Many factor VIII products available in the treatment of hemophilia A: an embarrassment of riches?

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Abstract: Hemophilia A (HA) is a common bleeding disorder caused by the deficiency of factor VIII (FVIII) with an incidence of ~1 in 5000 male births. Replacement of FVIII is necessary to prevent and treat bleeding episodes. However, with multiple new drugs in addition to old standards, choosing among the different FVIII treatment options is harder than ever. There are FVIII products that are plasma derived or recombinant, FVIII products designed to extend the half-life of FVIII, and the first single-chain FVIII product, recombinant factor VIII single chain (rFVIII-SC). As development of inhibitors to FVIII continues to be a major problem in the care of HA patients, recent studies showing lower rates of inhibitor development with plasma-derived FVIII products versus recombinant FVIII products have made choosing among the many options now available even more complex. Although still unproven, extended half-life (EHL) products may provide the hope of decreased immunogenicity but need further testing in previously untreated patients (PUPs). This review highlights some of the differences between FVIII products currently available and hopefully assists the clinician to decide which FVIII product to choose for their patients.

Keywords: hemophilia, hemophilia A, factor VIII, recombinant factor VIII single chain, extended half-life factor VIII, immunogenicity, bleeding disorder, inhibitor development

Introduction

Hemophilia A (HA) is an X-linked bleeding disorder, first reported in medical literature from the 1800s, in which the deficiency of factor VIII (FVIII) results in problems with hemostasis.1 Before therapy with FVIII infusions became available, hemophilia patients suffered from prolonged bruising and bleeding with manifestations ranging from severe and recurrent joint hemorrhages to life-threatening bleeding episodes and had a decreased life expectancy. HA is defined as mild when endogenous FVIII levels are >5%, moderate when between 1% and 5%, and severe when <1%. Treatment for HA with replacement factor first started with the use of fresh frozen plasma in the 1950s followed by cryoprecipitate in the 1960s. Finally, in the 1970s, lyophilized FVIII derived from pooled plasma became available and revolutionized treatment for HA, even allowing home infusion therapy. The HA community then suffered from the contamination of factor supplies with the HIV in the 1980s. This tragedy stimulated research and technical advances such as heat inactivation and solvent detergent treatment to generate safer plasma-derived FVIII products. The cloning of the gene for FVIII in 1984 resulted in recombinant FVIII preparations becoming available in the 1990s. Currently, the standard of care in developed countries is to offer primary prophylactic
FVIII infusions to patients with severe HA and has led to dramatic increase in the quality of life for these patients.2,3

Now, a child born with HA is expected to have a normal lifespan as prophylactic infusions of FVIII permit normal activity without restrictions albeit with great financial burden ($150000–$300000 per year in the US4). However, due to the short half-life of FVIII, frequent infusions are required in order to maintain a trough level >1%. In addition, the development of inhibitors in 25–30% of HA patients leads to increased risk for bleeding and the need for expensive treatment regimens to induce immune tolerance or the use of bypassing agent treatments to control bleeding episode. Therefore, the development of factor products with decreased immunogenicity and longer half-lives is an important unmet need for the HA community.

Various approaches have been used for increasing the half-life of recombinant FVIII, which is a glycoprotein that is normally bound by its chaperone protein, Von Willebrand factor (VWF), which is also important in achieving hemo- stasis and influences the half-life of FVIII.5,6 Once FVIII is synthesized from the gene located on the X chromosome, it undergoes post-translational modification such as N-glycosylation and sulfation that are important for protein–protein interactions. Activation of FVIII is induced by thrombin and results in proteolysis of FVIII into its active form that is rapidly inactivated and cleared. The most common methods of prolonging the half-life of FVIII have been conjugation of FVIII with larger molecules such as polyethylene glycol (PEG), albumin, or to the human immunoglobulin Fc. Conjugation with albumin takes advantage of the natural half-life of albumin, whereas PEGylation protects FVIII from proteolytic enzymes. Fc conjugation takes advantage of the existence of the neonatal Fc receptor that allows protection and recirculation of the FVIII via this receptor pathway.3,6 Recently, a novel single-chain recombinant FVIII (Afstyla®) has been approved by the US Food and Drug Administration (US FDA) in May 2016. Recombinant factor VIII single chain (rFVIII-SC) has increased stability and affinity for VWF, extending its half-life with the possibility of decreased immunogenicity.7–11

Current factors available and extended half-life (EHL) factor development
Numerous plasma-derived FVIII products and recombinant FVIII products are now available (Table 1). First-generation recombinant FVIII products such as Recombine® were generated from the full FVIII gene and contained both human albumin and animal proteins in the production process. FVIII gene protein production results in the production of a single polypeptide that is modified and cleaved to form light and heavy chains. These light and heavy chains are held together by a labile metal ion bridge that is required for proper functioning of the protein.3 It was found that deleting most of the B domain of FVIII actually improved secretion from the cell in the recombinant process,12 and so B-domain-deleted (BDD) FVIII products such as ReFacto® were developed. Because of the concern about unknown pathogens, second-generation factors such as Kogenate® removed human albumin as a stabilizing agent. Third-generation products, introduced over the last 5 years, no longer use any animal or human products in the cell culture or production of FVIII products. The recent availability of EHL products adds to the choices of agents available to the HA community (Table 1). With the many choices now available, it is becoming more difficult and complicated to determine which FVIII product to use for which patients. This review focuses on the many new FVIII products now available to help in choosing between the different options available.

Currently, most of our patients with HA are offered prophylactic infusions with third-generation products, with some patients transitioning to the newer third-generation or EHL products now available. Because third-generation products do not use any human or animal proteins in the production of their products, these are often requested by the families. Two newly approved third-generation FVIII products with standard half-life are now available: turoctogog alfa (Novoeight®) is a BDD FVIII produced in the Chinese hamster ovary (CHO) cell line and octocog alfa (Kovaltry®) is a full-length FVIII product that is produced in the baby heavy chains. These light and heavy chains are held together by a labile metal ion bridge that is required for proper functioning of the protein.3 It was found that deleting most of the B domain of FVIII actually improved secretion from the cell in the recombinant process,12 and so B-domain-deleted (BDD) FVIII products such as ReFacto® were developed. Because of the concern about unknown pathogens, second-generation factors such as Kogenate® removed human albumin as a stabilizing agent. Third-generation products, introduced over the last 5 years, no longer use any animal or human products in the cell culture or production of FVIII products. The recent availability of EHL products adds to the choices of agents available to the HA community (Table 1). With the many choices now available, it is becoming more difficult and complicated to determine which FVIII product to use for which patients. This review focuses on the many new FVIII products now available to help in choosing between the different options available.

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EHL FVIII products have brought exciting new options for the HA community, providing the possibility of fewer scheduled infusions. Despite the various reported half-lives found in the product brochures of FVIII products, it is
commonly understood that the half-life of standard FVIII products is between 8 and 12 hours with individual variation seen with different brands. There have been several strategies for extending FVIII half-life such as attaching a PEG moiety, while both Fc fragment and albumin conjugation have only resulted in a modest increase in the half-life of FVIII, likely due to the dependence on VWF and the neonatal Fc receptor pathway.6 Although both immunoglobulins and albumin normally have very long half-lives, these approaches have only resulted in a modest increase in the half-life of FVIII, likely due to the dependence on VWF that stabilizes the protein and prevents it from degradation. Normally, FVIII is composed of a heavy and a light chain held together by a disulfide bond and is composed of a heavy and a light chain held together by a disulfide bond. An added heavy chain protein (HCP) is added to stabilize the protein and prevent its degradation.

Many FVIII products available in the treatment of HA commonly stabilize the protein and prevent its degradation.

<table>
<thead>
<tr>
<th>Generation</th>
<th>Products</th>
<th>FVIII</th>
<th>Technology</th>
<th>Half-life*</th>
<th>Date of US FDA approval</th>
</tr>
</thead>
<tbody>
<tr>
<td>Plasma derived</td>
<td>Anti-hemophilic factor (Hemofil M®, Koate-DVI®, Monarc-M®, Monoclate-P®)</td>
<td>Full length</td>
<td>Pooled human plasma</td>
<td>14.8–17.5 hours</td>
<td>1966 (Hemofil M), 1974 (Koate-DVI)</td>
</tr>
<tr>
<td>Recombinant: first generation</td>
<td>Anti-hemophilic factor recombinant (Recombinate®)</td>
<td>Full length</td>
<td>BSA in culture and human albumin as stabilizer</td>
<td>14.6 ± 4.9 hours</td>
<td>December 1992</td>
</tr>
<tr>
<td>Recombinant: second generation</td>
<td>rFVIII-FS (Helixate®, Kogenate®)</td>
<td>Full length</td>
<td>Human plasma protein solution in culture</td>
<td>13.74 hours</td>
<td>June 2000</td>
</tr>
<tr>
<td>Recombinant: third generation</td>
<td>Anti-hemophilic factor recombinant (Advate®, Kovaltry®)</td>
<td>Full length</td>
<td>No human or animal protein added</td>
<td>12–14.2 hours</td>
<td>July 2003 (Advate), March 2016 (Kovaltry)</td>
</tr>
<tr>
<td>Recombinant: second generation</td>
<td>Moroctocog alfa (ReFacto®)</td>
<td>BDD</td>
<td>Human plasma protein solution in culture</td>
<td>14.5 ± 5.3 hours</td>
<td>March 2000</td>
</tr>
<tr>
<td>Recombinant: third generation</td>
<td>Moroctocog alfa (Xyntha®), Turoctocog alfa (Novoeight®)</td>
<td>BDD</td>
<td>No human or animal protein added</td>
<td>10.8–12 hours</td>
<td>February 2008 (Xyntha), October 2013 (Novoeight)</td>
</tr>
<tr>
<td>Recombinant: fourth generation</td>
<td>Simoctocog alfa (Nuvuviq®)</td>
<td>BDD</td>
<td>HEK cells to allow human glycosylation</td>
<td>17.1 ± 11.2 hours</td>
<td>September 2015</td>
</tr>
<tr>
<td>Recombinant: third-generation EHL</td>
<td>Octocog alfa pegol (Adynovate®)</td>
<td>BDD-PEGylated</td>
<td>PEGylation to parent drug Advate</td>
<td>14.69 ± 3.79 hours</td>
<td>December 2016</td>
</tr>
<tr>
<td>Recombinant: fourth-generation EHL</td>
<td>rFVIII–Fc (Eloctate®)</td>
<td>BDD-rFVIII–Fc</td>
<td>HEK cells to allow human glycosylation</td>
<td>19.7 ± 2.3 hours</td>
<td>June 2014</td>
</tr>
<tr>
<td>Recombinant: third-generation EHL</td>
<td>rFVIII–SC (Afostyla®)</td>
<td>EHL single chain</td>
<td>No human or animal protein added</td>
<td>14.2 hours</td>
<td>May 2016</td>
</tr>
</tbody>
</table>

Note: The half-life of the different factors was taken from the product brochures from the manufacturers and differs in how it was determined.

Abbreviations: FVIII, factor VIII; US FDA, US Food and Drug Administration; VWF, Von Willebrand factor; BSA, bovine serum albumin; rFVIII, recombinant factor VIII; BDD, B domain deleted; HEK, human embryonic kidney; EHL, extended half-life; PEG, polyethylene glycol; SC, single chain; rFVIII–FS, recombinant FVIII formulated with sucrose; rFVIII–Fc, anti-hemophilic factor (recombinant), Fc fusion protein; rFVIII–SC, anti-hemophilic factor (recombinant), single chain.

The US FDA approved both PEGylated FVIII (Adynovate®) and the Fc–FVIII fusion product (Eloctate®). Clinical studies thus far showed that rFVIII–Fc fusion product (Eloctate) has the longest half-life among the EHL products currently available with a half-life of 19.7 hours in adolescents and adults, although it decreased to 14.6 in patients 6–12 years old.20,21 Interestingly, when the same BDD FVIII gene is expressed using the HEK cell line allowing for human-like post-translational modifications, the resultant FVIII protein’s half-life also appears to be slightly increased. Simoctocog alfa (Nuvuviq) was developed with the hope to generate a less immunogenic FVIII product by eliminating potentially immunogenic glycosylation patterns found in non-human cell lines, and it appears that the humanized post-translational modifications may have also prolonged the half-life.22 Further studies, however, are needed to determine if these new products are indeed less immunogenic in previously untreated patients (PUPs).

Finally, using recombinant DNA technology, CSL Behring created a novel recombinant FVIII to modestly extend the half-life by increasing the affinity of FVIII to VWF, which stabilizes the protein and prevents it from degradation. Normally, FVIII is composed of a heavy and a light chain held together by a disulfide bond.
together by a labile metal ion bridge. CSL Behring developed a BDD recombinant FVIII molecule in which the heavy and light chains are covalently linked that has a twofold increased affinity to VWF.\textsuperscript{7,23} The increased stability provided by the single-chain covalent bond permits a higher efficiency manufacturing of the full-length factor and results in a more stable protein after reconstitution.\textsuperscript{24} Moreover, the enhanced affinity for VWF may accelerate the formation of VWF:FVIII complex that not only increases the half-life but also may result in decreased immunogenicity by inhibiting endocytosis of FVIII by dendritic cells.\textsuperscript{25,26} These modifications resulted in a FVIII molecule with an increased half-life that is comparable to the PEGylated EHL product (Adynovate).\textsuperscript{5,10,27,28} When FVIII is normally activated, the heavy and light chains are cleaved and released from VWF. Because these key thrombin cleavage sites in the single-chain FVIII are intact, once activated, rFVIII-SC becomes structurally identical to FVIII.

Preclinical testing in animals confirmed that recombinant rFVIII-SC had equal potency compared to other full-length and BDD FVIII products.\textsuperscript{11} In the AFFINITY clinical trial program, rFVIII-SC showed excellent hemostatic efficacy with a median annualized bleeding rate (ABR) of 1.14 that is comparable to other recently approved factor products.\textsuperscript{8} For example, rFVIII-Fc (Eloctate) reported a median ABR of 1.6, and simoctocog alfa (Nuwiq) reported a median ABR of 0.9. Furthermore, no participants (previously treated patients [PTPs] with >150 exposure days to other FVIII products) in the AFFINITY study developed an inhibitor, and the immunogenicity of rFVIII-SC is currently being studied in PUPs (#NCT02172950).\textsuperscript{8}

**Inhibitor development and immunogenicity of new EHL FVIII**

The most feared complication in the treatment of HA is the development of neutralizing anti-FVIII antibodies or “inhibitors” that occur in up to 25–30% of HA patients.\textsuperscript{29} The development of inhibitors results in the need for new treatments with bypassing agents and cumbersome and expensive immune tolerance induction protocols. Why the immunogenicity of FVIII products is so high is unclear, but it seems to be a combination of ethnicity, genetic mutation, and environmental factors.\textsuperscript{2} Also important are the differences in the post-translational modification patterns of FVIII – dependent on the cell line used, whether non-mammalian cell lines (e.g., CHO or BHK) or the human cell line (HEK) – and the role of VWF in both increasing the half-life of FVIII and reducing the endocytosis of FVIII by antigen-presenting cells.\textsuperscript{2}

With regard to ethnicity, Black HA patients have twice the rates of inhibitor development than White HA patients. It has been thought that this is due to the differences in genetic haplotypes of the patient’s FVIII gene versus the haplotypes that are found in various plasma-derived and recombinant FVIII products.\textsuperscript{30} Gunasekera et al\textsuperscript{11} reported that African Americans with intron-22 inversion had two- to threefold higher risk of inhibitor development than White HA patients with the same inversion. However, recent studies looking at the inhibitor risk in severe HA patients and ethnic haplotypes did not find a correlation between the haplotypes and inhibitor formation.\textsuperscript{31,32} Thus, whether inhibitor risk is increased due to ethnic differences in the FVIII gene compared to commercially available FVIII products remains unanswered.

Recently, the Research of Determinants of Inhibitor Development (RODIN) study group reported that second-generation FVIII products had higher inhibitor rates than third-generation products. This finding was confirmed by Calvez et al in their report of a French cohort of HA patients that also showed higher rates of inhibitor development (37.5%) with second-generation FVIII products\textsuperscript{33,34} and an adjusted hazard ratio of 1.58 when they compared the rates of inhibitor formation between second-generation and third-generation products. The authors speculated that this might be related to the cell line used for the production of the product: BHK cells (second generation) vs CHO cells (third generation) as the degree of product modifications such as sulfation and glycosylation differ between the two cell types.\textsuperscript{34}

Another potential benefit of the EHL FVIII products is the possibility of decreased immunogenicity of these new factor products. It must be emphasized that this is not yet proven and requires further studies in PUPs. However, it has long been demonstrated that conjugation of immunogenic peptides to a non-immunogenic protein carrier can induce tolerance and that immunoglobulin conjugation was superior to other protein carriers such as albumin.\textsuperscript{35} Recent case reports using the recombinant FVIII–Fc fusion protein in immune tolerance induction for HA patients with inhibitors also support this notion.\textsuperscript{36,37} Both Fc and albumin conjugations can potentially decrease the immunogenicity of FVIII by what is called antigen shielding, where the fusion moiety inhibits uptake of the FVIII product by the antigen-presenting cells needed to generate an immune response.\textsuperscript{38}

Conjugation of PEG to a molecule not only prolongs the half-life but also is thought to make it less immunogenic by inhibiting the binding of protein to endocytic receptors on antigen-presenting cells.\textsuperscript{39} This strategy has been utilized to
increase the half-life of multiple drugs now. For example, it is known that PEGylated asparaginase, which is used for the treatment of leukemia, causes fewer allergic reactions than the native asparaginase enzyme.40

With the advent of various EHL FVIII products now approved or in development utilizing the Fc fusion, albumin fusion, or PEGylation, further clinical trials are needed to determine if they also result in decreased immunogenicity. While animal studies support the notion that these new products have decreased immunogenicity, clinical studies with PUPs will be especially important. In addition, the effects of long-term exposure to a molecule such as PEG in HA patients still need to be evaluated.

**Immunogenicity of plasma-derived FVIII products versus recombinant FVIII products**

Recently reported results from the clinical trial called Study on Inhibitors in Plasma-Product Exposed Toddlers (SIPPET) trial are that plasma-derived FVIII products had lower rates of inhibitor formation than recombinant FVIII products in that population, complicating what we thought we knew was best for the treatment of patients with HA.42 While the use of recombinant FVIII has become the standard in most US and European clinics, the results of the SIPPET trial now further confuses the choice of factor that is best for each patient. What can be the reasons behind the decreased immunogenicity of plasma-derived FVIII products? One possible answer is that most recombinant FVIII products are produced using a non-human cell line and contain differences in the post-translational modifications of the resulting FVIII protein.54 Another potential reason is that preformed complexes containing VWF result in decreased immunogenicity, which is supported by animal studies.25 If human glycosylation decreases the immunogenicity of FVIII, use and studies of the newly approved FVIII products such as simoctocog alfa (Nuwiq) and rFVIII–Fc (Eloctate) should help to prove this point, as these are made using HEK cell line. It should be noted that pharmacokinetic studies showed that simoctocog alfa (Nuwiq) has increased half-life compared to Advate®, which is the same recombinant product made in non-human cell lines.25 It is thus tempting to speculate that human glycosylation pattern increases the affinity of FVIII to VWF, which not only increases the half-life but also decreases the immunogenicity of the FVIII.3 It should also be mentioned that plasma-derived FVIII products contain many other proteins as well that may influence FVIII immunogenicity.

With regard to the SIPPET trial, the medical and scientific advisory council (MASAC) of the National Hemophilia Foundation has highlighted the fact that the SIPPET trial differed from previous prospective studies comparing plasma-derived FVIII to recombinant FVIII such as RODIN and the European Pediatric Network trial (PedNet), which found no significant differences.43 Because the ethnicities in the SIPPET trial (mostly from Egypt, India, and Iran) differed significantly from the ethnicities of the patients in the other studies, they concluded that the results may not be applicable to the US HA community that are mostly Caucasian. In addition, they pointed out that in the US, majority of patients receive prophylactic infusions and that third-generation products are in use that differed from the study population. They recommended that the SIPPET study results may not be applicable to the US population and that the risk of inhibitor development must be weighed against the potential infectious risk of emerging viruses, which has not been tested with the current manufacturing processes.44

**Incorporating the EHL FVIII products into clinical practice**

Based on in vitro and animal studies done thus far, EHL FVIII products may also have a potential benefit of decreased immunogenicity and inhibitor development in addition to prolonged half-life and are very attractive to implement into clinical practice.39 However, it is worth noting that individualized dosing must be done using observed pharmacokinetic values, as the half-life of each product can differ between patients, and in individual cases, the use of EHL FVIII products may not prolong the half-life enough to justify the increased cost of these new factors. For example, despite the longest half-life advertised of any of the factors available, the actual half-life is much shorter in younger patients, and we have been unable to go to less than thrice-weekly infusions of rFVIII–Fc (Eloctate) in order to maintain a trough level >1%.

Nevertheless, it is exciting to have so many options in coming up with individualized treatment programs for HA patients.

So, what is the best approach to incorporate these new FVIII products into clinical practice? It has been suggested that the EHL products may be helpful in patients who have decreased compliance since these may decrease the frequency of infusions. In addition, in a patient who is having breakthrough bleeding with an already aggressive prophylactic regimen, it is reasonable to try an EHL product in an attempt to decrease breakthrough bleeds.4 Although there have been concerns about potential inhibitor development in switching FVIII products, there is little evidence that inhibitor rates...
increase with switching FVIII products. However, immuno-
nologically, recombinant products do have varying levels of
post-translational protein modifications due to production
in different cell lines that may provide potential epitopes for
immune reaction.\(^{38}\) Therefore, further studies are needed to
verify whether these new EHL products indeed decrease
immunogenicity. Unfortunately, it is likely that new clinical
studies in PUPs will be underpowered to detect a difference
in immunogenicity and will make it difficult to have convinc-
ing scientific basis to declare the least immunogenic product.
For PTPs, although it may make sense to switch to a product
that is generated in the same cell line so that glycosylation
patterns remain the same, the overwhelming evidence from
clinical studies shows that PTPs hardly ever develop an
inhibitor after a FVIII product switch.

**Conclusion**

Treatment options for HA have never been better with the
approval of several new recombinant FVIII products and the
new EHL FVIII products that are now even safer and may
offer the potential of decreased immunogenicity and longer
half-lives. Additional factor products that are still in develop-
ment (Table 2) will add to the choices that we will have
in the treatment of HA patients.\(^{4,19}\) However, there remains
much work to be done in terms of satisfying the unmet needs
for HA patients. The rate of inhibitor development is still the
scourge of HA treatment, and improvements are needed to
minimize the development of inhibitors while optimizing
immune tolerance induction protocols. Novel products cur-
rently in development continue to offer exciting possibilities.
New products such as rFVIII-VWF-XTEN in animal studies
allow subcutaneous dosing with fourfold increase in half-life,
and the development of a humanized bispecific antibody
that provides FVIII-mimetic function provides exciting new
possibilities in the treatment of HA.\(^{5,45,46}\) Perhaps, with the
introduction of many new EHL products to the market, the
cost of FVIII infusions will decrease enough to allow more
children in developing countries to receive prophylactic
therapy. Moreover, the recent success with gene therapy tri-
als for factor IX deficiency provides hope that we will soon
overcome the obstacles in developing an engineered gene
therapy construct for FVIII that will result in a cure.\(^ {47} \)

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National Military Medical Center, Uniformed Services Uni-
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scientific instrumentation, or organization is considered an
integral part of the scientific endeavor and does not consti-
tute endorsement or implied endorsement on the part of the
author, DoD, or any component agency. The author reports
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**Table 2** FVIII products in development

<table>
<thead>
<tr>
<th>Generation</th>
<th>Products</th>
<th>FVIII</th>
<th>Technology</th>
<th>Half-life (h)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Recombinant: third-generation EHL</td>
<td>N8-GP</td>
<td>BD-modified glycoPEGylated</td>
<td>GlycoPEGylated addition to FVIII</td>
<td>18.4–19.04</td>
</tr>
<tr>
<td>Novel bispecific antibody</td>
<td>Emicizumab</td>
<td>None</td>
<td>Bispecific antibody with FVIII-mimetic function</td>
<td>weeks</td>
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</table>

**References**

2. Coppola A, Di Capua M, Di Minno MN, et al. Treatment of hemo-
3. Orlova NA, Kovnir SV, Vorobiev II, Gabilov AG, Vorobiev AI. Blood
5. Pipe SW, Montgomery RR, Pratt KP, Lenting PJ, Lillicrap D. Life in
the shadow of a dominant partner: the FVIII-VWF association and its clini-
6. Shapiro AD. Long-lasting recombinant factor VIII proteins for
hemophilia A. Hematology Am Soc Hematol Educ Program.
7. Bensen-Kennedy D. Bringing new therapy options to the hemophilia
rVIII-SingleChain: results of a phase 1/3 multicenter clinical trial in
9. Pabinger-Fasching I. The story of a unique molecule in hemophilia
A: recombinant single-chain factor VIII. Thromb Res. 2016;141
(suppl 3):S2–S4.
and pharmacodynamics of rVIII-SingleChain, a novel recombinant single-chain
11. Zollner SB, Raquet E, Müller-Cohrs J, et al. Preclinical efficacy and
safety of rVIII-SingleChain (CSL627), a novel recombinant single-chain


44. MASAC: MASAC Recommendation on SIPPET (Survey of Inhibitors in Plasma-Product-Exposed Toddlers): RESULTS and Recommendations for Treatment Products for Previously Untreated Patients with Hemophilia A. MASAC; 2016.

