

Shear-induced unfolding triggers adhesion of von Willebrand factor fibers

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von Willebrand factor (VWF), a protein present in our circulatory system, is necessary to stop bleeding under high shear-stress conditions as found in small blood vessels. The results presented here help unravel how an increase in hydrodynamic shear stress activates VWF's adhesion potential, leading to the counterintuitive phenomena of enhanced adsorption rate under strong shear conditions. Using a microfluidic device, we were able to mimic a wide range of bloodflow conditions and directly visualize the conformational dynamics of this protein under shear flow. In particular, we find that VWF displays a reversible globule-stretch transition at a critical shear rate $\dot{\gamma}_{\text{crit}}$ in the absence of any adsorbing surface. Computer simulations reproduce this sharp transition and identify the large size of VWF's repeating units as one of the keys for this unique hydrodynamic activation. In the presence of an adsorbing collagen substrate, we find a large increase in the protein adsorption at the same critical shear rate, suggesting that the globule unfolding in bulk triggers the surface adsorption in the case of a collagen substrate, which provides a sufficient density of binding sites. Monitoring the adsorption process of multiple VWF fibers, we were able to follow the formation of an immobilized network that constitutes a "sticky" grid necessary for blood platelet adhesion under high shear flow. Because areas of high shear stress coincide with a higher chance for vessel wall damage by mechanical forces, we identified the shear-induced increase in the binding probability of VWF as an effective self-regulating repair mechanism of our microvascular system.

blood flow | mechanical activated proteins | polymer physics

Our circulatory system is exposed to an amazingly wide span of shear rates, ranging from 1 to 10^5 s^{-1} (1). This range calls for adhesion mechanisms during blood clotting that are different for different regimes of shear rates. At small shear rates, which are found in rather wide vessels, objects such as vesicles or cells bind to vessel walls once the contact area and the adhesion strength is large enough (2). At high shear rates, which are found in small arteries, hydrodynamic lift forces inhibit formation of a sufficiently large contact area, which makes adsorption of soft objects from the blood onto vessel walls increasingly difficult. In fact, theoretical analyses predict that adhering compact and soft objects (such as vesicles or platelets) will roll and detach from a surface at a particular shear stress and, more importantly, will remain unbound if the shear rate is increased further (3, 4). Experiments on leukocyte adhesion confirmed these theoretical predictions quite nicely (2). However, the experimental finding for blood platelet adhesion in small arteries contradicts this scenario: von Willebrand factor (VWF)-mediated platelet adhesion, which is necessary to stop bleeding in small vessels, is strongly enhanced under high shear-flow conditions (5). Therefore, this example of shear-induced adsorption must be explained by a distinct mechanism. Under normal conditions, the glycoprotein VWF assembles into multimeric polymers that, when stretched, can reach a length of up to $100 \mu\text{m}$ (6). The repeating unit is unusually large ($\approx 70\text{-nm}$ long axis, 10-nm short axis) (7) and represents a dimer containing 5,100 aa (8). A number of recent studies on VWF suggest that a conformational

transition, presumably triggered by an increase in shear, may be necessary for the initiation of platelet adhesion (5, 9–11). Former studies on the dynamics of uncollapsed biopolymers in good solvent conditions predict mechanically induced transitions for shear rates in the range of a few 10 s^{-1} (12). In contrast, typical shear rates in small arteries, where procoagulatory activity of VWF is predominant, are in the range of $1,000 \text{ s}^{-1}$. Until now, there have been no studies of which we are aware that mimic the hydrodynamic conditions of our microcirculatory system and directly monitor the conformational dynamics of unbound and conformationally unrestricted VWF when exposed to shear flow. Therefore, the origin of VWF's peculiar hydrodynamic behavior and related function remained unclear on both dynamical and molecular grounds.

Here, we unravel VWF's response to hydrodynamic stress by exploiting a microfluidic flow chamber incorporated directly on a chip. We found that the application of unusually high shear flows ($>1,000 \text{ s}^{-1}$) results in an abrupt elongation of VWF fibers in solution. Computer simulations clearly indicate that the molecular origin of VWF's extraordinarily response to mechanical stress can be tracked back to its extremely large repeating unit size. The conformational transition takes place in a reversible manner and on very short time scales. It must therefore not be mistaken with protein denaturation. The results presented here comprise a thorough investigation of VWF's behavior under strong shear flow and enable us to bridge the gap between physical forces, physiological function, and VWF-related diseases.

Results and Discussion

Shear-Induced Stretching. VWF was diluted in PBS (including 1 mM Mg and 1 mM Ca) that was then spread over the hydrophilic track of the flow chamber chip described in *Materials and Methods* (Fig. 1). Images of VWF at concentrations typical for human blood ($c \approx 2 \mu\text{g/ml}$) are presented in Fig. 2. At shear rates of $\dot{\gamma}_{\text{crit}} \approx 10\text{--}1,000 \text{ s}^{-1}$, the biopolymer exhibits a compact conformation (Fig. 2a Left). The size of the VWF globules was estimated within our fluorescence setup to be $d \approx 2 \mu\text{m}$, clearly showing that it consists of more than one repeating unit. This compact conformation remains unchanged as long as the shear rate is maintained below a certain (critical) value $\dot{\gamma}_{\text{crit}} \approx 5,000 \text{ s}^{-1}$. Increasing the shear rate above $\dot{\gamma}_{\text{crit}}$ induces a shape transformation of the VWF fibers from a collapsed conformation to a stretched conformation with a length, l , of $\approx 15 \mu\text{m}$ (Fig.

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Abbreviation: VWF, von Willebrand factor.

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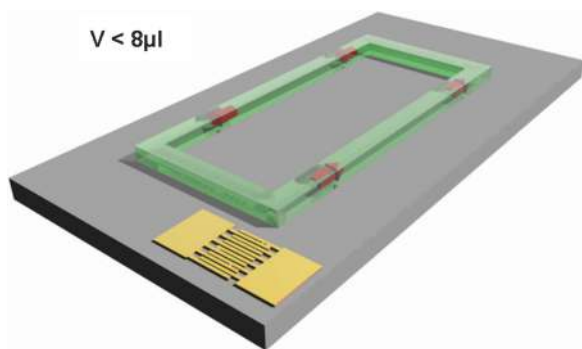


Fig. 1. The planar microfluidic device. The blood vessel is modeled by a planar hydrophilic track on an otherwise hydrophobic piezoelectric substrate. A surface acoustic wave, excited electrically, eventually interacts with the confined liquid at the solid–liquid interface and drives the liquid to flow (acoustic streaming). The surface wave basically acts as a localized pump, because its mechanical energy is absorbed by the liquid over only a few micrometers. Due to the small scales of this microfluidic system, it creates a homogenous laminar flow (low Reynold’s number) along the channel, mimicking the blood flow in arteries or capillaries.

2a Right). Fig. 2b shows that this transition is reversible, because the protein immediately relaxed to its compact conformation when the flow was turned off (at time $t = 0$ ms).

The abrupt nature of the transition can also be seen as a sudden increase in the average extension of the fibers as a function of shear rate, as shown in Fig. 3. These features (reversibility and critical stretching) make VWF a mechanically switchable protein, which, as will be shown, is crucial to its ability to stop bleeding.

The fact that the transition occurs at such a high shear rate cannot be understood from previous studies on linear chains under good solvent conditions (13, 14). For example, it has been shown that DNA of a length that is roughly the same as the VWF fibers studied here will exhibit drastic changes of elongation for shear rates as low as 30 s^{-1} at a viscosity of 1 cP (1 P = 0.1 Pa·sec) (i.e., water or the PBS used here) (12). Our studies, however, indicate that significant changes in conformation occur only at shear rates $\dot{\gamma}_{\text{crit}} \geq 5,000 \text{ s}^{-1}$. This value is more than two orders of magnitude higher than that reported for DNA. To explain this large discrepancy, it is necessary to account for the presence of attractive interactions between repeating units (presumably hydrophobic or H bonds) that would hold each VWF fiber tightly together, even under strong shear conditions. Therefore, we propose that a single VWF multimer forms a compact or collapsed structure similar to a folded protein (Fig. 2a Left). The difference to a regular protein is that here, all of the repeating units are attracted to each other most likely without any sequence-specific binding preference, assuring the reversible character of the transition, as experimentally confirmed in Fig. 2b.

Hydrodynamic Computer Simulations. To gain more insight into the unfolding mechanism, we perform hydrodynamic simulations of a single polymer in shear flow under different solvent conditions. Our simulation method incorporates hydrodynamics and solvent flow effects (15). The effective monomer–monomer attraction is modeled by a Lennard–Jones potential (16). Note that the monomers forming the polymer in our simulation are spherical beads of radius a and correspond to the repeating unit forming the VWF in our experiment. In the simulations, the only parameters are three dimensionless numbers, namely the rescaled shear rate $\dot{\gamma}\tau$ (where $\tau = 6\pi\eta a^3/k_B T$ is the characteristic diffusion time of a single bead); N , the number of repeating units; and the interaction potential strength, ΔU , between two

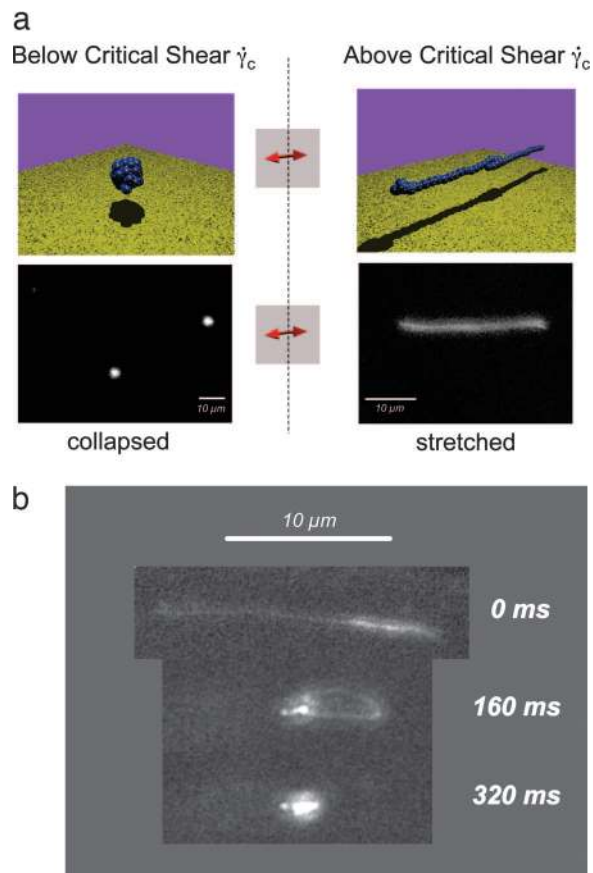


Fig. 2. Dynamic conformational change of VWF under shear. (a) Cartoon (Upper) and selected fluorescence (Lower) images of a video sequence (SI Movie 1, acquired at 25 frames per second) of VWF below and above the critical shear rate of a few thousand s^{-1} . Above $\dot{\gamma}_{\text{crit}}$, the protein is in an elongated conformation. The little yellow spots on the protein are meant to represent a variety of known binding sites. To assure that only unbound VWF was detected, the focus of the microscope was set $\approx 10 \mu\text{m}$ above the surface of the chip in the middle of the channel. (b) Relaxation of VWF. Selected fluorescence images of a video sequence (SI Movie 1, acquired at 25 frames per second). Once the hydrodynamic shear force is reduced below $\dot{\gamma}_{\text{crit}}$ at 0 ms, the protein immediately relaxes back to its compact state. The unfolding process is reversible and presumably does not include protein denaturation.

beads (in units of $k_B T$). Our results for the normalized extension (i.e., the average extension divided by the contour length) of a 50-mer are presented in Fig. 3d Inset. The left (open circles) and right (open squares) curves correspond to a polymer in the coil and globular states, respectively. The attractive well depth (in units of $k_B T$) between monomers for the coil and collapsed case are $\Delta U = 0.416$ and $\Delta U = 2.08$, respectively. On the lower horizontal axis in Fig. 3d Inset, we show the rescaled units for the shear rate, $\dot{\gamma}\tau$, that we use in our simulations. Note that the monomer radius a and solvent viscosity η only enter this rescaled shear rate, and thus our simulations are valid for arbitrary monomer radius a and solvent viscosity η . In the upper horizontal axis in Fig. 3d Inset, we restore physical units by making a specific choice for these parameters, namely a repeating unit or radius of $a = 80 \text{ nm}$ in a fluid with a viscosity similar to that used in the experiments, i.e., $\eta = 1 \text{ cP}$. We have purposely chosen a relatively large value for the bead (monomer) size so that the critical shear rate for the collapsed polymer is close to the experimental value. As shown by the upper scale in Fig. 3d Inset, the extension of the coiled polymer (circles) grows for $\dot{\gamma} > 10 \text{ s}^{-1}$ (vertical dotted line) and saturates at a value of approximately

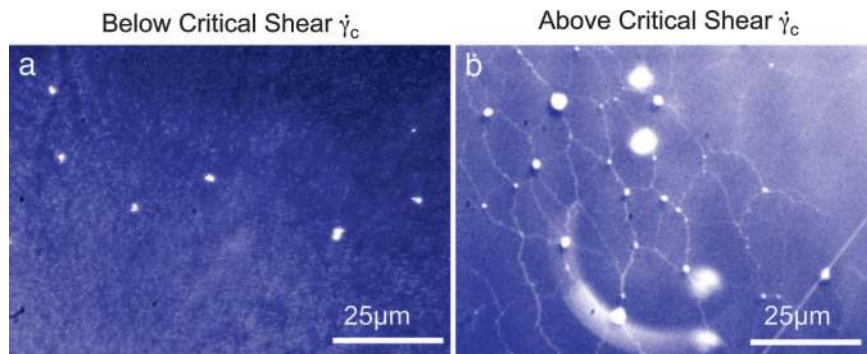


Fig. 4. Adhesion of VWF. (a) Below the critical shear rate $\dot{\gamma}_{crit}$, only few VWF coils adhere to the collagen coated surface. (b) VWF adhered at high shear rates forms a spider web like network, which represents a very adhesive substrate for blood platelets.

The qualitative behavior of the binding process (a rather sharp transition at a certain shear rate) and the order of magnitude of $\dot{\gamma}_{crit}$ is similar to the VWF–collagen interaction presented in Fig. 4. However, our first results clearly indicate that the absolute value of $\dot{\gamma}_{crit}$ is reduced by at least 25% ($<3,500 \text{ s}^{-1}$), probably due to an increased dragging force once platelets start to bind to the protein surface. The formation of aggregates between blood platelets and soluble VWF raised the question of why (to our knowledge) no such aggregates have been found under physiological conditions. There are probably two reasons. From a hydrodynamical point of view, the high shear flow conditions necessary to initiate the blood platelet–VWF associations are rather rare away from the surface, decreasing the probability to find the protein in the haemostatically active (stretched) conformation. From a biochemical perspective, the presence of the VWF degrading enzyme ADAMTS13 may avoid aggregate formation by cleaving VWF (20). Indeed, when we added ADAMTS13 into our microfluidic channel and exposed it to high shear, the formation of an adhesive VWF web was significantly decreased [see supporting information (SI) Fig. 6]. Taking both arguments together, we conclude that the *in vivo* association of VWF with blood platelets under bulk conditions is a rather unlikely scenario when compared with immobilized VWF–platelet association. However, under rather pathophysiological conditions (e.g., ADAMTS13 deficiency or stenotic conditions) the probability of VWF–platelet association is significantly increased (21).

Conclusion

To summarize, we have identified VWF fibers as hydrodynamically activated biopolymers, being haemostatically active only under high shear rates. We point out that the observed critical activation threshold ($\dot{\gamma}_{crit} \approx 5,000 \text{ s}^{-1}$) falls in the same range as the wall shear rate found in small vessels $\dot{\gamma}_{crit} > 1,000 \text{ s}^{-1}$ (1). Therefore, the probability for VWF to undergo a discontinuous, conformational transition and thereby become hemostatically

active increases dramatically under rather harsh hydrodynamic conditions, for which mechanically induced vessel damage due to high wall shear rates is more likely. This finding unravels the dynamics of VWF as the foundation of a self-adjusting repair mechanism for vascular injury under strong shear conditions and provides clues for understanding vessel and clotting dysfunctions, such as arteriosclerosis, thrombosis, and von Willebrand disease.

Generalizing these results, we envision the possibility for engineering functional polymers that could serve, for example, as local shear stress detectors. A second application would be smart materials that, as additives, would automatically detect and seal small holes in pipes at sights of high mechanical stress.

Materials and Methods

The Flow Chamber Chip. To study the hydrodynamic response of VWF, we used a planar microfluidic device (described in Fig. 1) (22) that mimics blood flow in small blood vessels (SI Movie 4). Applying a high-frequency signal to a set of electrodes (interdigital transducers) on a piezoelectric substrate (LiNbO_3) excites a surface acoustic wave. When the wave hits the solid–liquid interface, it transmits its acoustic energy into the liquid over a very small distance. The resulting pressure gradient acts like a localized nanopump, a phenomenon called acoustic streaming (23, 24).

To confine the water laterally, a hydrophilic channel on an otherwise hydrophobic substrate (silanized with octadecyltrichlorosilane) is created by optical lithography. In this way, we are able to design a microfluidic flow chamber on the chip surface. To keep the standard protocols for surface biofunctionalization, the chip surface is sputtered with a 20-nm-thick layer of glass (SiO_2). Because of the fast attenuation of the surface acoustic wave once it enters the water channel, this nanopump does not modify the conformation of molecules adsorbed or grafted to the substrate surface. Additional experimental advantages are the exact tunability of the applied shear rate (we checked by

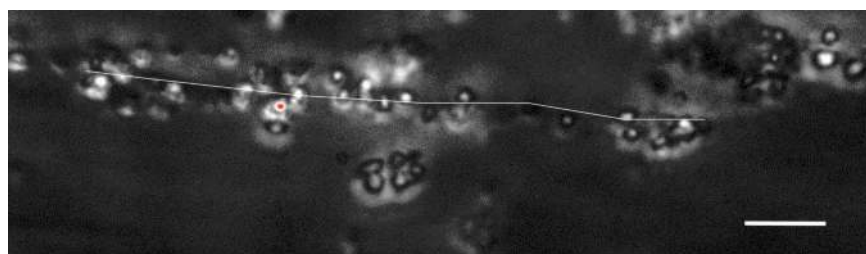


Fig. 5. Blood platelet–VWF aggregation under high shear flow ($\dot{\gamma} \approx 4,000 \text{ s}^{-1}$). Blood platelets ($\approx 1 \mu\text{m}$) are immobilized on the free end of a VWF fiber only when $\dot{\gamma} \approx 3,500 \text{ s}^{-1}$. The white line indicates the underlying VWF and is given as a guide to the eye (see also SI Fig. 7). (Scale bar, $10 \mu\text{m}$.)

measuring the shear rate with tracer particles), and the optical transparency of the piezoelectric substrate.

By mounting the microfluidic chip on a fluorescent microscope, we are able to directly visualize and track the conformation of individual fluorescently labeled VWF multimers freely flowing in the channel (SI Movie 1).

Labeling of VWF. A tetrafluorophenyl ester was used to attach the fluorophore Alexa Fluor 488 (Invitrogen, Karlsruhe, Germany) to the primary amines of the protein. Dye and protein were incubated and dialyzed according to the Invitrogen protocol. VWF was purchased from Calbiochem (La Jolla, CA).

Fluorescence Microscopy and Image Acquiring. Fluorescence microscopy is performed by using an inverted microscope equipped with a mercury lamp (Zeiss, Göttingen, Germany) and a CCD camera (Hamatsu, Photonics Deutschland, Hersching, Germany). Usually, images are taken at a rate of 25 frames per second. To avoid artifacts when determining the protein length while acquiring the image, small amounts of fluorescently labeled beads of a different absorption wavelength than VWF are added. From the measured speed of the beads and their spread

in the microscope image (caused by the finite exposure time), the actual protein length without any artifacts due to the finite exposure time is deduced.

The rate of adhesion is measured by simply counting the increase of light intensity on a fixed area of the chip surface as a function of time (intensity/m²·s) and define 1.0 as the maximum increase in fluorescence intensity per area and time.

Buffers. All experiments are performed in PBS (including 1 mM Mg and 1 mM Ca).

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