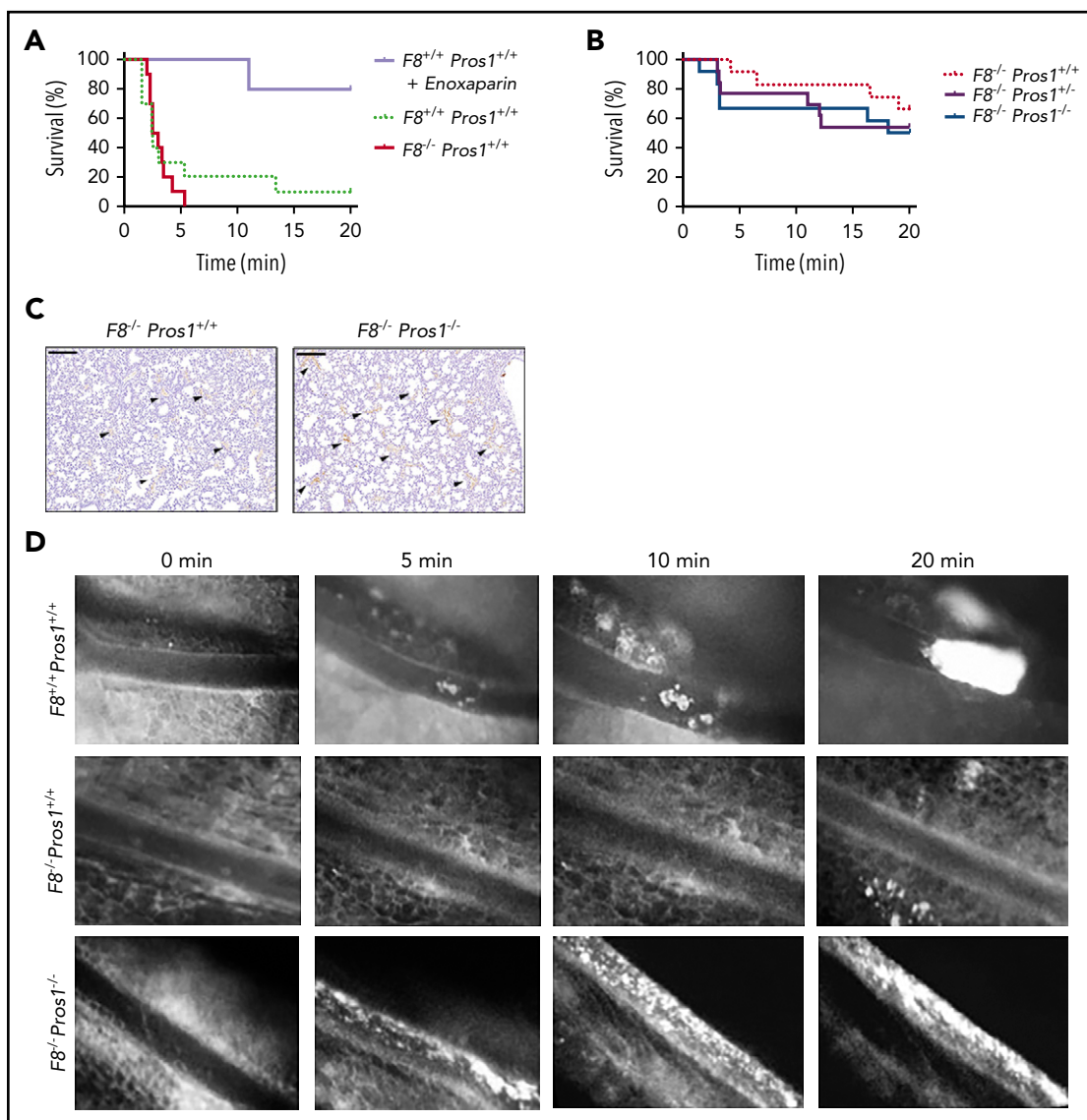
Figure 2. Murine models of thrombosis. (A-C) TF-induced venous thromboembolism in $F8^{+/+} Pros1^{+/+}$, $F8^{-/-} Pros1^{+/+}$, $F8^{-/-} Pros1^{+/-}$, and $F8^{-/-} Pros1^{-/-}$ mice (n = 10 per genotype). Anesthetized mice were injected intravenously via the inferior vena cava with different doses of recombinant TF (Innovin): 1/2 dilution (~4.3 nM TF) in panel A and 1/4 dilution (~2.1 nM TF) in panels B-C. In panel A, one group of $F8^{+/+}Pros1^{+/+}$ mice received an injection of the low molecular weight heparin (enoxaparin 60 μg/g s.c.). The time to onset of respiratory arrest that lasted at least 2 minutes was recorded. Experiments were terminated at 20 minutes. Kaplan-Meier survival curves (A-B). (C) Two minutes after onset of respiratory arrest or at the completion of the 20-minute observation period, lungs were excised and investigated for fibrin clots (indicated by arrowheads, immunostaining for insoluble fibrin, monoclonal antibody clone 102-10) Scale bars: 200 μm. (D) Thrombus formation in FeCl$_3$-injured mesenteric arteries recorded by intravital microscopy in $F8^{+/+} Pros1^{+/+}$, $F8^{-/-} Pros1^{+/+}$, and $F8^{-/-} Pros1^{-/-}$ mice, representative experiment (n = 3 per genotype).

difference between the genotypes might be due at least partly to the fact that in the presence of FVIIIa, FIXa is more effectively inhibited by PS.[27]

Pros1 targeting limits but does not abrogate tail bleeding in mice with HA

The bleeding phenotype was assessed by tail transection using a mild or a severe bleeding model.

In both models, blood loss was reduced in $F8^{-/-} Pros1^{-/-}$ mice in comparison with $F8^{-/-} Pros1^{+/+}$ mice (Figure 3A-B). When challenged by the mild model, $F8^{-/-} Pros1^{+/-}$ mice bled less did than $F8^{-/-} Pros1^{+/+}$ mice (Figure 3A). In contrast, when exposed to the severe model, $F8^{-/-} Pros1^{-/-}$ and $F8^{-/-} Pros1^{+/-}$ mice displayed

comparable blood loss (Figure 3B). However, $F8^{-/-}Pros1^{-/-}$ mice bled more than did $F8^{+/-}Pros1^{+/+}$ and $F8^{+/+}Pros1^{+/+}$ mice in both models (Figure 3A-B), indicating that the loss of *Pros1* in $F8^{-/-}$ mice did only partially correct the bleeding phenotype of $F8^{-/-}$ mice.

Then, a PS-neutralizing antibody[15] was used to investigate how inhibition of PS activity alters tail bleeding in $F8^{-/-}Pros1^{+/-}$ mice. This antibody limited blood loss in $F8^{-/-}Pros1^{+/-}$ mice (Figure 3C) to the same degree as complete genetic loss of *Pros1* (Figure 3B).

Pros1 targeting or PS inhibition fully protects HA and HB mice from acute hemarthrosis

Although bleeding may appear anywhere in hemophilia patients, most hemorrhages occur in the joints.[29,30] To determine
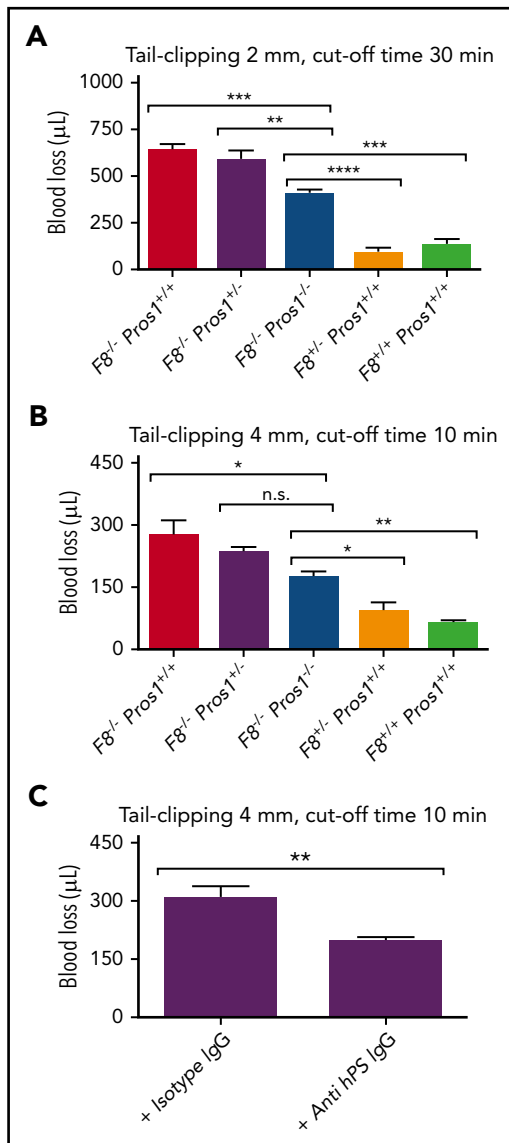
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Figure 3. Tail bleeding models. Blood was collected after 2-mm (A) and 4-mm (B) tail transections for 30 minutes (A) and 10 minutes (B) in a fresh tube of saline; total blood loss (in microliters) was then measured. $F8^{+/-}Pros1^{+/+}$ (white columns) and $F8^{+/+}Pros1^{+/+}$ (green columns) mice served as controls (n = 5 for all groups in panel A; n = 6 for all groups in panel A). (C) An anti-human PS antibody altered tail bleeding after a 4-mm transection. $^{*}P < .05$, $^{**}P < .01$, $^{***}P < .001$, $^{****}P < .0001$; n.s., not significant.

whether *Pros1* loss prevents hemarthrosis in hemophilic mice, we applied an acute hemarthrosis (AH) model to $F8^{-/-}Pros1^{+/+}$, $F8^{-/-}Pros1^{+/-}$, $F8^{-/-}Pros1^{-/-}$, and $F8^{+/+}Pros1^{+/+}$ mice. Knee swelling after injury was reduced in $F8^{-/-}Pros1^{-/-}$ and $F8^{+/+}Pros1^{+/+}$ mice in comparison with $F8^{-/-}Pros1^{+/+}$ and $F8^{-/-}Pros1^{+/-}$ mice (Figure 4A). There was also no difference in knee swelling between $F8^{-/-}Pros1^{-/-}$ and $F8^{+/+}Pros1^{+/+}$ mice (Figure 4A). Bleeding was observed in the joint space and synovium of $F8^{-/-}Pros1^{+/+}$ mice (IBS = 2, n = 5) but not of $F8^{-/-}Pros1^{-/-}$ mice (IBS = 0, n = 5) and $F8^{+/+}Pros1^{+/+}$ mice (IBS = 0, n = 5) (Figure 4B). There was more fibrin in the joint space and synovium from $F8^{-/-}Pros1^{+/+}$ than in those from $F8^{-/-}Pros1^{-/-}$ and $F8^{+/+}Pros1^{+/+}$ mice (Figure 4B). Similar data were obtained with $F9^{-/-}Pros1^{+/+}$ and $F9^{-/-}Pros1^{-/-}$ mice (IBS = 0, n = 3, and IBS = 2, n = 3, respectively) (supplemental Figure 4A-B).

These results were confirmed by the continuous subcutaneous infusion during 4 days of a PS-neutralizing antibody or a control antibody in $F8^{-/-}Pros1^{+/-}$ mice (starting 1 day before AH induction) (knee swelling in the PS-neutralizing antibody group was 0.43 ± 0.07 vs 0.69 ± 0.09 mm in the control group; n = 9, P = .04). PS plasma level in the PS-neutralizing antibody group was 26% ± 6% vs 45% ± 3% in the control group (n = 5, P = .017). In addition, PS inhibition was alternatively achieved by intravenous injection of a murine PS (mPS) siRNA prior to the AH challenge in $F8^{-/-}Pros1^{+/-}$ and $F8^{-/-}Pros1^{+/+}$ mice (Figure 4C-D). The IBS assessment confirmed the lack of intra-articular bleeding in $F8^{-/-}Pros1^{+/+}$ mice treated with mPS siRNA (IBS = 0.5, n = 3) when compared with those treated with control siRNA (IBS = 2, n = 3) (Figure 4C). Importantly, PS expression was reduced by mPS siRNA both in plasma (26% ± 3% vs 84% ± 11% in controls; n = 3, P = .006) and in the synovium (Figure 5A).

Both PS and TFPI are expressed in the synovium of mice

To understand the prominent intra-articular hemostatic effect of the genetic loss of *Pros1* and PS inhibition in hemophilic mice, we immunostained knee sections for PS and TFPI. PS was mainly present at the lining layer of the synovial tissue of $F8^{-/-}Pros1^{+/+}$ mice with AH treated with control siRNA, whereas synovial staining for PS was remarkably reduced in $F8^{-/-}Pros1^{+/+}$ mice with AH that received mPS siRNA (Figure 5A). In contrast, TFPI staining was more prominent in synovial tissue from hemophilic mice that received the mPS siRNA than in tissue from mice treated with the control siRNA (Figure 5A). However, TFPI expression was comparable in the synovial lining layer of both $F8^{-/-}Pros1^{+/+}$ and $F8^{-/-}Pros1^{-/-}$ mice (Figure 5B). Interestingly, PS expression was not detected in the synovial tissue of $F8^{-/-}Pros1^{+/+}$ mice, which received hrFVIII infusion at the time of the intra-articular injury, both in the noninjured and injured knees (supplemental Figure 5).

To demonstrate further that PS is expressed by fibroblast-like synoviocytes (FLS), we performed western blots on conditioned media collected from $F8^{+/+}Pros1^{+/+}$, $F8^{-/-}Pros1^{+/+}$, and $F8^{-/-}Pros1^{-/-}$ FLS. As is shown in Figure 5C, media of $F8^{+/+}Pros1^{+/+}$ and $F8^{-/-}Pros1^{+/+}$ FLS displayed a band at a molecular weight ~75 kDa, comparable to PS and similar to the one observed in plasma and platelets. As was expected, no staining was detected in media obtained from $F8^{+/+}Pros1^{-/-}$ FLS (Figure 5C).

We also studied TFPI expression in $F8^{-/-}Pros1^{+/+}$ and $F8^{-/-}Pros1^{-/-}$ FLS conditioned media (Figure 5D). All media displayed a band at ~50 kDa similar to the one observed with placenta lysates. TFPI isoform expression was investigated following protein deglycosylation because fully glycosylated TFPIα and TFPIβ migrate at the same molecular weight.[31] Deglycosylated TFPI from FLS media migrated as a single band at the molecular weight of TFPIα, similar to placenta TFPI (positive control for TFPIα) (Figure 5D). This indicates that FLS express TFPIα but not TFPIβ. Moreover, PS and TFPI expression increased in $F8^{-/-}Pros1^{+/+}$ FLS after stimulation with thrombin (Figure 5E-F).

Gene expression of *Pros1* and *Tfpi* was quantified by real-time polymerase chain reaction in relation to the expression of glyceraldehyde-3-phosphate dehydrogenase. *Pros1* and *Tfpi* transcripts were expressed by FLS. Expression of *Pros1* was
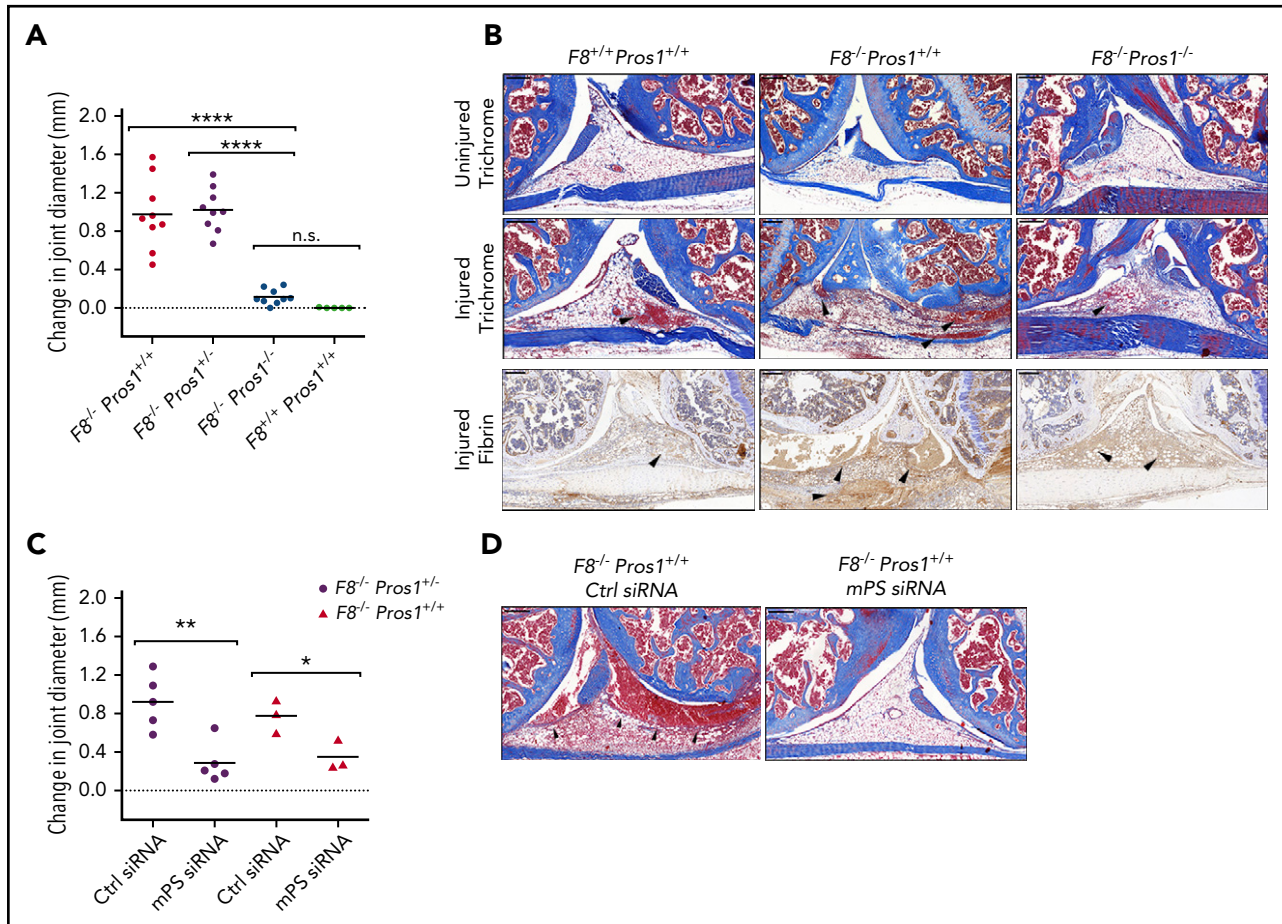
Figure 4. Acute hemarthrosis model. (A) Difference between the knee diameter 72 hours after the injury and before the injury in $F8^{-/-}Pros1^{+/+}$, $F8^{-/-}Pros1^{+/-}$, $F8^{-/-}Pros1^{-/-}$, and $F8^{+/+}Pros1^{+/+}$ mice. (B) Microscopic evaluation (Masson's trichrome stain and immunostaining for insoluble fibrin) of the knee intra-articular space of a representative, uninjured leg and injured leg after 72 hours in $F8^{+/+}Pros1^{+/+}$, $F8^{-/-}Pros1^{+/+}$, and $F8^{-/-}Pros1^{-/-}$ mice. (C) In vivo mPS silencing using specific siRNA: evaluation of the joint diameter 72 hours after injury in $F8^{-/-}Pros1^{+/-}$ and $F8^{-/-}Pros1^{+/+}$ mice treated with a single intraperitoneal infusion of mPS siRNA or control siRNA. (D) Microscopic evaluation (Masson's trichrome stain) of the knee intra-articular space of a representative injured leg after 72 hours in $F8^{-/-}Pros1^{+/+}$ mice previously treated with mPS siRNA or control siRNA. Arrowheads on Masson's trichrome staining indicate red blood cell extravasation, whereas in the immunohistochemistry, they indicate the deposit of insoluble fibrin. Measurements are presented as means ± SEMs. *$P < .05$; **$P < .005$; ****$P < .0001$. ctrl, control; n.s., not significant. Scale bars: 200 μm.

comparable in $F8^{+/+}Pros1^{+/+}$ and $F8^{-/-}Pros1^{+/+}$ FLS, and *Tfpi* expression was similar in $F8^{+/+}Pros1^{+/+}$, $F8^{-/-}Pros1^{+/+}$, and $F8^{-/-}Pros1^{-/-}$ FLS (supplemental Figure 6).

Both PS and TFPI are expressed in the synovium of patients with HA or HB

Human HA, HB, and osteoarthritis knee synovial tissues were then analyzed for both PS and TFPI (Figure 6A). A strong signal was found for TFPI and PS in the synovial lining and sublining layers of HA patients on demand (n = 7). By contrast, immunostaining for both PS and TFPI was decreased in HA patients under prophylaxis (n = 5). HB patients on demand displayed less signal for both PS and TFPI in the synovial lining and sublining layers (n = 4) than did HA patients on demand. Sections from osteoarthritis patients (n = 7) did not show an intense staining for TFPI and PS, similarly to hemophilic patients under prophylaxis. To evaluate which isoform of TFPI is expressed by human FLS, we performed western blotting on conditioned media of human FLS isolated from healthy subjects and patients with osteoarthritis. As with murine FLS, human FLS expresses TFPIα but not TFPIβ (Figure 6B). These data are consistent with murine observations

in which PS was not detected in the synovium after hrFVIII infusion (supplemental Figure 5).

Loss of *Pros1* is responsible for the lack of TFPI-dependent PS activity and resistance to APC in HA mice

The full protection against AH in HA or HB mice lacking *Pros1* or in which PS was inhibited could be explained at least partly by the lack of PS cofactor activity for APC and TFPI in the joint. However, the reason for a partial hemostatic effect of the lack of *Pros1* or PS inhibition in HA mice challenged in the tail bleeding models needs to be further investigated.

Ex vivo TF-initiated thrombin-generation testing has shown a correlation between the capacity of plasma to generate thrombin and the clinical severity of hemophilia.[32-34] Therefore, we investigated the impact of *Pros1* loss on thrombin generation in the plasma of HA mice. TFPI-dependent PS activity was not assessed in platelet-free plasma (PFP) but in platelet-rich plasma (PRP) because TFPI cofactor activity of PS cannot be demonstrated in mouse plasma using thrombin-generation tests.[15] This
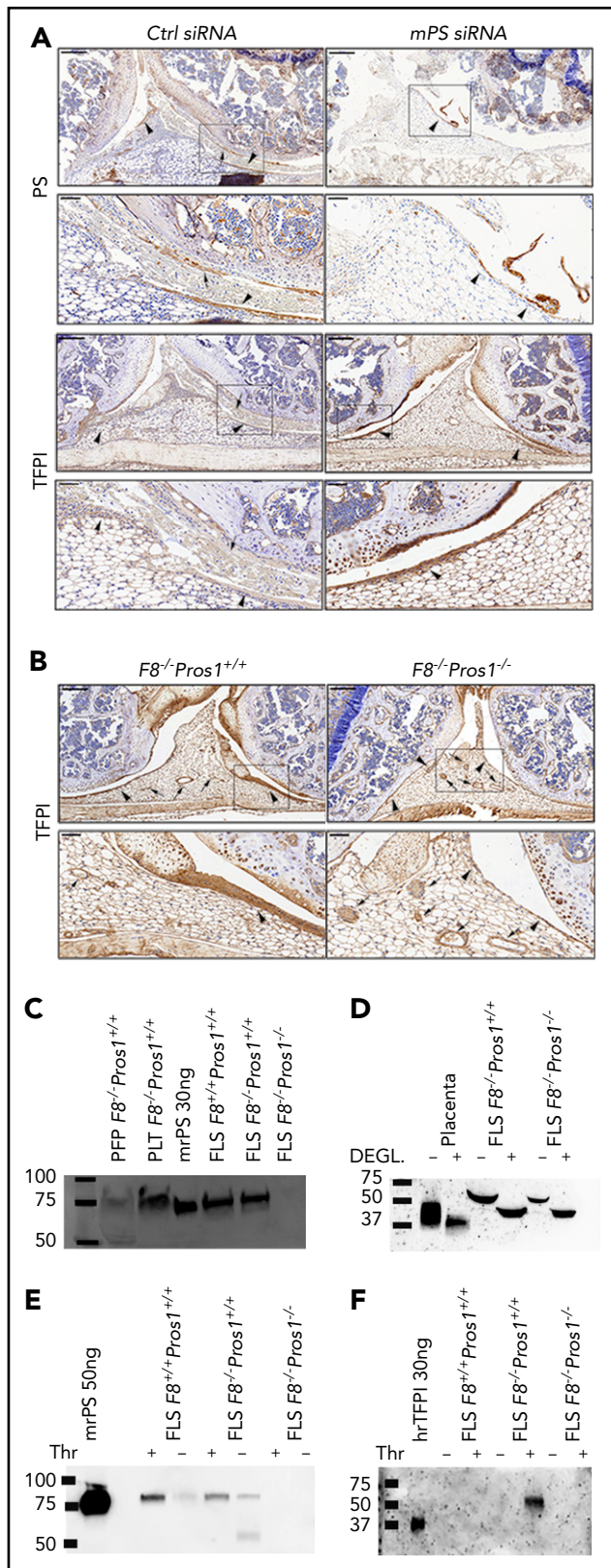
Figure 5. Both PS and TFPI are expressed in murine synovium. (A) Immunostaining for PS and TFPI in the knee intra-articular space of injured knees from $F8^{-/-}$ $Pros1^{+/+}$ mice previously treated with control siRNA or mPS siRNA. Arrowheads point to synovial tissue and arrows to vascular structures, all positive for both PS and TFPI. The boxed areas in the upper subpanels (scale bars: 200 μm) show the area enlarged in the subpanel below (scale bars: 50 μm). (B) Immunostaining for TFPI in the knee

is explained by the lack of TFPIα in mouse plasma and its presence in mouse platelets.[35]

Both thrombin peak and endogenous thrombin potential (ETP) were significantly higher in $F8^{-/-}Pros1^{-/-}$ PRP than in $F8^{-/-}$ $Pros1^{+/+}$ PRP in response to 1 pM TF (1072 ± 160 vs 590 ± 10 nmol/L·min, n = 3 per group, P = .04), suggesting the lack of PS TFPI cofactor activity in $F8^{-/-}Pros1^{-/-}$ PRP (Figure 7A). Consistent with previous work,[15] both thrombin peak and ETP were comparable in PFP of $F8^{-/-}Pros1^{+/+}$ and $F8^{-/-}Pros1^{-/-}$ mice in the presence of 1, 2.5, or 5 pM TF (data not shown).

To assess whether $F8^{-/-}Pros1^{-/-}$ mice exhibited defective functional APC-dependent PS activity, we used thrombin-generation testing in Ca^{2+} ionophore-activated PRP in the absence of APC, in the presence of WT recombinant APC, or in the presence of a mutated (L38D) recombinant mouse APC (L38D APC, a variant with ablated PS cofactor activity).[36] In this assay, APC titration showed that the addition of 8 nM of WT APC was able to reduce ETP by 90% in activated PRP of WT mice, whereas the same concentration of L38D APC diminished ETP by only 30% (data not shown). On the basis of these data, we recorded thrombin-generation curves for activated PRP (3 mice/assay). The calculated APC ratio ($ETP_{+APC\ WT}/ETP_{+APC\ L38D}$) indicated an APC resistance in $F8^{-/-}Pros1^{-/-}$ plasma but not in $F8^{-/-}Pros1^{+/+}$ plasma (0.87 ± 0.13 vs 0.23 ± 0.08, respectively; P = .01) (Figure 7B).

APC-dependent PS activity was also tested in PFP from $F8^{-/-}Pros1^{+/+}$ and $F8^{-/-}Pros1^{-/-}$ mice (2 mice/assay) in the presence of 2 nM of WT APC and L38D APC. The calculated APC ratio showed an APC resistance in $F8^{-/-}Pros1^{-/-}$ mice but not in $F8^{-/-}Pros1^{+/+}$ mice (1.08 ± 0.04 vs 0.25 ± 0.09, respectively; P = .0003) (Figure 7B).

Improved fibrin network in HA mice lacking *Pros1*

Tail-bleeding mouse models are sensitive not only to platelet dysfunction but also to coagulation[37] and fibrinolysis[38] alterations. To understand the differences between studied genotypes regarding tail bleeding, we used scanning electron microscopic imaging to investigate fibrin structure (Figure 7C). Clots from $F8^{+/+}Pros1^{+/+}$ and $F8^{-/-}Pros1^{-/-}$ plasma showed a denser network of highly branched fibrin fibers than did those from $F8^{-/-}Pros1^{+/+}$ plasma clots (supplemental Figure 7A-B). In contrast, clots from $F9^{+/+}Pros1^{+/+}$ and $F9^{-/-}Pros1^{-/-}$ plasma did not display a network denser than that of $F9^{-/-}Pros1^{+/+}$ plasma clots, but a trend for augmented fibers branching (supplemental Figure 7C-D).

Figure 5 (continued) intra-articular space of uninjured knees from $F8^{-/-}Pros1^{+/+}$ and $F8^{-/-}Pros1^{-/-}$ mice. Arrowheads point to synovial tissue and arrows to vascular structures, all positive for both PS and TFPI. The boxed areas in the upper subpanels (scale bars: 200 μm) show the area enlarged in the subpanel below (scale bars: 50 μm). (C-D) Western blot analysis of conditioned media from primary murine fibroblast-like synoviocyte (FLS) cultures using anti-PS (C) and anti-TFPI (D) antibodies. Platelet-free plasma, protein lysates from platelets, and murine PS were used as positive controls (C). TFPI isoform expression was determined by comparing the molecular weights of deglycosylated TFPI and of fully glycosylated TFPI. Murine placenta was used as a positive control for TFPIα. (E-F) Western blot analysis of total protein lysates isolated from FLS after 24 hours of culture in the presence of thrombin (+) or of a vehicle (−) using anti-TFPI (E) and anti-PS (F) antibodies. Human recombinant TFPI full length was used as a positive control for TFPIα. Blots are representative of 3 independent experiments. hrTFPI, human recombinant TFPI; PLT, protein lysates from platelets; Thr, thrombin.
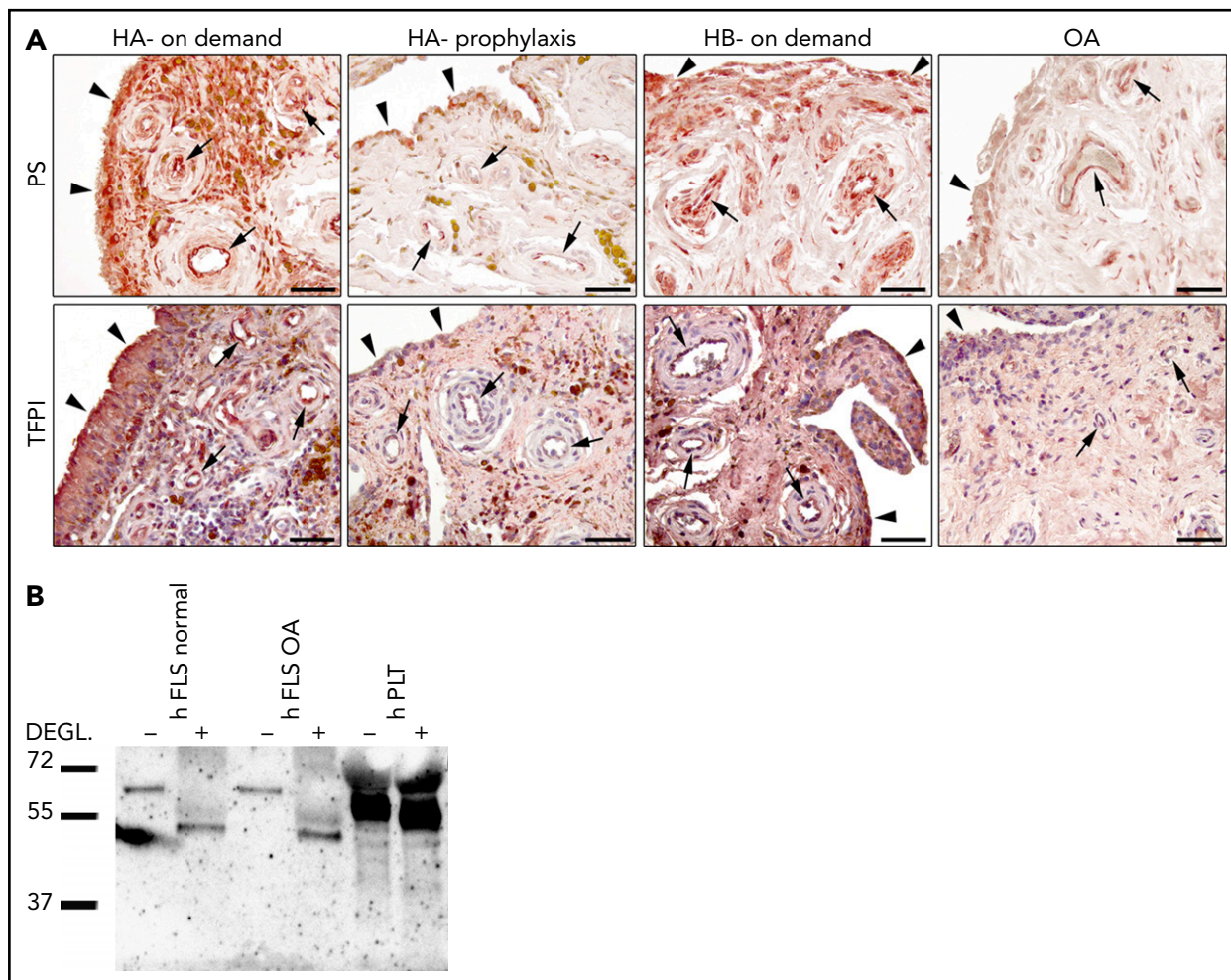
Figure 6. PS and TFPI in human synovium. (A) PS and TFPI are expressed in synovial tissue of patients with HA (on demand and on prophylaxis), HB on demand or osteoarthritis (OA). Arrowheads point to synovial lining layer and arrows to vascular structures in the sublining layer, all positive for both PS and TFPI. Scale bars: 50 μm. (B) Western blot analysis of conditioned media of primary human FLS cultures from a healthy individual and an OA patient before and after deglycosylation using anti-TFPI antibody. Human platelet lysate was used as a positive control for TFPIα. Blots are representative of 3 independent experiments. DEGL., deglycosylation; hFLS, human FLS; hPLT, human platelet lysate; OA, osteoarthritis.

Fibrin fibers from $F8^{-/-}Pros1^{-/-}$ and $F8^{-/-}Pros1^{+/+}$ mice, and from $F9^{-/-}Pros1^{-/-}$ and $F9^{-/-}Pros1^{+/+}$ mice, displayed a larger diameter than did fibers from $F8^{+/+}Pros1^{+/+}$ mice or $F9^{+/+}Pros1^{+/+}$ mice, respectively. Nevertheless, the fiber surface of $F8^{-/-}Pros1^{-/-}$ and $F9^{-/-}Pros1^{-/-}$ mice showed less porosity than did that of $F8^{-/-}Pros1^{+/+}$ or $F9^{-/-}Pros1^{+/+}$ mice, respectively, suggesting that $F8^{-/-}Pros1^{-/-}$- and $F9^{-/-}Pros1^{-/-}$-derived fibers might be less permeable and thereby more resistant to fibrinolysis than are $F8^{-/-}Pros1^{+/+}$- or $F9^{-/-}Pros1^{+/+}$-derived fibers.[39] These data, complementing both TFPI and APC cofactor activity results (Figure 7A-B), help to explain why tail bleeding in $F8^{-/-}Pros1^{-/-}$ mice was improved when compared with $F8^{-/-}Pros1^{+/+}$ mice but not completely corrected as in $F8^{+/+}Pros1^{+/+}$ mice.

PS inhibition in plasma restores thrombin generation in patients with HA

We then examined the effect of PS inhibition on thrombin generation in human HA plasma. ETP in PFP increased 2- to 4-fold in the presence of a PS-neutralizing antibody. Similar results were obtained using an anti-human TFPI antibody against the C-terminal domain for efficient FXa inhibition, even in the presence of an FVIII inhibitor (Figure 7D-E). PS inhibition had a remarkable effect in PRP samples, in which it increased ETP more than 10 times (1912 ± 37 and 1872 ± 64 nM ∗ min) (Figure 7F and Figure 7G, respectively). Thus, PS inhibition completely restored ETP in hemophilic plasma (for comparison, ETP in normal plasma is 1495 ± 2 nM ∗ min). Similar results were obtained using the anti-TFPI antibody (Figure 7D-G). These data confirm in humans the improvement of thrombin generation in HA PFP and PRP driven by PS inhibition that we observed in mice.

Discussion

PS being a key regulator of thrombin generation, we considered that targeting PS could constitute a potential therapy for hemophilia.

Extensive studies in mice provide proof-of-concept data that support a central role for PS and TFPI contributing to bleeding and serious joint damage in hemophilic mice. Targeting *Pros1* or inhibiting PS has the ability to ameliorate hemophilia in mice, as judged by the in vivo improvement of the bleeding phenotype in the tail-bleeding assays and the full protection against hemarthrosis
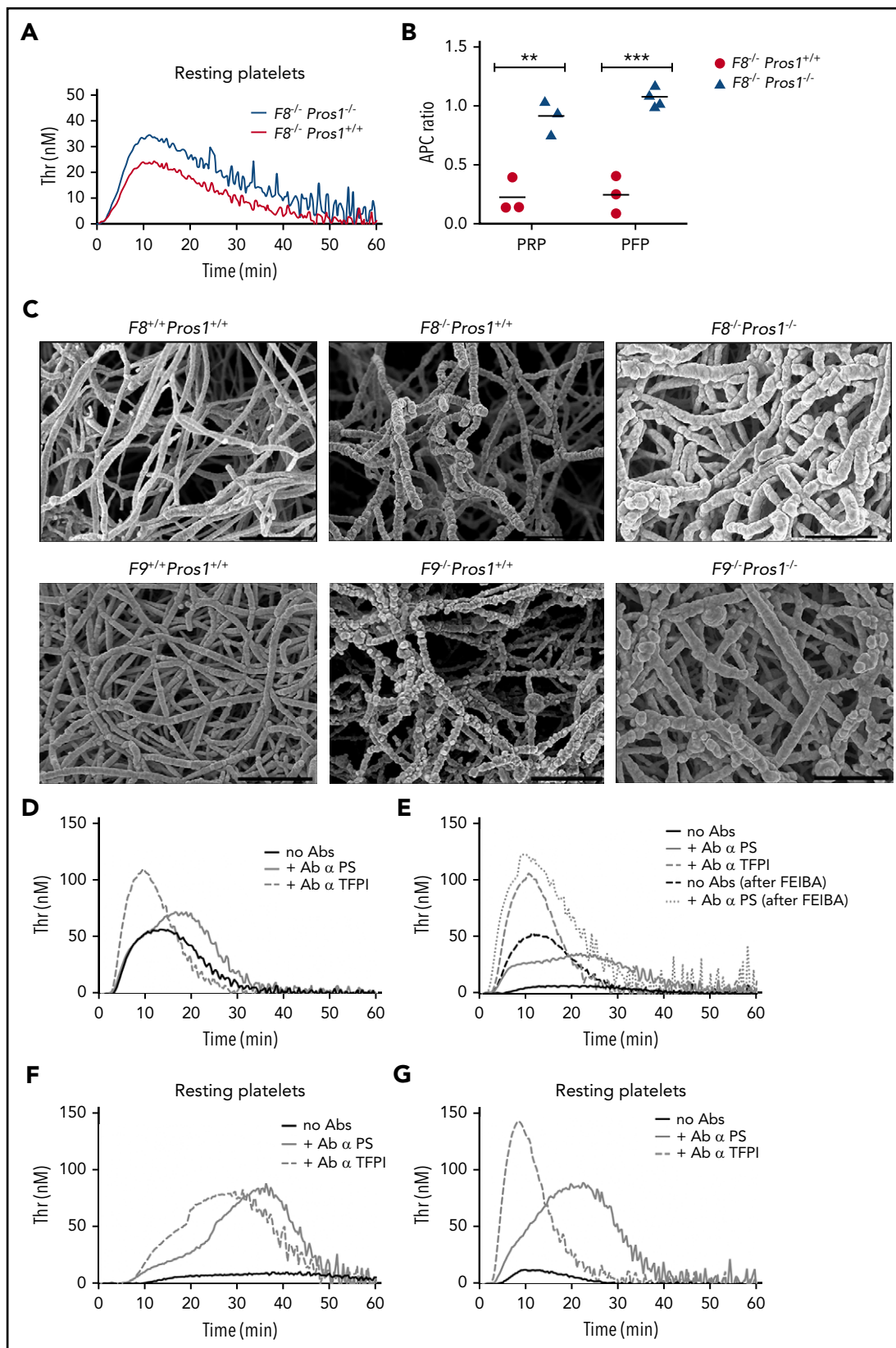
Figure 7. Thrombin generation and fibrin network in hemophilia. (A) TF-induced (1 pM) thrombin generation in PRP from $F8^{-/-}$ $Pros1^{+/+}$ and $F8^{-/-}$ $Pros1^{-/-}$ mice depicting TFPI-dependent PS activity. (B) APC-dependent PS activity in PRP and PFP from $F8^{-/-}$ $Pros1^{+/+}$ and $F8^{-/-}$ $Pros1^{-/-}$ mice. (C) Representative scanning electron microscopy images from $F8^{+/+}$ $Pros1^{+/+}$, $F8^{-/-}$ $Pros1^{+/+}$, and $F8^{-/-}$ $Pros1^{-/-}$ fibrin structure and from $F9^{+/+}$ $Pros1^{+/+}$, $F9^{-/-}$ $Pros1^{+/+}$, and $F9^{-/-}$ $Pros1^{-/-}$ fibrin structure (Scale bars: 1 μm). (D-G) Thrombin generation triggered by low TF concentration (1 pM) in PFP (D-E) and PRP (F-G) from severe HA patients (FVIII < 1%) without (D,F) and with (E,G) a high titer of inhibitor. Measurements are presented as means ± SEM. **$P < .005$; ***$P < .0005$. Ab, antibody; FEIBA, factor eight inhibitor bypassing activity.

(Figures 3A-C and 4). Because joints display a very weak expression of TF[40] and synovial cells produce a high amount of TFPIα and PS (Figure 5), the activity of the extrinsic pathway is greatly reduced intra-articularly, predisposing hemophilic joints to bleeding. Moreover, both thrombomodulin (TM) and endothelial protein C receptor (EPCR) are expressed by FLS,[41,42] suggesting that the TM-thrombin complex activates EPCR-bound PC to generate the very potent anticoagulant, APC, in the context of AH. Importantly, the expression of TFPIα is upregulated by thrombin (Figure 5F). Thus AH, which usually results in marked local inflammation and joint symptoms that can last for days to weeks, also promotes the local generation and secretion of multiple anticoagulants, namely APC, TFPIα, and their mutual cofactor PS, which could help explain the pathophysiology of joint damage in hemophilia.

Observations using clinical samples from hemophilic patients are consistent with the lessons learned from murine studies. In humans, blocking PS in plasma from patients with HA with or without inhibitors normalizes the ETP (Figure 7D-G). Patients with HB display less intra-articular expression of TFPI and PS than do patients with HA, consistent with current knowledge that patients with HB bleed less than do those with HA[22-26,43,44] (Figure 6). Moreover, patients with HA receiving prophylaxis display less TFPI and PS synovial expression than do patients receiving FVIII concentrates only in the context of bleeding, that is, "on-demand therapy" (Figure 6A). These data confirm the lack of PS synovial expression after hrFVIII infusion in F8^{−/−} Pros1^{−/−} mice (supplemental Figure 5). Finally, human FLS secrete both TFPIα and PS as observed in mice, thus strengthening the extrapolation of murine hemophilia data to humans.

The extensive findings in this report lead us to propose that targeting PS may potentially be translated to therapies useful for hemophilia. PS in human and murine joints is a novel pathophysiological contributor to hemarthrosis and constitutes an attractive potential therapeutic target especially because of its dual cofactor activity for both APC and TFPIα within the joints. In the presence of PS, hemarthrosis increases TFPIα expression in the synovia. Targeting PS in mice protects them from hemarthrosis. Thus, we propose that TFPIα and its cofactor PS, both produced by FLS, together with the TM-EPCR-PC pathway, comprise a potent intra-articular anticoagulant system, which has an important pathologic impact on hemarthrosis. The murine PS-silencing RNA that we successfully used in hemophilic mice (Figures 4H-I and 5A) are a therapeutic approach that we would develop for hemophilic patients. The advantage of silencing RNA over current factor replacement therapy is its longer half-life, reducing the frequency of the injections and its possible subcutaneous administration route. Future studies are needed to assess the merits of this new concept for targeting simultaneously the multiple anticoagulant cofactor activities of PS, which involve both APC and TFPI.

Authorship

Contribution: R.P., L. Bologna, M.M., D.M., I.R., N.D., P.A., C.Q., M.D.R.C., H.-U.S., L.I.-M., S.C., and A.A.-S. designed and performed the experiments; L. Bologna, R.P., M.M., D.M., I.R., N.D., S.S., P.A., J.A.F., L. Burnier, C.Q., M.D.R.C., Y.M., J.A.K.H., J.H.G., H.-U.S., L.I.-M., F.S., S.C., and A.A.-S. interpreted the data; R.P., L. Bologna, J.H.G., S.C., and A.A.-S. wrote the manuscript; and J.A.F., L. Burnier, Y.M., J.A.K.H., J.H.G., and F.S. prepared the reagents.

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ORCID profiles: M.M., 0000-0003-3956-8480; L. Burnier, 0000-0001-7767-0334; J.A.K.H., 0000-0002-1300-7135; H.-U.S., 0000-0002-9404-7736; F.S., 0000-0001-5255-2630; A.A.-S., 0000-0003-2872-4863.

Correspondence: Anne Angelillo-Scherrer, Department of Hematology and Central Hematology Laboratory, Inselspital, Bern University Hospital, University of Bern, CH-3010 Bern, Switzerland; e-mail: anne.angelillo-scherrer@insel.ch.

REFERENCES

1. Bolton-Maggs PH, Pasi KJ. Haemophilias A and B. Lancet. 2003;361(9371):1801-1809.

2. Hoyer LW. Hemophilia A. N Engl J Med. 1994;330(1):38-47.

3. Mannucci PM, Tuddenham EG. The hemophilias—from royal genes to gene therapy. N Engl J Med. 2001;344(23):1773-1779.

4. Berntorp E, Shapiro AD. Modern haemophilia care. Lancet. 2012;379(9824):1447-1456.

5. Hartmann J, Croteau SE. 2017 clinical trials update: innovations in hemophilia therapy. Am J Hematol. 2016;91(12):1252-1260.

6. Polderdijk SG, Adams TE, Ivanciu L, Camire RM, Baglin TP, Huntington JA. Design and characterization of an APC-specific serpin for the treatment of hemophilia. Blood. 2017;129(1):105-113.

7. DiScipio RG, Davie EW. Characterization of protein S, a gamma-carboxyglutamic acid containing protein from bovine and human plasma. Biochemistry. 1979;18(5):899-904.

8. Walker FJ. Regulation of activated protein C by a new protein. A possible function for

bovine protein S. *J Biol Chem.* 1980;255(12): 5521-5524.

9. Hackeng TM, Seré KM, Tans G, Rosing J. Protein S stimulates inhibition of the tissue factor pathway by tissue factor pathway inhibitor. *Proc Natl Acad Sci USA.* 2006;103(9): 3106-3111.

10. Ndonwi M, Broze G Jr. Protein S enhances the tissue factor pathway inhibitor inhibition of factor Xa but not its inhibition of factor VIIa-tissue factor. *J Thromb Haemost.* 2008;6(6): 1044-1046.

11. Mahasandana C, Suvatte V, Marlar RA, Manco-Johnson MJ, Jacobson LJ, Hathaway WE. Neonatal purpura fulminans associated with homozygous protein S deficiency. *Lancet.* 1990;335(8680):61-62.

12. Comp PC, Esmon CT. Recurrent venous thromboembolism in patients with a partial deficiency of protein S. *N Engl J Med.* 1984; 311(24):1525-1528.

13. Schwarz HP, Fischer M, Hopmeier P, Batard MA, Griffin JH. Plasma protein S deficiency in familial thrombotic disease. *Blood.* 1984; 64(6):1297-1300.

14. Burstyn-Cohen T, Heeb MJ, Lemke G. Lack of protein S in mice causes embryonic lethal coagulopathy and vascular dysgenesis. *J Clin Invest.* 2009;119(10):2942-2953.

15. Saller F, Brisset AC, Tchaikovski SN, et al. Generation and phenotypic analysis of protein S-deficient mice. *Blood.* 2009;114(11): 2307-2314.

16. Bi L, Lawler AM, Antonarakis SE, High KA, Gearhart JD, Kazazian HH Jr. Targeted disruption of the mouse factor VIII gene produces a model of haemophilia A. *Nat Genet.* 1995; 10(1):119-121.

17. Lin HF, Maeda N, Smithies O, Straight DL, Stafford DW. A coagulation factor IX-deficient mouse model for human hemophilia B. *Blood.* 1997;90(10):3962-3966.

18. Weiss EJ, Hamilton JR, Lease KE, Coughlin SR. Protection against thrombosis in mice lacking PAR3. *Blood.* 2002;100(9):3240-3244.

19. Maroney SA, Cooley BC, Ferrel JP, et al. Absence of hematopoietic tissue factor pathway inhibitor mitigates bleeding in mice with hemophilia. *Proc Natl Acad Sci USA.* 2012;109(10):3927-3931.

20. Øvlisen K, Kristensen AT, Valentino LA, Hakobyan N, Ingerslev J, Tranholm M. Hemostatic effect of recombinant factor VIIa, NN1731 and recombinant factor VIII on needle-induced joint bleeding in hemophilia A mice. *J Thromb Haemost.* 2008;6(6): 969-975.

21. Hakobyan N, Valentino LA, Cong L, et al. Haemarthrosis model in mice: BSS— Bleeding Severity Score assessment system. *Haemophilia.* 2016;22(5):790-798.

22. Biss TT, Chan AK, Blanchette VS, et al; Canadian Association of Nurses in Hemophilia Care (CANHC). The use of prophylaxis in 2663 children and adults with haemophilia: results of the 2006 Canadian National Haemophilia Prophylaxis Survey. *Haemophilia.* 2008;14(5): 923-930.

23. Melchiorre D, Linari S, Manetti M, et al. Clinical, instrumental, serological and histological findings suggest that hemophilia B may be less severe than hemophilia A. *Haematologica.* 2016;101(2):219-225.

24. Nagel K, Walker I, Decker K, Chan AK, Pai MK. Comparing bleed frequency and factor concentrate use between haemophilia A and B patients. *Haemophilia.* 2011;17(6):872-874.

25. Schulman S, Eelde A, Holmström M, Ståhlberg G, Odeberg J, Blombäck M. Validation of a composite score for clinical severity of hemophilia. *J Thromb Haemost.* 2008;6(7): 1113-1121.

26. Tagariello G, Iorio A, Santagostino E, et al; Italian Association Hemophilia Centre (AICE). Comparison of the rates of joint arthroplasty in patients with severe factor VIII and IX deficiency: an index of different clinical severity of the 2 coagulation disorders. *Blood.* 2009; 114(4):779-784.

27. Chattopadhyay R, Sengupta T, Majumder R. Inhibition of intrinsic Xase by protein S: a novel regulatory role of protein S independent of activated protein C. *Arterioscler Thromb Vasc Biol.* 2012;32(10):2387-2393.

28. Gailani D, Renné T. Intrinsic pathway of coagulation and arterial thrombosis. *Arterioscler Thromb Vasc Biol.* 2007;27(12):2507-2513.

29. Fischer K, Collins P, Björkman S, et al. Trends in bleeding patterns during prophylaxis for severe haemophilia: observations from a series of prospective clinical trials. *Haemophilia.* 2011;17(3):433-438.

30. Stephensen D, Tait RC, Brodie N, et al. Changing patterns of bleeding in patients with severe haemophilia A. *Haemophilia.* 2009; 15(6):1210-1214.

31. Piro O, Broze GJ Jr. Comparison of cell-surface TFPIalpha and beta. *J Thromb Haemost.* 2005;3(12):2677-2683.

32. Beltrán-Miranda CP, Khan A, Jaloma-Cruz AR, Laffan MA. Thrombin generation and phenotypic correlation in haemophilia A. *Haemophilia.* 2005;11(4):326-334.

33. Dargaud Y, Béguin S, Lienhart A, et al. Evaluation of thrombin generating capacity in plasma from patients with haemophilia A and B. *Thromb Haemost.* 2005;93(3):475-480.

34. Santagostino E, Mancuso ME, Tripodi A, et al. Severe hemophilia with mild bleeding phenotype: molecular characterization and global coagulation profile. *J Thromb Haemost.* 2010; 8(4):737-743.

35. Maroney SA, Ferrel JP, Pan S, et al. Temporal expression of alternatively spliced forms of tissue factor pathway inhibitor in mice. *J Thromb Haemost.* 2009;7(7):1106-1115.

36. Preston RJ, Ajzner E, Razzari C, et al. Multifunctional specificity of the protein C/activated protein C Gla domain. *J Biol Chem.* 2006;281(39):28850-28857.

37. Dejana E, Quintana A, Callioni A, de Gaetano G. Bleeding time in laboratory animals. III—Do tail bleeding times in rats only measure a platelet defect? (the aspirin puzzle). *Thromb Res.* 1979;15(1-2):199-207.

38. Flight SM, Johnson LA, Du QS, et al. Textilinin-1, an alternative anti-bleeding agent to aprotinin: importance of plasmin inhibition in controlling blood loss. *Br J Haematol.* 2009; 145(2):207-211.

39. He S, Blombäck M, Jacobsson Ekman G, Hedner U. The role of recombinant factor VIIa (FVIIa) in fibrin structure in the absence of FVIII/FIX. *J Thromb Haemost.* 2003;1(6): 1215-1219.

40. Drake TA, Morrissey JH, Edgington TS. Selective cellular expression of tissue factor in human tissues. Implications for disorders of hemostasis and thrombosis. *Am J Pathol.* 1989;134(5):1087-1097.

41. Dargaud Y, Simpson H, Chevalier Y, et al. The potential role of synovial thrombomodulin in the pathophysiology of joint bleeds in haemophilia. *Haemophilia.* 2012;18(5): 818-823.

42. Xue M, March L, Sambrook PN, Fukudome K, Jackson CJ. Endothelial protein C receptor is overexpressed in rheumatoid arthritic (RA) synovium and mediates the anti-inflammatory effects of activated protein C in RA monocytes. *Ann Rheum Dis.* 2007;66(12): 1574-1580.

43. Melchiorre D, Milia AF, Linari S, et al. RANK-RANKL-OPG in hemophilic arthropathy: from clinical and imaging diagnosis to histopathology. *J Rheumatol.* 2012;39(8):1678-1686.

44. Quick AJ, Hussey CV. Hemophilia B (PTC deficiency, or Christmas disease). *AMA Arch Intern Med.* 1959;103(5):762-775.

45. Mann KG, Jenny RJ, Krishnaswamy S. Cofactor proteins in the assembly and expression of blood clotting enzyme complexes. *Annu Rev Biochem.* 1988;57(1):915-956.

46. Broze GJ Jr. The role of tissue factor pathway inhibitor in a revised coagulation cascade. *Semin Hematol.* 1992;29(3):159-169.

47. Repke D, Gemmell CH, Guha A, Turitto VT, Broze GJ Jr, Nemerson Y. Hemophilia as a defect of the tissue factor pathway of blood coagulation: effect of factors VIII and IX on factor X activation in a continuous-flow reactor. *Proc Natl Acad Sci USA.* 1990;87(19): 7623-7627.