



HAL
open science

Von Willebrand Disease: What does the future hold?

Cécile V Denis, Sophie Susen, Peter J. Lenting

► **To cite this version:**

Cécile V Denis, Sophie Susen, Peter J. Lenting. Von Willebrand Disease: What does the future hold?. Blood, 2021, 137 (17), pp.2299-2306. 10.1182/blood.2020008501 . hal-03267627

HAL Id: hal-03267627

<https://hal.science/hal-03267627>

Submitted on 22 Jun 2021

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

Von Willebrand Disease: What does the future hold?

Cécile V. Denis¹, Sophie Susen², and Peter J. Lenting¹,

¹Laboratory for Hemostasis, Inflammation & Thrombosis, Unité Mixed de Recherche S1176, Institut National de la Santé et de la Recherche Médicale, Université Paris-Saclay, 94276 Le Kremlin-Bicêtre, France

²CHU Lille, Université de Lille, Institut National de la Santé et de la Recherche Médicale Unité 1011, Institut Pasteur de Lille, Lille, France; CHU Lille, Hematology and Transfusion, Lille, France

Correspondence: Peter J. Lenting, INSERM U1176, 80 rue du General Leclerc, 94276 Le Kremlin-Bicêtre, France
Tel: +33149595651; Fax: +33146719472;
Email: peter.lenting@inserm.fr

Short title: Von Willebrand disease: What does the future hold?

Article type: Perspective

Abstract: 242 words

Main text (without figure legends): 4000 words

Figures: 2

References: 86

1 **Abstract**

2 Von Willebrand disease (VWD) is characterized by its heterogeneous clinical manifestation,
3 which complicates its diagnosis and management. The clinical management of VWD has
4 remained essentially unchanged over the last 30 years or so, using von Willebrand factor (VWF)
5 concentrates, desmopressin and anti-fibrinolytic agents as main tools to control bleeding. This
6 is in contrast to hemophilia A, for which a continuous innovative path has led to novel
7 treatment modalities.

8 Despite current VWD management being considered effective, quality-of-life studies
9 consistently reveal a higher than anticipated burden of VWD on patients, which is particularly
10 true for women. Apparently, despite our perceived notion of current therapeutic efficiency,
11 there is space for innovation with the goal to reach superior efficacy. Developing innovative
12 treatments for VWD is complex, especially given the heterogeneity of the disease and the
13 multifunctional nature of VWF. In this perspective article, we describe several potential
14 strategies that could provide the basis for future VWD treatments. These include genetic
15 approaches such as gene therapy using dual-vector adeno-associated virus and transcriptional
16 silencing of mutant alleles. Furthermore, protein-based approaches to increase FVIII levels in
17 VWD-type 3 or 2N patients will be discussed. Finally, antibody-based options to interfere with
18 VWF degradation (for congenital VWD-type 2A or acquired Von Willebrand syndrome-type 2A)
19 or increase endogenous VWF levels (for VWD-type 1) will be presented. By highlighting these
20 potential strategies, we hope to initiate an innovative path, which ultimately would allow us
21 to better serve VWD patients and their specific needs.

22

23

1

2 **Introduction**

3 Von Willebrand disease (VWD) is an inherited bleeding disorder resulting from quantitative or
4 qualitative deficiencies in von Willebrand factor (VWF), a plasma multimeric glycoprotein
5 controlling platelet adhesion and aggregation.¹ Although VWD is expected to affect men and
6 women equally, there are more women diagnosed with VWD due to gynecological-related
7 hemostatic challenges. The real prevalence of VWD has been a matter of debate for many
8 years, ranging from as high as 1.3% when taking into account epidemiological studies² to 100
9 cases per million when considering patients with clinically relevant bleeding symptoms.³
10 VWD is traditionally classified in three types. VWD-type 1 is characterized by a partial
11 quantitative deficiency in VWF. VWD-type 2 refers to qualitative VWF defects with a number
12 of subtypes (2A, 2B, 2N and 2M) according to the patients' functional defect. Finally, VWD-
13 type 3, also called severe VWD, is manifested by a virtually complete deficiency in VWF.¹
14 Bleeding symptoms are usually considered less serious than hemophilia-related symptoms.
15 Although that may be true for mild VWD patients, specific comparative studies revealed that
16 the bleeding-profile between VWD-type 3 and severe hemophilia A is more similar than
17 expected.^{4,5} Nevertheless, prophylactic treatment remains relatively rare in VWD patients.
18 To develop alternative treatments for managing VWD in the 21st century, it will be crucial to
19 understand the challenges faced by VWD patients and to take into account the disease
20 specificities.

21

22 **Challenges in the diagnosis of VWD**

23 Given the heterogeneity of the disease that is observed sometimes within the same family or
24 even in the same patient throughout his/her lifetime, VWD has been and remains to this day,
25 a complex entity to diagnose⁶ and to manage, partially explaining the difference between
26 prevalence and actually diagnosed patients. Laboratory testing of VWD outside highly
27 specialized centers is a challenge in itself, and the link between mild phenotypes and a VWF
28 defect is not always straightforward. In addition, bleeding history relies on a subjective
29 interpretation by the patients, their parents as well as the clinician. This is particularly true
30 with regard to menstrual bleeding, a subject not easily discussed both inside and outside the
31 family circle. Drs. Weyand and James recently noted that while 76% of men with VWD are
32 diagnosed by 10 years, 50% of the women are not diagnosed by 12 years of age.⁷ Although

1 this could partially be explained by a first bleeding episode at different ages, it has also been
2 shown that the diagnostic delay following the first bleeding episode is longer in women than
3 men.⁸ Delayed diagnosis may result in missed treatment opportunities and increased
4 morbidity (both medical and psychological). It is thus important to increase knowledge and
5 awareness of VWD among (general) clinicians and public.

6

7 **VWD and quality of life**

8 What objective tools do we actually rely on to define that VWD is a mild bleeding disorder in
9 most cases? First and primary is the clinical management of patients, since many VWD
10 patients are followed in specialized hemophilia treatment centers. The health care
11 professionals managing patients are highly-trained hematologists who have the necessary
12 expertise to evaluate the disease burden. However and as stated maybe provocatively by Dr.
13 Kouides “VWD on clinical grounds is primarily a disease of females”.⁹ We are now increasingly
14 learning how gender bias may have unwillingly affected patient’s care in many areas.¹⁰⁻¹²
15 Whether such an issue has contributed to underestimate VWD severity in terms of physical
16 and psychological consequences is unclear. Yet, there seems to be a parallel between the
17 increasing number of female hematologists and the greater recognition of significant VWD-
18 associated morbidities, especially when related to women reproductive health.¹³⁻¹⁵
19 The second tool to evaluate disease severity are health-related quality of life (HR-QoL) studies,
20 widely accepted as crucial instruments of patient-centered care.¹⁶ For VWD, a few studies
21 have addressed this issue using either 1) validated HR-QoL questionnaires, 2) questionnaires
22 not officially validated or 3) through interviews. All studies have reported significantly
23 decreased quality of life in VWD patients. The largest study (509 patients) was performed in
24 the Netherlands using the generic Short Form (SF)-36 and reported decreased vitality (score
25 measuring energy and fatigue) in both men and women with VWD.¹⁷ Smaller studies
26 performed in Canada (102 patients, SF-36 questionnaire), Finland (47 patients, Medical
27 Outcomes Study-36 questionnaire), Sweden (30 female patients, SF-36 questionnaire),
28 Canada again (28 patients, Health Utility Index Mark-2 and Mark-3) or United Kingdom (57
29 female patients, general questionnaire) not only confirmed a lower quality of life for VWD
30 patients compared to the general population, but for studies in which both men and women
31 were included, most reported a difference between both genders.¹⁸⁻²² Studies focusing
32 primarily on women clearly indicate how the burden of VWD is borne disproportionately by

1 women. Heavy menstrual bleeding and its associated pain (observed in 32-100% of patients),²³
2 post-partum hemorrhage (in 15-60% of patients),¹³ and hysterectomy (performed in 10-26%
3 of patients),²⁴ are common in VWD women.^{15,25} Qualitative interview-based studies about HR-
4 QoL or lived experiences are even more striking with women reporting a considerable
5 impairment in their lives.²⁶⁻²⁸ Obviously not all VWD patients are affected by severe bleeding
6 symptoms, a fact that can mitigate some of the HR-QoL studies where only a modest impact
7 was reported, such as in the large Dutch study. Focusing on subgroups can probably give
8 better ideas of the situation. A perfect example is illustrated in another Dutch study focusing
9 on VWD patients with joint bleeds.²⁹ The authors show that 23% of patients from the Dutch
10 VWD cohort self-reported joint bleeds. Not only is this percentage higher than anticipated,
11 but joint bleeds were reported in VWD-type 1 as well as in VWD-type 2 or VWD-type 3, even
12 if factor VIII (FVIII) levels exceeded 10%. Joint bleeds were associated with increased
13 consumption of factor concentrates, joint pain and lower HR-QoL, suggesting a “significant
14 burden of arthropathy in VWD”.²⁹

15 Another factor that can impact the results of the HR-QoL studies is the absence of disease-
16 specific instruments.²³ All the validated HR-QoL questionnaires used so far are generic
17 instruments. A VWD-specific QoL has been developed,³⁰ but although it was mentioned that
18 the questionnaire was validated in a German study (Wil-QoL) and a French study (WISH-QoL),
19 no publications are yet available.

20 Altogether, these different studies illustrate the higher than anticipated burden of VWD on
21 patients and suggest that VWD classification in types 1, 2 and 3, although crucial to understand
22 the physiopathology of the disease is not always a good proxy for clinical severity. The actual
23 patient’s phenotype is not only driven by the VWD phenotype, but also by burden in terms of
24 hospitalizations, surgical interventions, drug consumption, gynecological visits, sick leaves and
25 other objective parameters. This burden is only in part included in the bleeding score used to
26 characterize the disease (Bleeding Assessment Tool of the International Society on
27 Thrombosis and Hemostasis). However, this tool is cumulative and more adapted to diagnosis
28 rather than reevaluation of the bleeding tendency. To better understand the evolution of the
29 patient’s phenotype and associated burden, an alternative score could be designed to
30 determine spontaneous symptomatology in the presence and absence of treatment.
31 Undoubtedly, despite our perceived notion of therapeutic efficacy of current VWD treatments,

1 improved management of patients is necessary to address patient's needs in an optimal
2 fashion.

3

4 **Current treatment**

5 Over the last 30 years, many reviews related to the clinical management of VWD have been
6 published (see *eg.* ^{1,24,31-38}) and guidelines were recently published.³⁹ Without extensively
7 revisiting the topic, some relevant issues can be highlighted.

8 The most relevant message is that the toolkit for the treatment of VWD has remained
9 essentially unchanged over these 30 years. Already in 1991, Dr. Aledort described the use of
10 intermediate-purity factor concentrates and desmopressin as main approaches for the
11 management of VWD, as well as adjunctive anti-fibrinolytic and estrogenic therapy.³¹ It then
12 took until 2005 for the development of a purer plasma-derived VWF concentrate, poor in FVIII
13 activity.⁴⁰ As for the first recombinant VWF concentrate, it is only since 2015 that it became
14 available in the USA and 2017 in Europe.⁴¹

15 When establishing a parallel with progresses made in the clinical management of hemophilia
16 A over the same period of time, the contrast is particularly striking (Figure 1). While a parallel
17 development was observed in terms of intermediate-purity concentrates (which contain both
18 VWF and FVIII) and desmopressin in the 1980s, a real divergence started in the 1990s with the
19 development of monoclonal antibody-purified and recombinant FVIII concentrates. At that
20 time, who would have guessed that it would take another 20 years for comparable products
21 to be available for VWD? But research did not stand still in the hemophilia field and while
22 VWD management was only trying to fill the gap, nothing short of a revolution was looming
23 on the other side of the VWF/FVIII partnership. Indeed, during the last decade, hemophilia A
24 treatment has soared with several novel therapeutic strategies being available or in late
25 clinical development.⁴²⁻⁴⁵ In particular, gene therapy is reaching maturity and therefore for
26 the first time in history, a real cure for the disease is within sight.

27 Thus, innovation in VWD treatment has basically stopped and the limitations of current
28 treatments have not been addressed. Of course, the arrival of a recombinant VWF
29 concentrate is an important step forward. However, without neglecting its specific merits, the
30 availability of this recombinant VWF concentrate is not changing the essentials of VWD
31 management, and prophylactic treatment will still require multiple intravenous infusions *per*
32 week. The invasive nature of intravenous administration is associated with an increased risk

1 of infections, it is inconvenient, and it is challenging in the very young and old to find adequate
2 venous access for repeated injections. Furthermore, prophylactic treatment using
3 concentrates is expensive and not affordable world-wide. Finally, a major hindrance to the
4 use of prophylaxis is that the criteria for introduction of prophylaxis for VWD patients
5 (particularly women with heavy menstrual bleeding) are not as well defined as for patients
6 with hemophilia. As a consequence, the benefit or need for prophylaxis are less well
7 recognized.⁴⁶

8 The advantages of desmopressin are numerous, and validate its status as the “treatment of
9 choice” in VWD-type 1 patients. It can be administered in different ways (IV, subcutaneous or
10 intranasal), is relatively cheap, and without risk of transmittable disease. It will transiently
11 achieve adequate post-treatment levels of VWF and FVIII in 80% of cases.^{47,48} However, there
12 are also some limitations. It has variable effectiveness in type 2A & 2M, and is usually
13 considered for minor bleeding/surgery.⁴⁷ Moreover, this medication is contra-indicated for
14 VWD-type 2B, while its application in VWD-type 3 is ineffective since no FVIII or VWF is
15 released from the endothelial storage-organelles.⁴⁷ Importantly, desmopressin is primarily a
16 antidiuretic, with VWF release being an off-target effect. A number of side-effects
17 concomitant to desmopressin use have also been reported (transient headaches, facial
18 flushing, hypotension, hyponatremia and mild tachycardia), which are generally mild and well-
19 tolerated.^{49,50} Nevertheless, a recent VWD coping survey presented by Kalot *et al.* revealed
20 that patients/caregivers display more concern about the side effect profile of desmopressin
21 compared to health care professionals, and seem to prefer a prophylactic approach.⁵¹
22 Furthermore, the repetitive use of desmopressin may result in reduced responsiveness
23 (tachyphylaxis) due to exhaustion of the VWF storage-organelles.⁵² Finally, it is relevant to
24 mention that desmopressin is a monopolized product by Ferring Pharmaceuticals, which could
25 compromise availability in case of production issues (see eg.
26 [https://www.ferringusa.com/press/ferring-us-issues-voluntary-nationwide-recall-of-ddavp-
27 nasal-spray-10-mcg-0-1ml-desmopressin-acetate-nasal-spray-10-mcg-0-1ml-stimate-nasal-
28 spray-1-5-mg-ml-due-to-superpotency/](https://www.ferringusa.com/press/ferring-us-issues-voluntary-nationwide-recall-of-ddavp-nasal-spray-10-mcg-0-1ml-desmopressin-acetate-nasal-spray-10-mcg-0-1ml-stimate-nasal-spray-1-5-mg-ml-due-to-superpotency/)).

29 Hormonal treatment is commonly used to reduce heavy menstrual bleeding in VWD. However,
30 as for desmopressin, increased VWF/FVIII levels are a mere secondary effect of hormonal
31 treatment, which in itself has its own side effects and contra-indications. These are not always

1 minor, as for example the use of progesterone has been associated with an increased risk of
2 sphenoidal osteomeningiomas.

3 In conclusion, YES there are some efficient treatments for VWD, but NO this does not mean
4 that we should stop trying to improve these treatments with the goals of reaching superior
5 efficacy, improving quality of life of the patients and finally curing the disease.

6

7 **The challenge of innovation in VWD management**

8 What can be done to improve VWD treatment? Unsurprisingly, there is not a simple answer
9 to that question. Increasing prophylactic treatment would be a first step, as the introduction
10 of prophylaxis for VWD patients appears to significantly reduce the rate of bleeding episodes
11 and the need for hospitalization.^{53,54} Nevertheless, prophylactic treatment did not fully
12 prevent against the occurrence of bleeding episodes.⁵⁴ In addition, the current concentrates
13 have similar issues as the FVIII concentrates that spurred the development of novel treatment
14 strategies for hemophilia, including a relatively short half-life requiring multiple intravenous
15 infusions *per week*. Designing VWF variants with a prolonged half-life could be an option to
16 reduce infusion frequency. Such a strategy would not be such a major challenge to implement
17 but besides an abstract published in 2006 which showed its validity, no follow-up
18 development has been put in place.⁵⁵

19 However, to be truly innovative, we as a scientific community need to think of alternative
20 ways that are adapted to the patients' situation, rather than being limited to VWF
21 concentrates or desmopressin. It is with such novel mindset that we will be able to break the
22 glass ceiling and come up with the VWD-specific equivalents of fitusiran or emicizumab. In
23 order to move in this direction, we first need to identify the patients who will better benefit
24 from a more personalized approach, a concept increasingly recognized as relevant.^{37,56} As part
25 of this identification process, we should realize that severe VWD is not necessarily restricted
26 to VWD-type 3. Patients with VWD-type 1 or 2 may sometimes experience bleeding
27 phenotypes similar in severity than some VWD-type 3 patients.⁵⁷ Knowing the VWD subtype
28 of the patients will be crucial to adapt treatment but should not represent a limit in our
29 perception and appreciation of disease severity. There are patients who experience major
30 bleeds and some who have frequent minor bleeds and our challenge is to address the different
31 needs of the bleeders.

1 In the following sections, we will describe a number of potential approaches that could change
2 the current paradigm of how VWD is treated or can be treated in the upcoming years. Of note,
3 we will not address specifically the platelet dysfunction described in some patients with VWD-
4 type 2B⁵⁸ since targeting the VWF gene defect would also resolve the platelet phenotype.

5

6 **Novel strategies: genetic approaches**

7 *Gene therapy*

8 To really cure VWD, gene therapy represents the ultimate goal, especially as it seems to be
9 successful for both hemophilia A and B. However, the challenges to express VWF via gene
10 therapy appear quite daunting. A first obstacle relates to its multimerization. Several mouse
11 studies have shown that functional VWF can be efficiently expressed in hepatocytes of VWF-
12 deficient mice, either transiently⁵⁹⁻⁶² or for long-term expression.^{63,64} However, hepatocytes
13 are not well-equipped to produce sufficiently multimerized VWF, and over time the
14 hemostatically active VWF multimers seem to disappear.⁶³ So targeting the endothelium will
15 be necessary for efficient protein processing.

16 Size represents a second hurdle: the VWF cDNA is too large to be incorporated in adeno-
17 associated viruses (AAV), the type of vectors most used in clinical trials. Successful
18 incorporation into lentiviruses has been reported, but data on efficient *in vivo* expression is
19 yet lacking.^{65,66}

20 In an attempt to overcome both obstacles, our team in collaboration with Federico Mingozzi's
21 team has designed a dual-AAV vector approach, in which the cDNA of VWF is divided over two
22 AAV-particles, which in turn are adapted to specifically target endothelial cells. We show that
23 such an approach is technically feasible although the current expression levels are too low to
24 be clinically meaningful.⁶⁷

25 But even if we overcome these obstacles, there is another hurdle that we cannot ignore: if
26 VWF is finally expressed at reasonable levels in the endothelial cells of the patient, how can
27 we avoid the incorporation of dominant-negative mutants into the VWF multimer, when being
28 expressed in the same cells as the transgene VWF? Indeed, Bowman *et al.* reported that the
29 inheritance pattern for VWD-type 3 proved co-dominant in approximately 50% of the families
30 included in their study.⁶⁸

31 Perhaps expression of full-length VWF is not always necessary. In VWD-type 2N, and also some
32 VWD-type 3 patients, the bleeding complications are associated with the low levels of FVIII,

1 and increasing FVIII levels would help to reduce bleeding in joints and soft tissues. For those
2 patients, a gene therapy approach using the N-terminal portion of VWF (D1-D2-D'-D3 domains)
3 would result in normalization of FVIII levels, as the presence of the dimeric D'D3-fragment
4 protects FVIII from degradation and premature clearance.⁶⁹ Advantage of this approach is that
5 multimerization is not a requirement, the cDNA (about 3.8 kb) easily fits into the AAV-vector
6 and the protein can be expressed in hepatocytes. Preliminary studies in mice showed that
7 such an approach is viable from a technical point of view.⁶⁹

8

9 *Transcriptional silencing/gene-editing*

10 Rather than trying to express functional VWF, another approach would be to silence
11 expression of mutant VWF. This makes even more sense given the dominant-negative
12 character of many of the VWF mutations. A first example of this approach was reported by
13 Casari *et al.*, who demonstrated that siRNA-mediated targeting of the mRNA breakpoint of an
14 in-frame deletion mutant improved protein expression and multimerization.⁷⁰ Recently, we
15 applied this siRNA-approach to a mouse model of VWD and confirmed the restoration of VWF
16 multimers in plasma by silencing the dominant-negative allele.⁷¹ To find a broader application
17 for this technology, de Jong *et al.* designed siRNAs targeting common single-nucleotide
18 polymorphisms in the *VWF* mutant allele, while leaving expression of the wild-type allele
19 unaffected.⁷²

20 Alternatively, another option could be to correct the *VWF* gene itself, using gene-editing
21 approaches such as Crispr-Cas9. Little information on this approach in view of VWD currently
22 exists, with the exception of a study investigating the bi-allelic deletion of the *VWF* gene in
23 endothelial colony forming cells (ECFCs, also known as blood outgrowth endothelial cells).⁷³

24 Both siRNA and gene-editing approaches are attractive, but one of the technical difficulties
25 includes the specific targeting of sufficient numbers of endothelial cells. Currently, we do not
26 know how many endothelial cells would need to be targeted to achieve sufficient production
27 of functional VWF multimers. Additional research on these promising strategies is clearly
28 needed at this point.

29

30 *Cellular therapy*

31 *Ex vivo* cellular therapy represents another method to introduce cells expressing functional
32 VWF. This method was previously described by De Meyer and colleagues.⁶⁵ Briefly, canine

1 ECFCs were transduced using lentiviral vectors containing the human VWF cDNA. One day
2 after transduction, cells indeed expressed functional multimerized VWF at levels 3-fold higher
3 compared to normal ECFCs. However, expression levels to decreased 7-fold over a 4-week
4 period during the expansion phase, which could be due to *in vitro* culturing of these cells.
5 While a similar approach could also be applied using endothelial cells derived from induced
6 pluripotent stem cells, human somatic cell therapy approaches are currently limited by the
7 ability to generate sufficient amounts of cells to achieve physiological relevant expression
8 levels after transplantation, the uncertain life-span of transplanted cells and the occurrence
9 of genetic changes in the cells.⁷⁴ Progress on cellular therapies in other disease will certainly
10 be helpful for the future development of this approach regarding VWD.

11

12 **Novel strategies: protein-based strategies**

13 *Compensating FVIII-deficiency in VWD*

14 Some of the bleeding complications in VWD-type 2N and type 3 patients are related to low
15 FVIII levels, and improving FVIII levels could thus be beneficial. Infusing standard recombinant
16 FVIII is not a viable option, since FVIII will be extremely-short lived in the absence of protection
17 conferred by VWF. On the other hand, using FVIII/VWF concentrates in type 2N patients could
18 lead to an excess of VWF, potentially posing a thrombotic risk.^{75,76} An alternative solution may
19 soon become available with the development of BIVV001, a novel FVIII-fusion variant which
20 circulates independently of endogenous VWF, having a half-life 3-4 times longer than
21 recombinant FVIII.^{77,78} Although being developed for hemophilia A, this molecule could
22 provide a novel treatment option for VWD-type 2N patients to correct FVIII-deficiency without
23 increasing VWF levels.

24 Another approach was described by Weyand and colleagues, related to a VWD-type 3 patient
25 who had developed allo-antibodies against both FVIII and VWF and experienced recurrent
26 bleeding episodes.⁷⁹ To compensate for FVIII-deficiency in this patient, treatment with
27 emicizumab was initiated. Emicizumab is a bispecific antibody that mimics to some extent the
28 cofactor function of FVIII.^{43,80} Emicizumab prophylaxis resulted in improved bleed prevention
29 and reduced treatment burden.⁷⁹

30

31 *Increasing endogenous VWF levels*

1 Desmopressin is often the first choice of treatment in VWD-type 1, as it will temporarily
2 normalize endogenous levels of the otherwise functional VWF molecule. Due to the
3 limitations associated with desmopressin detailed earlier, it would be of interest to develop
4 novel strategies to increase endogenous VWF levels in a more sustainable approach. Various
5 options could be designed in this respect. For instance, after the discovery that interleukin-11
6 infusion was associated with an increase in VWF levels in both mouse and canine models, its
7 clinical benefit was evaluated in two Phase II studies.⁸¹⁻⁸⁴ Although some clinical benefit
8 following subcutaneous application was observed, the increase in VWF levels was modest
9 (1.1-1.5 fold).^{83,84} No additional clinical studies seem to be registered at Clinicaltrials.gov.
10 Another strategy includes the use of single-domain antibodies (nanobodies) fused to an
11 albumin-binding moiety (*eg.* peptide or nanobody), for instance, a VWF-binding nanobody
12 that is fused to an albumin-binding peptide.⁸⁵ In this approach, such fusion protein would
13 bridge circulating VWF to albumin, taking advantage of the long albumin half-life, and induce
14 a prolonged half-life for VWF and consequently, increase VWF levels. Preliminary mouse
15 studies revealed that pre-incubating VWF with a VWF-binding nanobody fused to an albumin-
16 binding peptide (KB-VWF-013-ABP) resulted in a 7-fold increased VWF half-life. Moreover,
17 subcutaneous application of this fusion protein was associated with a 4-8 fold increase in
18 VWF- and FVIII-levels over a period of 7 days.⁸⁵ This approach thus holds great promises but
19 needs to be further confirmed in different preclinical models.

20

21 *Preventing VWF degradation*

22 Excessive degradation of VWF is a particular problem in congenital VWD-type 2A(IIA), but also
23 in some variants of acquired von Willebrand syndrome (aVWs), for instance in those having
24 severe aortic stenosis or patients requiring mechanical circulatory support. While congenital
25 VWD-type 2A could be treated using VWF concentrates, this is trickier in aVWs since it seems
26 conceivable that the infused VWF will also be susceptible to degradation while passing
27 through the pumps. To reduce VWF degradation, antibodies can be used to interfere with
28 ADAMTS13-VWF interactions. Importantly, inhibition of VWF degradation should be limited
29 to avoid thrombotic side effects as seen in thrombotic thrombocytopenic purpura. A few years
30 ago, we described an anti-VWF antibody (Mab508) targeting the ADAMTS13-docking site at
31 the VWF-D4 domain.⁸⁶ This antibody interferes with excessive VWF degradation, while leaving
32 basal degradation intact. Indeed, the presence of Mab508 led to the conservation of high

1 molecular weight-multimers in *ex vivo* whole-blood perfusions systems.⁸⁶ Given the
2 advantages of antibody-based treatment over concentrate-based treatments (*eg.* long half-
3 life, eventual subcutaneous administration), such antibodies could be valuable assets in
4 managing patients characterized by excessive VWF degradation.

5

6 **Forward-looking perspective**

7 The heterogeneous nature of VWD implies that a “one size fits all” approach will never have
8 the potential to address all the patients’ needs. However, the management of VWD patients
9 currently follows such a trend. As highlighted in the last few paragraphs, research has not
10 stood still and various solutions have been or are under investigation, although more could
11 definitely be done in this regard. The main roadblock, in our view, appears at the next level,
12 the translation of research results in potential industrial partnerships, allowing development
13 of future treatments. So what are the biggest obstacles? Market size is certainly a major hurdle
14 with few patients likely to benefit from each of the approaches highlighted previously.
15 Another difficulty comes from the difference in perception of disease burden between health
16 care professionals and the patients themselves. Maybe it is time to stop comparing bleeding
17 in hemophilia and bleeding in VWD. All HR-QoL studies tell us that VWD patients have a
18 significantly impaired quality of life. Patient’s advocacy has therefore a critical role to play in
19 order to bring all stakeholders around the same table in order to prepare the future. No
20 patient should be left behind. Our main aspiration is that ten years from now, if we were to
21 write a review about VWD management, such a review will be quite different than the ones
22 published for the last 30 years.

23

24 **Acknowledgements:**

25 Financial support by the Agence Nationale de la Recherche (ANR-17-RHUS-0011) is gratefully
26 acknowledged. This perspective could not have been written without the input of Dr. Olivier
27 Christophe and Dr. Caterina Casari.

28

29 **Author contributions:**

30 All authors contribution to the conception, writing and editing of this manuscript.

31

32 **Disclosures:**

1 CVD, SS and PJJ are inventors on patents owned by Inserm related to bleeding disorders. CVD,
2 SS and PJJ are co-founders of Laelaps Therapeutics.

3

4 **Legends**

5 **Figure 1: Evolution of treatment modalities for hemophilia A and VWD**

6 A schematic overview of when novel treatment options became available for hemophilia A
7 and VWD. High-purity and recombinant concentrates (orange) arrived 20 years later for VWD
8 compared to hemophilia A. Extended half-life variants, non-factor therapies and gene therapy
9 (red) have found their way to the clinic or are in late-stage clinical development for hemophilia
10 A, whereas they are not yet at the horizon with regard to VWD.

11

12

13 **Figure 2: Potential avenues to enrich the arsenal of treatment options for VWD**

14 Examples of potential novel genetic- or protein-based strategies for the management of VWD
15 are highlighted. Some approaches could be used to serve all VWD patients, whereas others
16 will be type-specific. Abbreviations used in the figure: AAV: adeno-associated virus; ECs:
17 endothelial cells; ECFCs: endothelial colony forming cells (also referred to as blood outgrowth
18 endothelial cells); ECMO: extra-corporal membrane oxygenation; FVIII: factor VIII; IL-11:
19 interleukin-11; LVAD: left ventricular assist device; TALENs: transcription activator-like
20 effector nucleases; VWD: von Willebrand disease; VWs: von Willebrand syndrome. Portions
21 of this figure were created using Biorender.com.

22

23

1 **References**

2

3 1. Leebeek FW, Eikenboom JC. Von Willebrand's Disease. *N Engl J Med.*
4 2016;375(21):2067-2080.

5 2. Nichols WL, Hultin MB, James AH, et al. von Willebrand disease (VWD): evidence-based
6 diagnosis and management guidelines, the National Heart, Lung, and Blood Institute
7 (NHLBI) Expert Panel report (USA). *Haemophilia.* 2008;14(2):171-232.

8 3. Sadler JE, Mannucci PM, Berntorp E, et al. Impact, diagnosis and treatment of von
9 Willebrand disease. *Thromb Haemost.* 2000;84(2):160-174.

10 4. Lak M, Peyvandi F, Mannucci PM. Clinical manifestations and complications of
11 childbirth and replacement therapy in 385 Iranian patients with type 3 von Willebrand
12 disease. *Br J Haematol.* 2000;111(4):1236-1239.

13 5. Silwer J. von Willebrand's disease in Sweden. *Acta Paediatr Scand Suppl.* 1973;238:1-
14 159.

15 6. James PD, Connell NT, Ameer B, et al. ASH ISTH NHF WFH 2021 guidelines on the
16 diagnosis of von Willebrand disease. *Blood Adv.* 2021;published online 12 January
17 2021. DOI 10.1182/bloodadvances.2020003265.

18 7. Weyand AC, James PD. Sexism in the management of bleeding disorders. *Res Pract*
19 *Thromb Haemost.* 2020;5(1):51-54.

20 8. Atiq F, Saes JL, Punt MC, et al. Major differences in clinical presentation, diagnosis and
21 management of men and women with autosomal inherited bleeding disorders.
22 *EClinicalMedicine.* 2021; doi.org/10.1016/j.eclinm.2021.100726.

23 9. Kouides PA. Females with von Willebrand disease: 72 years as the silent majority.
24 *Haemophilia.* 1998;4(4):665-676.

25 10. Greenwood BN, Carnahan S, Huang L. Patient-physician gender concordance and
26 increased mortality among female heart attack patients. *Proc Natl Acad Sci U S A.*
27 2018;115(34):8569-8574.

28 11. Regitz-Zagrosek V. Sex and gender differences in health. Science & Society Series on
29 Sex and Science. *EMBO Rep.* 2012;13(7):596-603.

30 12. Roter DL, Hall JA, Aoki Y. Physician gender effects in medical communication: a meta-
31 analytic review. *JAMA.* 2002;288(6):756-764.

- 1 13. Castaman G, James PD. Pregnancy and delivery in women with von Willebrand disease.
2 *Eur J Haematol.* 2019;103(2):73-79.
- 3 14. Laffan M, Sathar J, Johnsen JM. von Willebrand disease: Diagnosis and treatment,
4 treatment of women, and genomic approach to diagnosis. *Haemophilia.* 2020;doi:
5 10.1111/hae.14050.
- 6 15. Lavin M, Aguila S, Dalton N, et al. Significant gynecological bleeding in women with low
7 von Willebrand factor levels. *Blood Adv.* 2018;2(14):1784-1791.
- 8 16. Snyder CF, Jensen RE, Segal JB, Wu AW. Patient-reported outcomes (PROs): putting
9 the patient perspective in patient-centered outcomes research. *Med Care.* 2013;51(8
10 Suppl 3):S73-79.
- 11 17. de Wee EM, Mauser-Bunschoten EP, Van Der Bom JG, et al. Health-related quality of
12 life among adult patients with moderate and severe von Willebrand disease. *J Thromb*
13 *Haemost.* 2010;8(7):1492-1499.
- 14 18. Barr RD, Sek J, Horsman J, et al. Health status and health-related quality of life
15 associated with von Willebrand disease. *Am J Hematol.* 2003;73(2):108-114.
- 16 19. Govorov I, Ekelund L, Chaireti R, et al. Heavy menstrual bleeding and health-associated
17 quality of life in women with von Willebrand's disease. *Exp Ther Med.*
18 2016;11(5):1923-1929.
- 19 20. Kadir RA, Sabin CA, Pollard D, Lee CA, Economides DL. Quality of life during
20 menstruation in patients with inherited bleeding disorders. *Haemophilia.*
21 1998;4(6):836-841.
- 22 21. Solovieva S. Clinical severity of disease, functional disability and health-related quality
23 of life. Three-year follow-up study of 150 Finnish patients with coagulation disorders.
24 *Haemophilia.* 2001;7(1):53-63.
- 25 22. Xu Y, Deforest M, Grabell J, Hopman W, James P. Relative contributions of bleeding
26 scores and iron status on health-related quality of life in von Willebrand disease: a
27 cross-sectional study. *Haemophilia.* 2017;23(1):115-121.
- 28 23. Von Mackensen S. Quality of life in women with bleeding disorders. *Haemophilia.*
29 2011;17 Suppl 1:33-37.
- 30 24. Kujovich JL. von Willebrand's disease and menorrhagia: prevalence, diagnosis, and
31 management. *Am J Hematol.* 2005;79(3):220-228.

- 1 25. Kouides PA, Phatak PD, Burkart P, et al. Gynaecological and obstetrical morbidity in
2 women with type I von Willebrand disease: results of a patient survey. *Haemophilia*.
3 2000;6(6):643-648.
- 4 26. Kirtava A, Drews C, Lally C, Dilley A, Evatt B. Medical, reproductive and psychosocial
5 experiences of women diagnosed with von Willebrand's disease receiving care in
6 haemophilia treatment centres: a case-control study. *Haemophilia*. 2003;9(3):292-297.
- 7 27. Marshall AL, Dasari H, Warner ND, Grill DE, Nichols WL, Pruthi RK. Self-reported
8 reproductive health experiences in women with von Willebrand disease: a qualitative
9 interview-based study. *J Obstet Gynaecol*. 2019;39(2):288-290.
- 10 28. Arya S, Wilton P, Page D, et al. « Everything Was Blood When it Comes to Me”:
11 Understanding the Lived Experiences of Women with Inherited Bleeding Disorders. *J*
12 *Thromb Haemost*. 2020;18(12):3211-3221.
- 13 29. van Galen KPM, Meijer K, Vogely HC, et al. Joint surgery in von Willebrand disease: a
14 multicentre cross-sectional study. *Haemophilia*. 2016;22(2):256-262.
- 15 30. Von Mackensen S, Federici A. Quality of life assessment in patients with von
16 Willebrand’s disease: development of a first disease-specific instrument (VWD-QOL).
17 *Poster presentation at the annual congress of the Society of Thrombosis and*
18 *Haemostasis Research, Dresden, 21-25 February 2007, P126*. 2007.
- 19 31. Aledort LM. Treatment of von Willebrand's disease. *Mayo Clin Proc*. 1991;66(8):841-
20 846.
- 21 32. Kruse-Jarres R, Johnsen JM. How I treat type 2B von Willebrand disease. *Blood*.
22 2018;131(12):1292-1300.
- 23 33. Lavin M, O'Donnell JS. New treatment approaches to von Willebrand disease.
24 *Hematology Am Soc Hematol Educ Program*. 2016;2016(1):683-689.
- 25 34. Lavin M, O'Donnell JS. How I treat low von Willebrand factor levels. *Blood*.
26 2019;133(8):795-804.
- 27 35. Leebeek FWG, Atiq F. How I manage severe von Willebrand disease. *Br J Haematol*.
28 2019;187(4):418-430.
- 29 36. Mannucci PM. How I treat patients with von Willebrand disease. *Blood*.
30 2001;97(7):1915-1919.
- 31 37. Phua CW, Berntorp E. A personalized approach to the management of VWD. *Transfus*
32 *Apher Sci*. 2019;58(5):590-595.

- 1 38. Tosetto A, Castaman G. How I treat type 2 variant forms of von Willebrand disease.
2 *Blood*. 2015;125(6):907-914.
- 3 39. Connell NT, Flood V, Brignardello-Petersen R, et al. ASH ISTH NHF WFH 2021 guidelines
4 on the management of von Willebrand disease. *Blood Adv*. 2021;published online 12
5 January 2021. DOI 10.1182/bloodadvances.2020003264.
- 6 40. Goudemand J, Scharrer I, Berntorp E, et al. Pharmacokinetic studies on Wilfactin, a von
7 Willebrand factor concentrate with a low factor VIII content treated with three virus-
8 inactivation/removal methods. *J Thromb Haemost*. 2005;3(10):2219-2227.
- 9 41. Mannucci PM, Kempton C, Millar C, et al. Pharmacokinetics and safety of a novel
10 recombinant human von Willebrand factor manufactured with a plasma-free method:
11 a prospective clinical trial. *Blood*. 2013;122(5):648-657.
- 12 42. Chowdary P. Anti-tissue factor pathway inhibitor (TFPI) therapy: a novel approach to
13 the treatment of haemophilia. *Int J Hematol*. 2020;111(1):42-50.
- 14 43. Kitazawa T, Igawa T, Sampei Z, et al. A bispecific antibody to factors IXa and X restores
15 factor VIII hemostatic activity in a hemophilia A model. *Nat Med*. 2012;18(10):1570-
16 1574.
- 17 44. Polderdijk SG, Adams TE, Ivanciu L, Camire RM, Baglin TP, Huntington JA. Design and
18 characterization of an APC-specific serpin for the treatment of hemophilia. *Blood*.
19 2017;129(1):105-113.
- 20 45. Sehgal A, Barros S, Ivanciu L, et al. An RNAi therapeutic targeting antithrombin to
21 rebalance the coagulation system and promote hemostasis in hemophilia. *Nat Med*.
22 2015;21(5):492-497.
- 23 46. Miesbach W, Berntorp E. Translating the success of prophylaxis in haemophilia to von
24 Willebrand disease. *Thromb Res*. 2021;199:67-74.
- 25 47. Federici AB. The use of desmopressin in von Willebrand disease: the experience of the
26 first 30 years (1977-2007). *Haemophilia*. 2008;14 Suppl 1:5-14.
- 27 48. Mannucci PM. New therapies for von Willebrand disease. *Blood Adv*. 2019;3(21):3481-
28 3487.
- 29 49. Miesbach W, Krekeler S, Duck O, et al. Clinical assessment of efficacy and safety of
30 DDAVP. *Hamostaseologie*. 2010;30 Suppl 1:S172-175.
- 31 50. Stoof SC, Cnossen MH, de Maat MP, Leebeek FW, Kruijff MJ. Side effects of
32 desmopressin in patients with bleeding disorders. *Haemophilia*. 2016;22(1):39-45.

- 1 51. Kalot MA, Al-Khatib M, Connell NT, et al. An international survey to inform priorities
2 for new guidelines on von Willebrand disease. *Haemophilia*. 2020;26(1):106-116.
- 3 52. Mannucci PM, Bettega D, Cattaneo M. Patterns of development of tachyphylaxis in
4 patients with haemophilia and von Willebrand disease after repeated doses of
5 desmopressin (DDAVP). *Br J Haematol*. 1992;82(1):87-93.
- 6 53. Holm E, Carlsson KS, Lovdahl S, Lail AE, Abshire TC, Berntorp E. Bleeding-related
7 hospitalization in patients with von Willebrand disease and the impact of prophylaxis:
8 Results from national registers in Sweden compared with normal controls and
9 participants in the von Willebrand Disease Prophylaxis Network. *Haemophilia*.
10 2018;24(4):628-633.
- 11 54. Peyvandi F, Castaman G, Gresele P, et al. A phase III study comparing secondary long-
12 term prophylaxis versus on-demand treatment with vWF/FVIII concentrates in severe
13 inherited von Willebrand disease. *Blood Transfus*. 2019;17(5):391-398.
- 14 55. Turecek PL, Scheiflinger F, Siekmann J, et al. Biochemical and Functional
15 Characterization of PEGylated rVWF. *Blood*. 2006;108 (11): 1021.
- 16 56. Hazendonk H, Heijdra JM, de Jager NCB, et al. Analysis of current perioperative
17 management with Haemate((R)) P/Humate P((R)) in von Willebrand disease:
18 Identifying the need for personalized treatment. *Haemophilia*. 2018;24(3):460-470.
- 19 57. Federici AB, Bucciarelli P, Castaman G, et al. The bleeding score predicts clinical
20 outcomes and replacement therapy in adults with von Willebrand disease. *Blood*.
21 2014;123(26):4037-4044.
- 22 58. Casari C, Berrou E, Leuret M, et al. von Willebrand factor mutation promotes
23 thrombocytopenia by inhibiting integrin alphaIIb beta3. *J Clin Invest*.
24 2013;123(12):5071-5081.
- 25 59. De Meyer SF, Vandeputte N, Pareyn I, et al. Restoration of plasma von Willebrand
26 factor deficiency is sufficient to correct thrombus formation after gene therapy for
27 severe von Willebrand disease. *Arterioscler Thromb Vasc Biol*. 2008;28(9):1621-1626.
- 28 60. Golder M, Pruss CM, Hegadorn C, et al. Mutation-specific hemostatic variability in mice
29 expressing common type 2B von Willebrand disease substitutions. *Blood*.
30 2010;115(23):4862-4869.

- 1 61. Marx I, Christophe OD, Lenting PJ, et al. Altered thrombus formation in von Willebrand
2 factor-deficient mice expressing von Willebrand factor variants with defective binding
3 to collagen or GPIIb/IIIa. *Blood*. 2008;112(3):603-609.
- 4 62. Rayes J, Hollestelle MJ, Legendre P, et al. Mutation and ADAMTS13-dependent
5 modulation of disease severity in a mouse model for von Willebrand disease type 2B.
6 *Blood*. 2010;115(23):4870-4877.
- 7 63. Portier I, Vanhoorelbeke K, Verhenne S, et al. High and long-term von Willebrand
8 factor expression after Sleeping Beauty transposon-mediated gene therapy in a mouse
9 model of severe von Willebrand disease. *J Thromb Haemost*. 2018;16(3):592-604.
- 10 64. Quiviger M, Giannakopoulos A, Verhenne S, et al. Improved molecular platform for the
11 gene therapy of rare diseases by liver protein secretion. *Eur J Med Genet*.
12 2018;61(11):723-728.
- 13 65. De Meyer SF, Vanhoorelbeke K, Chuah MK, et al. Phenotypic correction of von
14 Willebrand disease type 3 blood-derived endothelial cells with lentiviral vectors
15 expressing von Willebrand factor. *Blood*. 2006;107(12):4728-4736.
- 16 66. Wang L, Rosenberg JB, De BP, et al. In vivo gene transfer strategies to achieve partial
17 correction of von Willebrand disease. *Hum Gene Ther*. 2012;23(6):576-588.
- 18 67. Barbon E, Kawecki C, Marmier S, et al. Development of a dual hybrid AAV vector for
19 endothelial-targeted expression of von Willebrand factor. *Gene Ther*. 2021.
- 20 68. Bowman M, Tuttle A, Notley C, et al. The genetics of Canadian type 3 von Willebrand
21 disease: further evidence for co-dominant inheritance of mutant alleles. *J Thromb*
22 *Haemost*. 2013;11(3):512-520.
- 23 69. Yee A, Gildersleeve RD, Gu S, et al. A von Willebrand factor fragment containing the
24 D'D3 domains is sufficient to stabilize coagulation factor VIII in mice. *Blood*.
25 2014;124(3):445-452.
- 26 70. Casari C, Pinotti M, Lancellotti S, et al. The dominant-negative von Willebrand factor
27 gene deletion p.P1127_C1948delinsR: molecular mechanism and modulation. *Blood*.
28 2010;116(24):5371-5376.
- 29 71. Campioni M, Legendre P, Loubiere C, et al. In vivo modulation of a dominant-negative
30 variant in mouse models of von Willebrand disease type 2A. *J Thromb Haemost*.
31 2021;19(1):139-146.

- 1 72. de Jong A, Dirven RJ, Boender J, et al. Ex vivo Improvement of a von Willebrand Disease
2 Type 2A Phenotype Using an Allele-Specific Small-Interfering RNA. *Thromb Haemost.*
3 2020;120(11):1569-1579.
- 4 73. Schillemans M, Kat M, Westeneng J, et al. Alternative trafficking of Weibel-Palade body
5 proteins in CRISPR/Cas9-engineered von Willebrand factor-deficient blood outgrowth
6 endothelial cells. *Res Pract Thromb Haemost.* 2019;3(4):718-732.
- 7 74. Halliwell J, Barbaric I, Andrews PW. Acquired genetic changes in human pluripotent
8 stem cells: origins and consequences. *Nat Rev Mol Cell Biol.* 2020;21(12):715-728.
- 9 75. Franchini M. Thrombotic complications in von Willebrand disease. *Hematology.*
10 2006;11(1):49-52.
- 11 76. Rietveld IM, Lijfering WM, le Cessie S, et al. High levels of coagulation factors and
12 venous thrombosis risk: strongest association for factor VIII and von Willebrand factor.
13 *J Thromb Haemost.* 2019;17(1):99-109.
- 14 77. Konkle BA, Shapiro AD, Quon DV, et al. BIVV001 Fusion Protein as Factor VIII
15 Replacement Therapy for Hemophilia A. *N Engl J Med.* 2020;383(11):1018-1027.
- 16 78. Seth Chhabra E, Liu T, Kulman J, et al. BIVV001, a new class of factor VIII replacement
17 for hemophilia A that is independent of von Willebrand factor in primates and mice.
18 *Blood.* 2020;135(17):1484-1496.
- 19 79. Weyand AC, Flood VH, Shavit JA, Pipe SW. Efficacy of emicizumab in a pediatric patient
20 with type 3 von Willebrand disease and alloantibodies. *Blood Adv.* 2019;3(18):2748-
21 2750.
- 22 80. Lenting PJ, Denis CV, Christophe OD. Emicizumab, a bispecific antibody recognizing
23 coagulation factors IX and X: how does it actually compare to factor VIII? *Blood.*
24 2017;130(23):2463-2468.
- 25 81. Denis CV, Kwack K, Saffaripour S, et al. Interleukin 11 significantly increases plasma
26 von Willebrand factor and factor VIII in wild type and von Willebrand disease mouse
27 models. *Blood.* 2001;97(2):465-472.
- 28 82. Olsen EH, McCain AS, Merricks EP, et al. Comparative response of plasma VWF in dogs
29 to up-regulation of VWF mRNA by interleukin-11 versus Weibel-Palade body release
30 by desmopressin (DDAVP). *Blood.* 2003;102(2):436-441.
- 31 83. Ragni MV, Jankowitz RC, Jaworski K, Merricks EP, Kloos MT, Nichols TC. Phase II
32 prospective open-label trial of recombinant interleukin-11 in women with mild von

- 1 Willebrand disease and refractory menorrhagia. *Thromb Haemost.* 2011;106(4):641-
2 645.
- 3 84. Ragni MV, Novelli EM, Murshed A, Merricks EP, Kloos MT, Nichols TC. Phase II
4 prospective open-label trial of recombinant interleukin-11 in desmopressin-
5 unresponsive von Willebrand disease and mild or moderate haemophilia A. *Thromb*
6 *Haemost.* 2013;109(2):248-254.
- 7 85. Kawecki C, Aymé G, Casari C, et al. Development of nanobodies fused to albumin
8 binding peptide: a tool to increase plasmatic levels of endogenous proteins. *Res Pract*
9 *Thromb Haemost.* 2017;1 (Suppl 1):OC-38.4.
- 10 86. Rauch A, Legendre P, Christophe OD, et al. Antibody-based prevention of von
11 Willebrand factor degradation mediated by circulatory assist devices. *Thromb*
12 *Haemost.* 2014;112(5):1014-1023.

Figure 1: Evolution of treatment modalities for hemophilia A and von Willebrand disease

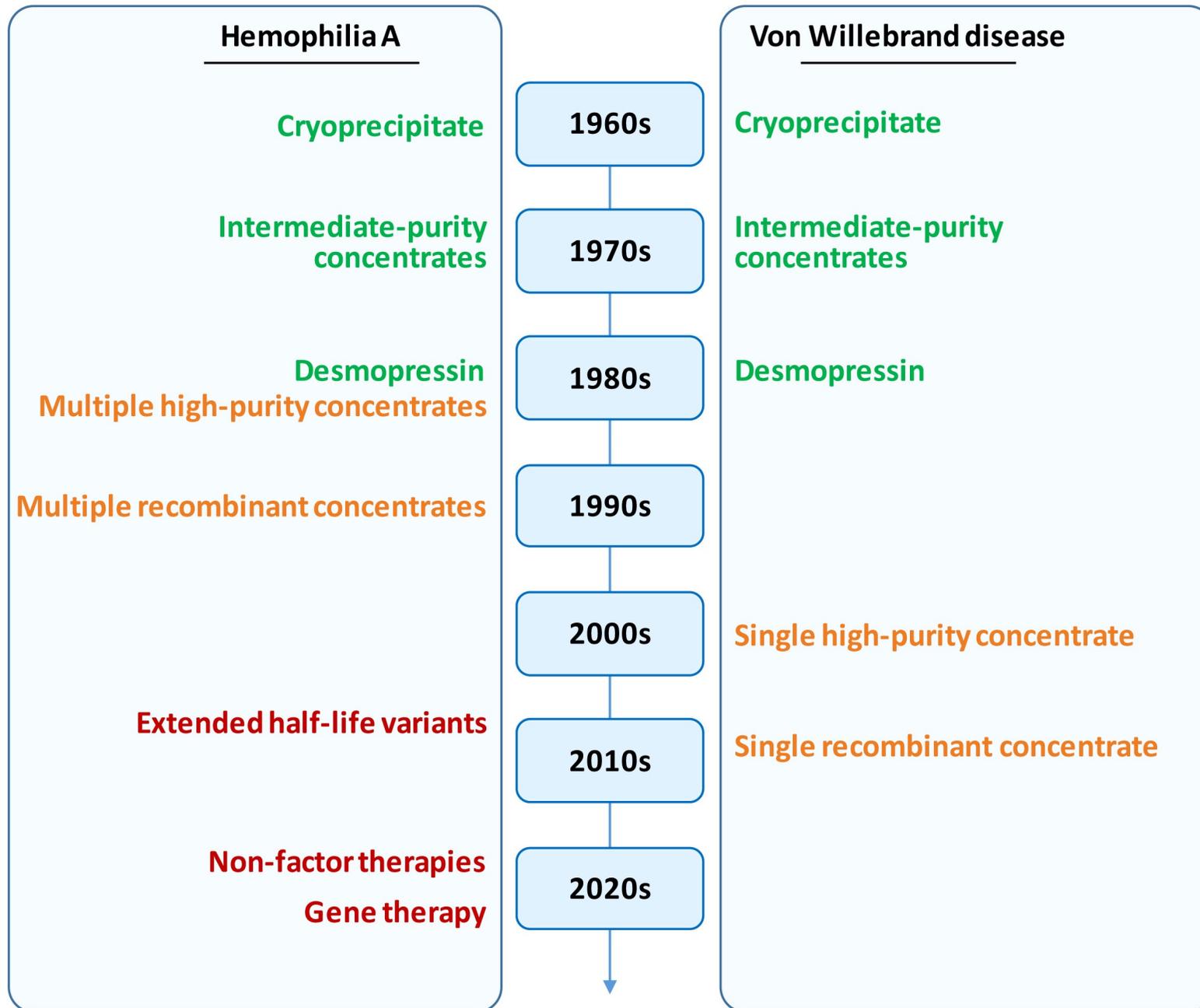
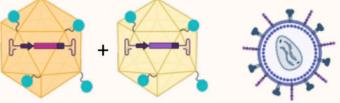
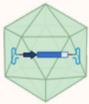
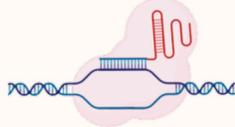
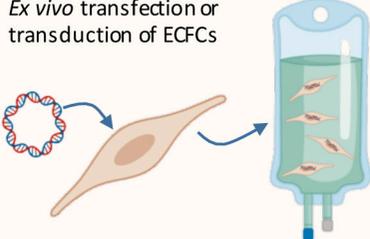
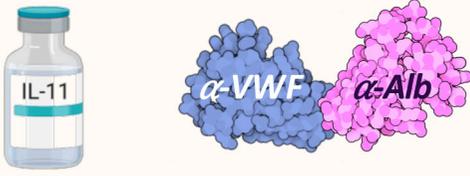
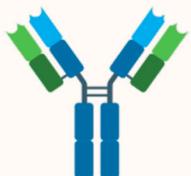


Figure 2: Potential avenues to enrich the arsenal of treatment options for VWD

Genetic approaches

| | |
|---|--|
| <p>Somatic gene therapy</p>  <p>EC-specific dual-vector AAV + Lentivirus targeting ECs</p> <ul style="list-style-type: none"> • Expression of full-length VWF in ECs to ensure optimal multimerization • All VWD-types, but absence of dominant-negative mutants in host cells is obligatory <hr/>  <p>Hepatocyte-specific AAV</p> <ul style="list-style-type: none"> • Expression of VWF D1-D2-D'-D3 fragment in hepatocytes to normalize FVIII levels • VWD-type 2N & type 3 | <p>Gene editing</p>  <p>Crispr-Cas9 TALENs Zinc-finger nucleases</p> <ul style="list-style-type: none"> • Mono-allelic gene editing for dominant-negative mutants • Bi-allelic gene editing for recessive mutants • All types of VWD |
| <p>Transcriptional silencing</p>  <p>siRNA</p> <ul style="list-style-type: none"> • Silencing of mutant alleles • All types of VWD | <p>Cellular therapy</p>  <p>Ex vivo transfection or transduction of ECFCs</p> <ul style="list-style-type: none"> • Expression of full-length VWF in ECs to ensure optimal multimerization • No co-multimerization with mutant VWF, but co-existence in circulation • All types of VWD |

Protein-based therapies

| | | |
|--|---|---|
| <p>Compensating FVIII deficiency</p>  <p>BIVV001 emicizumab</p> <ul style="list-style-type: none"> • BIVV001 is a FVIII-variant that circulates independently of endogenous VWF • Emicizumab is a bispecific antibody mimicking FVIII-cofactor function & can be given subcutaneously • VWD-type 2N and type 3 | <p>Increasing endogenous VWF levels</p>  <p>Interleukin-11 bispecific nanobodies</p> <ul style="list-style-type: none"> • IL-11 stimulates synthesis of VWF, resulting in a modest rise of VWF levels (1.1-1.5-fold) • can be given subcutaneously • Bispecific nanobodies may delay clearance of endogenous VWF & can be given subcutaneously • VWD-type 1 and type 2M | <p>Preventing excessive VWF degradation</p>  <p>Mab508</p> <ul style="list-style-type: none"> • Mab508 impairs VWF-ADAMTS13 interactions, thereby conserving high-molecular weight multimers • VWD-type 2A (IIA) • Acquired VWs-type 2A (including severe aortic stenosis, LVAD, ECMO,...) |
|--|---|---|