Identification and Functional Characterisation of von Willebrand Disease

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Abstract
The identification and functional characterisation of von Willebrand disease (VWD) is challenging due to clinical uncertainty and limitations in test processes and panels used by laboratories, and because the classification scheme does not always permit unequivocal assignment of subtype. This article reviews contemporary alternatives to classic diagnostic approaches, including the incorporation of extended core test panels inclusive of the collagen-binding assay and the potential for desmopressin (DDAVP) challenge not only to provide therapeutic information but also to assist the better characterisation of individuals with defects or deficiencies in von Willebrand factor (VWF). Supplementary assays such as the PFA-100® and the VWF propeptide assay following DDAVP challenge are also worth considering.

Keywords
Desmopressin (DDAVP), von Willebrand factor (VWF), von Willebrand disease (VWD), diagnosis, classification

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von Willebrand disease (VWD) is the most common inherited bleeding disorder and is characterised by low levels of and/or abnormal function in the plasma protein von Willebrand factor (VWF). Typically, laboratory investigation entails initial plasma testing of factor VIII coagulant (FVIII:C), VWF protein antigen (VWF:Ag) and VWF activity, which is classically assessed using the ristocetin co-factor (VWF:RCo) assay. Newer tests of VWF function include the collagen-binding assay (VWF:CB) and other putative activity (VWF:Act) assays. Depending on initial test patterns and local availability, supplementary laboratory testing may also employ VWF multimers, ristocetin-induced platelet agglutination (or aggregation) (RIPA), VWF–FVIII binding (VWF:FVIIIb) and, in some cases, genetic analysis.

Six types of VWD can be defined: types 1, 2A, 2B, 2M, 2N and 3. Type 1 VWD is a partial quantitative defect and is simply defined by a reduction in plasma VWF; thus, the presenting VWF is essentially ‘qualitatively normal’. Type 3 VWD is defined by (virtual) complete deficiency of VWF, and is diagnosed when there is essentially no measurable plasma VWF. Type 2 VWD defines qualitative defects of VWF. Type 2A VWD is defined by decreased VWF-dependent platelet adhesion and a selective deficiency of high-molecular-weight (HMW) VWF multimers, which can arise from either decreased production or increased plasma clearance. Type 2B VWD is defined by an increased affinity of VWF for its platelet receptor, glycoprotein Ib alpha (GPIba). This increased affinity typically (but not always) results in clearance of both HMW VWF and platelets from circulation, and thus (usually mild) thrombocytopenia. Type 2N VWD is characterised by markedly decreased binding affinity of VWF for factor VIII, and presents phenotypically like haemophilia A. Type 2M VWD is defined by decreased VWF-dependent platelet adhesion without selective deficiency of HMW VWF multimers. In practice, type 2M VWD comprises a composite of different functional VWF defects and essentially any qualitative defect that cannot otherwise be characterised within other type 2 VWD groups. The most common type 2M VWD variants so far identified display defective binding of VWF to GPIba, but essentially (near) normal collagen binding.

Laboratory Identification of von Willebrand Disease – Current Practice
The correct diagnosis of VWD requires both clinical and laboratory evaluation and evidence. An appropriate clinical evaluation is critical, and includes an assessment of personal and familial history of bleeding/bruising, evaluation of recent medication and a physical examination. Appropriate laboratory evaluation is also critical, but is often lacking. There are limitations in the tests used by most laboratories, test panels are often incomplete and interpretation of test data is often inadequate.

Laboratory Tests Used for the Identification and Characterisation of von Willebrand Disease
VWD is characterised by low levels of plasma VWF and/or abnormal VWF function. Ideally, a laboratory investigation would entail a panel of tests that would identify all possible presentations of VWD. Depending on local preferences, currently available test panels may include any combination of FVIII:C, VWF:Ag, VWF:RCo, VWF:C8, VWF:Act, VWF multimers, RIPA, VWF:FVIIIb and genetic analysis.

However, the actual tests, specific test methodologies and their combinations, as used by individual laboratories, vary widely and this will influence, according to the specific investigation, the
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appropriate diagnosis and typing of VWD or its exclusion. In general, the more extensive the test panel and the more thorough the investigation, the more likely the correct identification and typing of VWD. Alternatively, the use of limited test panels or poor test methodologies will compromise test accuracy and result in a high likelihood of incorrect diagnoses. This will psychologically affect individuals and compromise their therapeutic management.

von Willebrand Factor: Antigen
This is the most often used test in the investigation of VWD. It is an immunological test that detects all forms of VWF equally well (i.e. functional, dysfunctional and non-functional, high-, intermediate- and low-molecular weight). This test should not be used alone as it has no differential sensitivity to identify VWD subtypes. VWF:Ag is most commonly performed as an enzyme-linked immunosorbent assay (ELISA), by enzyme-linked immunofluorescent assay (ELFA) or by automated immunoturbidimetric procedures such as latex-immuno-assay (LIA). ELISA and ELFA assays suffer fewer technical problems, but require additional instrumentation and with a few exceptions (e.g. Vidas, bioMerieux) are unsuitable for urgent testing. LIA-based assays are suited to urgent testing and can be performed using most modern coagulation analysers, but suffer from potential technical limitations (e.g. with testing of lipaemic and icteric specimens, as well as falsely elevated values in the presence of rheumatoid factor). VWF:Ag assays show moderate inter-assay or inter-laboratory variations (~10–15%), and LIA-based results tend to be slightly higher than those obtained by ELISA. Both ELISA and LIA methods show a lower limit of assay sensitivity of around 5–10U/dl.

von Willebrand Factor: Ristocetin Co-factor
This functional assay assesses the ability of VWF to bind to GPIba in the presence of ristocetin, and is most commonly performed as a quantitative agglutination procedure using fixed or lyophilised platelets with an aggregometer or automated coagulation analyser. This assay has the ability to preferentially identify HMW and intermediate-molecular-weight VWF, and so results tend to be lower than those of VWF:Ag when these VWF forms are lacking (i.e. types 2A and 2B VWD). VWF:RCo is also lower than VWF:Ag when there is a specific defect in VWF binding to platelet GPIba (i.e. with some forms of type 2M VWD). Thus, the test combination of VWF:Ag and VWF:RCo should identify and partially distinguish all types of VWD except type 2N, with type 3 VWD showing an absence of VWF. While their role in VWD diagnostics is largely unclear and evolving, they should not be used as surrogate assays for VWF:RCo.

von Willebrand Factor: Collagen-binding Assay
These assays identify another functional property of VWF, namely its ability to bind to subendothelial matrix components, i.e. collagen in this case. Most commonly performed as an ELISA procedure, a well-optimised VWF:CB assay will demonstrate better preferential binding to HMW VWF than VWF:RCo, so that results tend to be lower than those of both VWF:Ag and VWF:RCo when there is an absence of HMW VWF forms (i.e. types 2A and 2B VWD). In theory, the test combination of VWF:Ag and VWF:CB should identify and partially distinguish all types of VWD but type 2N and possible 2M, with type 3 VWD showing an absence of VWF using both tests, type 1 showing low but concordant test results with both and types 2A and 2B yielding proportionally less VWF:CB compared with VWF:Ag (i.e. evidence of VWF functional discordance). Nevertheless, VWF:CB assays are not surrogates for VWF:RCo, as both assays detect distinct functional properties of VWF. Therefore these assays should be utilised as complementary assays. VWF:CB has a moderate inter-assay or inter-laboratory variation of around 10–20%, and a similar limit of assay sensitivity to VWF:Ag (~2U/dl). Such favourable technical features make the VWF:CB more practically ‘useful’ than the VWF:RCo assay in VWD diagnostics; however, there remains a significant lack of standardisation and because of inappropriate formulations most commercial VWF:CB assays do not preferentially recognise HMW VWF.

von Willebrand Factor: Activity
This term refers to alternative (purported) activity assays for VWF, most of which utilise a monoclonal antibody to VWF, typically directed against a functional binding site. These assays are either ELISA- or LIA-based and, in fact, are not true activity assays, although they may show some preferential binding to HMW VWF and thus may yield discordant patterns compared with VWF:Ag when testing types 2A and 2B VWD. While their role in VWD diagnosis is largely unclear and evolving, they should not be used as surrogate assays for VWF:RCo.

Factor VIII: Coagulant
As VWF is the physiological carrier of FVIII in vivo, FVIII:C testing is mandatory in the process of identification of VWD and will help guide further characterisation. FVIII:C is most commonly performed as a one-stage clotting assay, although some laboratories use a chromogenic assay. Although FVIII:C will be approximately equal to or somewhat higher than the detected level of VWF in most individuals with VWD, FVIII:C levels cannot be easily predicted. A FVIII:C level lower than that of VWF may suggest either haemophilia A or type 2N VWD.

von Willebrand Factor: Multimer Analysis
This is a time-consuming and relatively skilled procedure that aims to identify the relative distribution of molecular-weight forms of VWF, as well as assessing qualitative defects of VWF multimer structure, but that few laboratories perform. Although of potential use in the diagnosis of VWD, the appropriate identification and characterisation of the major proportion of VWD cases does not require multimer analysis.
Table 1: Expected Laboratory Phenotypic von Willebrand Factor Assay Findings in von Willebrand Disease*

<table>
<thead>
<tr>
<th>Assay Parameter</th>
<th>VWD Subtype</th>
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<tbody>
<tr>
<td></td>
<td>1</td>
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<tr>
<td>VWF:Ag</td>
<td>Low (&lt;50%)</td>
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<tr>
<td>VWF:RCo</td>
<td>Low (&lt;50%)</td>
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<tr>
<td>VWF:CB</td>
<td>Low (&lt;50%)</td>
</tr>
<tr>
<td>FVIII:C</td>
<td>Normal (&gt;0.7)</td>
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<tr>
<td>VWF:CB</td>
<td>Normal (&gt;0.7)</td>
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<tr>
<td>VWF:RCo</td>
<td>Normal (&gt;0.7)</td>
</tr>
<tr>
<td>FVIII:VWF</td>
<td>Normal (&gt;0.7)</td>
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*Absolute values noted in this table should be treated as a guide only; different laboratories utilise different values based on internal or differentially published studies. **Results in type 2M von Willebrand disease (VWD) depend on the specific mutation and defect defined. Most cases identified to date are platelet-binding dysfunctional (2M (p)) and show low von Willebrand factor (VWF):antigen (Ag), low (but concordant) VWF:collagen binding (CB), and lower (discordant) VWF:RCo, with resultant normal CB/Ag but low ristocetin co-factor (RCo)/Ag. Some rarer type 2M VWD cases with specific CB defects have also been identified (2M (c)). NA = not applicable; C = coagulant.

**von Willebrand Factor: Factor VIII:**
Performed to specifically identify type 2N VWD and to distinguish this from haemophilia A, the assay is typically performed by ELISA and performed in parallel with a standard VWF:Ag ELISA assay. VWF:FVIII assesses the ability of VWF to bind FVIII, and thus shows abnormal test results where there is a VWF:FVIII binding defect (i.e. type 2N VWD). Thus, concordant levels of VWF:Ag and VWF:FVIII indicate normal FVIII binding and exclude type 2N VWD, whereas a proportionally low level of VWF:FVIII to VWF:Ag is indicative of type 2N VWD. This assay is best performed by a VWD expert laboratory, not because the assay is difficult to perform but because its interpretation is problematic.

**Ristocetin-induced Platelet Agglutination:**
Usually performed as part of a platelet function study, this assay involves challenging patient platelets with sequential concentrations of ristocetin. Responsiveness with low concentrations of ristocetin (typically ≤0.5mg/ml) is characteristic of type 2B or pseudo (platelet-type [PT]) VWD. Alternatively, a poor response to high concentrations of ristocetin (i.e. >1.5mg/ml) would suggest (severe) type 1 VWD or types 2A or 2M VWD, whereas no response would suggest either type 3 VWD or Bernard-Soulier Syndrome. Each of these alternative possibilities would require further study for a definitive diagnosis.

**Using Laboratory Tests to Identify and Characterise von Willebrand Disease**

**Type 3 von Willebrand Disease**
Type 3 VWD is defined where plasma VWF is ‘virtually’ absent. In practice, the measured levels of plasma VWF (assayed by any VWF assay) should be <5%, although this may not always be apparent with some assays due to a lower limit of assay sensitivity (see Table 1). As plasma VWF protects and stabilises FVIII:C, plasma levels of FVIII:C are also typically low and usually also <5%. Type 3 VWD is a severe form of VWD, and patients may present clinically with a bleeding diathesis that resembles a combination of mucosal bleeding typical of VWD plus haemophilia A-like symptoms. The main difficulties with misidentification of type 3 VWD largely relate to inappropriate test panels, assay variation and lower limit of sensitivity parameters.

**Type 1 von Willebrand Disease**
This is a quantitative disorder that presents with low levels of ‘normal-functioning’ VWF. In practice, the presenting plasma levels of VWF would be similar, irrespective of the assay used to identify VWF (i.e. VWF:Ag, VWF:FVIIIc, or VWF:Act; see Table 1). Thus, the ratio of any VWF test to an alternative VWF test would be approximately 1.0 (in practice, the range of results will be in the region of 0.7–1.5). The severity of bleeding symptoms in type 1 VWD will correlate to the presenting level of plasma VWF. Our laboratory defines “severe” type 1 VWD in individuals with less than ~16% of normal plasma levels of VWF and ‘moderate’ type 1 VWD in those with levels between ~16 and ~35% VWF. The concept of ‘mild’ type 1 VWD is contentious, and some will define individuals with plasma VWF from ~35% to the lower limit of the normal reference range as having ‘mild’ type 1 VWD, whereas others may identify these individuals as having a borderline–low VWF and, while recognising such VWF ‘deficiency’ as a potential risk factor for bleeding, would not label these individuals as having VWD. In practice, VWF genetic defects are difficult to identify in individuals with presumed type 1 VWD where VWF levels exceed 35%.

**Type 2 von Willebrand Disease**
This is a qualitative disorder, with individuals presenting with dysfunctional forms of VWF and the type of dysfunction defining the VWD type.

**Type 2A von Willebrand Disease**
This defines a specific deficiency of HMW VWF, and affected individuals will therefore present with relatively lower levels of VWF:FVIIIc and VWF:CB than VWF:Ag (see Table 1). In general, this functional VWF discordance is defined by ratios of VWF:FVIIIc/VWF:Ag (RCo/Ag) and VWF:CB/VWF:Ag (CB/Ag) below ~0.7. Although VWF multimer analysis can confirm the loss of HMW VWF, in practice this is usually not required, and our own preference would be to perform a RIPA analysis and, if indicated, a DDAVP trial (as explained in later sections of this article).

**Type 2B von Willebrand Disease**
This defines individuals with hyper-adhesive VWF, who will also typically present with relatively lower levels of VWF:RCo and VWF:CB compared with VWF:Ag, or functional VWF discordance (similar to that for 2A VWD) as defined by ratios of RCo/Ag and CB/Ag below ~0.7. Although VWF multimer analysis can also be used to confirm the loss of HMW VWF, in practice this is usually not required, and would not enable its differentiation from type 2A VWD. The definitive phenotypic test for identifying type 2B VWD is RIPA. Type 2B VWD can be distinguished from the phenotypically similar PT-VWD using RIPA-mixing studies or by specific genetic analysis of the VWF and platelet GPIba genes.

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Figure 1: Simplified Algorithm Describing a Possible Approach to Improving the Identification and Functional Characterisation of von Willebrand Disease

VWD = von Willebrand disease; VWF = von Willebrand factor; Ag = antigen; RCo = ristocetin co-factor; CB = collagen binding; C = coagulant; RIPA = ristocetin-induced platelet agglutination; DDAVP = desmopressin; Mult = multimer.

**Type 2N von Willebrand Disease**

Type 2N VWF defines dysfunctional VWF to FVIII binding and individuals will present lower relative levels of plasma FVIII:C to VWF. The definitive phenotypic test for type 2N VWD, and to enable its discrimination from haemophilia A, is the VWF:FVIIIB assay, with genetic testing a feasible option.2,7

**Type 2M von Willebrand Disease**

Type 2M VWD comprises a variety of VWF defects, characterised by some dysfunction of VWF that does not fit within the other type 2 VWD categories. The majority of type 2M VWD cases so far described show a selective VWF to GPIIb-binding defect without a corresponding collagen-binding defect. Accordingly, these tend to present with relatively lower VWF:RCo than VWF:Ag (or low RCo/Ag ratios), but relatively normal (or near normal) CB/Ag. However, a small number of type 2M VWD cases showing a collagen-binding defect (with relatively normal GPIIb binding) have been reported, and these will show the opposing discordance pattern, i.e. low relative CB/Ag but normal (or near normal) RCo/Ag ratios.

A simplified algorithm highlighting the above patterns to permit a generalised means of identifying and characterising VWD can be seen in Figure 1. Our laboratory performs VWF:Ag, VWF:CB and FVIII:C testing on all cases being evaluated for possible VWD. VWF:RCo testing is also performed where there is appropriate clinical evidence of significant muco-cutaneous bleeding and/or when VWF:Ag and/or VWF:CBA ≤ 74%.

**Evident Problems with Current Identification and Laboratory Diagnosis of von Willebrand Disease**

Given appropriate selection and application of test methodologies and panels, most cases of VWD will be appropriately identified and characterised. Nevertheless, the diagnosis of VWD remains problematic for a proportion of test cases due to the under-recognised heterogeneity of VWD and because of current test limitations.1,4,8,12 For example, many investigators have
recently reported on diagnostic inconsistencies for VWD, including a series of genetic/phenotypic studies using samples derived from ‘expert’ VWD laboratories,10–12 which identified that between 15 and 33% of cases originally identified as type 1 VWD could be reclassified as type 2. Thus, a large number of type 2 VWD cases can be misidentified as type 1 VWD based on testing predominantly using a core three-test panel of FVIII, VWF:Ag and VWF:RCo, even in VWD ‘expert’ laboratories. This situation mimics that identified by analysis of testing in ‘real-world’ laboratories comprising predominantly ‘non-expert’ pathology-based clinics, and where cases are derived from a mixed clinical referral base.13–15,20–22 A similar matter regarding the misdiagnosis of type 1 VWD as type 2 VWD, and type 3 VWD as type 1 or type 2 VWD or vice versa, can also be identified.14,20–22 Most misdiagnoses occur in these investigations either because of laboratory test panel limitations or because of misinterpretations of test findings.

Such misdiagnoses can have both therapeutic implications and psychological effects on affected patients. Although misdiagnoses and misclassifications have multiple causes, errors are in part likely due to the recognised limitations in the otherwise standard panel of VWF:Ag, VWF:RCo and FVIII:C as used by most laboratories for most investigations. This would include the previously mentioned assay limitations, including the lower test sensitivity limit, plus high relative assay variability or poor reproducibility, particularly for the VWF:RCo assay. It has also previously been identified that the addition of VWF:CB testing to such a test panel will consistently reduce error rates in VWD diagnosis.22–24,45–47 Additional examples related to the misdiagnosis of VWD, and explanatory reasons are provided elsewhere.4 In brief, type 2A, 2M and 2B VWD are often misidentified as type 1 VWD because discordance of VWF:RCo and VWF:Ag (or a low RCo/Ag ratio) is not always apparent in single testing, and type 1 VWD is sometimes misidentified as type 2A or 2M VWD because false VWF functional discordance (i.e. false low RCo/Ag ratio) is sometimes identified. Occasionally, type 3 VWD is misidentified as type 1 or 2 VWD or haemophilia A. Interestingly, laboratories and clinicians sometimes misinterpret laboratory test data.6 Contrary to popular misconceptions, performance of multimer analysis is not a similar matter regarding the misdiagnosis of type 1 VWD as type 2 VWD, and type 3 VWD as type 1 or type 2 VWD or vice versa, can also be identified.14,20–22 Most misdiagnoses occur in these investigations either because of laboratory test panel limitations or because of misinterpretations of test findings.

Towards a New Paradigm to Better Define von Willebrand Disease

Expansion to a Comprehensive Test Panel that Includes the von Willebrand Factor: Collagen Binding Assay

Due to diagnostic problems, laboratories and clinicians need additional strategies to ensure the appropriate identification of VWD. In the ‘real-world VWD testing’ setting, the addition of a VWF:CB assay to the core three-test panel of FVIII, VWF:Ag and VWF:RCo as typically used by most laboratories will substantially reduce the diagnostic error rate. For example, the misidentification of type 2 VWD as type 1 VWD can be

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### Figure 2: Summary of Discriminatory Post-desmopressin Changes in Hemophilia A and in Types 1, 2A and 2M von Willebrand Disease

#### A. Pre- and post-desmopressin (DDAVP) values for factor VIII (FVIII) coagulant (C) and various VWF parameters. Dashed horizontal line indicates a nominal ‘normal’ cut-off value of 50U/dl. VWD-1s = ‘severe’ type 1 von Willebrand disease (VWD) patient group (baseline von Willebrand factor [VWF] values <16U/dl); VWD-1m = ‘mild’ type 1 VWD patient group (baseline VWF values 16–35U/dl); VWD-1p = ‘possible mild’ type 1 VWD patient group (baseline VWF values 36–65U/dl); H-A = haemophilia A patient group; VWD-2A and VWD-2M = types 2A VWD and 2M VWD patient groups, respectively. Ratios for VWF:CB:VWF:Ag and ristocetin co-factor (RCo):VWF:Ag for the same patient groups identified in A. Dashed horizontal line indicates a nominal ‘normal’ cut-off value of 0.7 as discriminatory for functional VWF discordance (i.e. <0.7 is suggestive of a functional VWD pattern). 

#### B. Pre- and post-DDAVP PFA-100® closure-times (CTs) for the same patient groups identified in A. Dashed horizontal line indicates a nominal ‘normal’ cut-off value of 200 seconds in brief, type 2A VWD is typically characterised by non-correction of PFA-100 CTs (C) and good incremental rises in FVIII:C and VWF:Ag only (A); thus, CB/Ag and RCo/Ag both tend to remain low (B). In contrast, type 2A VWD is typically characterised by non-correction of PFA-100 CTs (C) and good incremental rises in FVIII:C and VWF:Ag only (A); thus, CB/Ag and RCo/Ag both tend to remain low (B).
A progressive process for the characterisation of individuals with von Willebrand disease (VWD). With each step comes an increasing diagnostic certainty and a more comprehensive characterisation of that individual's VWD. VWF = von Willebrand factor; Ag = antigen; RCo = ristocetin co-factor; CB = collagen-binding; C = coagulant; RPA = ristocetin-induced platelet-aggregation; DDAVP = desmopressin.

reduced by >50% (i.e. from a background of around 22% to typically <10%). Use of a VWF:CB will also reduce the error rates associated with other types of VWD misdiagnoses. This is largely because optimised VWF:CB assays better discriminate HMW VWF, show a better lower limit of sensitivity and have a lower inter-assay variability than standard VWF:RCo assays. Performance of VWF:C test can also negate the need to perform mutimer analysis in many test cases.

Unfortunately, some assay standardisation concerns remain with broadly applied and commercially available VWF:CB assays that currently prevent the more universal translation of these findings.

Use of Desmopressin Challenge to Help Identify von Willebrand Disease Subtypes

Desmopressin (DDAVP) is a non-transfusional therapy often applied to individuals with VWD that acts to release endogenous (endothelial) stores of VWF. DDAVP is particularly useful for type 1 VWD, to individuals with VWD that acts to release endogenous (endothelial) stores of VWF. DDAVP is particularly useful for type 1 VWD, to

The above process can be extended to utilise additional test parameters, such as the PFA-100 (Siemens, Marburg, Germany). Similar observations were reported by our laboratory for type 1 VWD over 10 years ago using a small number of patients within a single institution study, and have also been reported by other workers employing the same extended test panel. There have been a number of recent guidelines published on the diagnosis and management of VWD, and although all mention DDAVP therapy, current recommendations are to monitor primarily using the VWF:RCo assay. This author believes that existing data provide strong support for the incorporation of VWF:C testing within this context; furthermore, use of the four-test panel noted previously (i.e. FVIII:C, VWF:Ag, VWF:CB and VWF:RCo) will assist the future identification and functional characterisation of various patients with differing types of VWD, and the better discrimination of type 1, 2A, and 2M VWD. In other words, use of this extended test panel will lessen the currently significant identification error rate otherwise obtained using the more limited but more often used test panel of FVIII:C, VWF:Ag and VWF:RCo.

In brief, the PFA-100 is very sensitive to the presence of plasma VWF, and accordingly is highly sensitive to VWD. The PFA-100 gives a single end-point value called the closure time (CT), and individuals with VWD provide prolonged CTs, in part according to the
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severity and type of VWD. Our laboratory reported many years ago in a small pilot study that in type 1 VWD, DDAVP tended to normalise all of the VWF test parameters (i.e. VWF:Ag, VWF:RCo, and VWF:CB), and also tended to correct the prolonged PFA-100 CT. In contrast, in type 2A VWD DDAVP tended to normalise only VWF:Ag, but not the functional VWF test parameters (i.e. VWF:RCo and VWF:CB), and also failed to correct the prolonged PFA-100 CT. The working hypothesis at that time was that normalisation of the PFA-100 required normalisation of functional VWF.

As an extension to the previously noted study, an evaluation of the PFA-100 has also been recently undertaken in the context of DDAVP responsiveness, and in part as a follow-up to findings recently reported by others. It was found, using a larger number of VWD cases, that in type 1 VWD normalisation of the PFA-100 CT was dependent on normalisation of functional VWF, particularly that identified by the VWF:CB assay. In contrast, in type 2A and 2M VWD normalisation of the PFA-100 was rarely achieved, consistent with the usual finding that correction of functional VWF also failed to occur. In total, the composite data (i.e. PFA-100 CT and plasma tests for VWF:Ag, VWF:RCo, VWF:CB and FVIII:C) were considered to provide additional discriminatory power for the functional identification of VWD.

Using the von Willebrand Factor Propeptide Assay to Help Identify von Willebrand Disease Subtypes

Another assay that has been evaluated within the context of the functional characterisation of VWD during DDAVP therapy is the VWF propeptide (VWFpp) assay. Before de novo synthesised VWF leaves the endothelial cell, it undergoes endoproteolytic cleavage of its propeptide (VWFpp). The processed VWF and VWFpp are released either constitutively or, following activation of the endothelium, through a regulated pathway. Of interest, the plasma half-life of mature VWF and VWFpp differs several-fold (around eight to 12 and two to three hours, respectively). This property can be exploited to evaluate the potential for reduced VWF half-life or increased VWF clearance, which has been suggested as a pathogenic mechanism in some forms of VWD. In this context, it is usual to assess the relative elimination half-lives of either or both VWFpp and VWF post-DDAVP or to calculate VWFpp/VWF ratios, either at steady state or post-DDAVP.

For example, the VWFpp/VWF ratio has been shown to be dramatically increased in type Vicenza VWD compared with normal subjects, whereas it is typically normal in most type 1 VWD patients, except for those carrying specific VWF mutations, such as C1130F, C1149R, C2671Y, S2179F and W1144G. Similarly, a very short half-life for VWF can be observed in type Vicenza VWD and patients carrying these specific VWF mutations, while most type 1 VWD patients are reported to have a half-life similar to that of normal individuals. A significant inverse correlation can be shown between VWFpp ratio and VWF half-life in both VWD patients and normal subjects. Accordingly, several workers have proposed that the VWFpp/VWF ratio is useful for distinguishing between type 1 VWD cases with a normal and a reduced VWF survival, as well as for identifying type Vicenza VWD.

The Molecular and Clinical Markers for Diagnosis and Management of 1 VWD (MCDM-1VWD) cohort have also reported on a group of type 1 VWD patients, and identified that a substantially increased ratio of steady-state VWFpp/VWF predicted a reduced VWF half-life in patients with markedly decreased VWF:Ag levels. They concluded that the systematic assay of both plasma VWF:Ag and VWFpp in moderately severe type 1 VWD patients may identify patients with a reduced VWF survival phenotype.

However, the consistency of findings in patients with type 1 VWD has recently been questioned. Thus, although post-DDAVP clearance of VWF was increased by approximately three-fold in a type 1 VWD cohort overall, this was not shown to consistently associate with steady-state VWF:Ag levels. Furthermore, increased VWF clearance was not consistently associated with increased ratios of VWFpp/VWF, indicating that a normal ratio does not necessarily reflect normal post-DDAVP survival in type 1 VWD patients. This may reflect complex heterogeneity in pathogenic mechanisms within type 1 VWD.

Genetic Analysis in von Willebrand Disease Diagnosis

There has been a recent explosion of genetic studies into VWD, including the previously mentioned studies into presumed type 1 VWD. The generally increased awareness of genetic testing and its availability leads to the undesirable situation that clinicians, often keen to exploit newly developed tests to assist in the diagnosis of patients under their care, will request such tests at odds to their true clinical utility and cost-effectiveness. In general, although recent studies are invaluable in terms of furthering our understanding of VWD, there are several limitations when attempting to translate research findings into diagnostically useful test strategies. As the VWF gene is large and complex, genetic testing for VWD is not foolproof and is typically expensive. VWD can arise from genetic events unrelated to the VWF gene, and the expression of VWF and the clinical severity in individual patients can be influenced by several epigenetic events. Currently, most of these additional complexities remain unknown.

In type 1 VWD, the search for a causative mutation may require an exhaustive and costly analysis of the entire gene, which will remain fruitless in nearly half of test cases. A significant proportion of type 1 VWD cases where a presumptive mutation is identified will also prove to be non-causal or ‘innocent’ polymorphisms. Thus, the search for mutations in the vast majority of presumptive type 1 VWD investigations cannot be encouraged. The search for mutations in presumptive type 2 VWD cases will typically be more fruitful, but will still be clinically useful only in select cases where phenotypic testing has failed to provide diagnostic clarity. Excluding potential utility for pre-natal assessment in some cases of type 3 VWD, genetic investigations in type 3 VWD will otherwise also unlikely prove to be diagnostically or clinically useful.

As the VWF gene is large and complex, genetic testing for von Willebrand disease is not foolproof and is typically expensive.

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also encouraged to consider the views of experts in the field, inclusive of recent guidelines.1,3

Conclusion
Historically, the diagnosis of VWD can be seen as a dynamically evolving process,22 beginning with the global tests of coagulation, several screening tests of platelet function that included the skin-bleeding time and progression to detection of FVIII:C and VWF:Ag. During those years, the identification of VWD remained less than optimal. The next tests of significance to emerge were those based on ristocetin (VWF:RCo and RIPA), which appeared in the 1970s. Although these permitted a sort of revolution in the investigation of VWD, diagnosis remained problematic, given the poor reproducibility of the VWF:RCo and the poor sensitivity of RIPA for VWD. The VWF:CB was originally described in 1986, and although now over 21 years of age, it still has to mature into a universally useful assay, largely because of standardisation matters.1 Interestingly, several international mutational VWD studies, which were expected to provide some definitive answers regarding phenotype-genotype correlations in VWD, have instead shown a greater complexity within VWD than previously recognised, and have also highlighted significant error rates in the diagnosis and classification of VWD among expert laboratories.23 Errors within the real world of VWD investigation continue to cause misidentification of type 2 VWD as type 1 or 3 VWD, misidentification of type 1 VWD as type 2A or type 2M VWD, and use of the VWFpp assay may permit the differential functional characterisation of individuals with types 1, 2A or 2M VWD, and use of the VWFpp assay may permit the identification of individuals with reduced VWF survival. Accordingly, perhaps it is time to consider a paradigm shift to enable the appropriate diagnosis and functional characterisation of an individual’s VWD, as depicted in Figure 3, and incorporating the use of a DDAPV challenge to identify functional changes in VWF over time. Combined use of the VWF:CB and VWF:RCoR assay may better permit the differential functional characterisation of individuals with types 1, 2A or 2M VWD, and use of the VWFpp assay may permit the identification of individuals with reduced VWF survival. Finally, the potential use of the FPIA-100 in this setting also warrants further investigation.


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