Impact of novel hemophilia therapies around the world

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Abstract
Hemophilia A and B are hereditary bleeding disorders, characterized by factor VIII or IX deficiencies, respectively. For many decades, prophylaxis with coagulation factor concentrates (replacement therapy) was the standard-of-care approach in hemophilia. Since the 1950s, when prophylaxis started, factor concentrates have been improved with virus inactivation and molecule modification to extend its half-life. The past years have brought an intense revolution in hemophilia care, with the development of nonfactor therapy and gene therapy. Emicizumab is the first and only nonreplacement agent to be licensed for prophylaxis in people with hemophilia A, and real-world data show similar efficacy and safety from the pivotal studies. Other nonreplacement agents and gene therapy have ongoing studies with promising results. Innovative approaches, like subcutaneous factor VIII and lipid nanoparticles, are in the preclinical phase. These novel agents, such as extended half-life concentrates and emicizumab, have been available in resource-constrained countries through the constant efforts of the World Federation of Haemophilia Humanitarian Aid Program. Despite the wide range of new approaches and therapies, the main challenge remains the same: to guarantee treatment for all. In this article, we discuss the evolution of hemophilia care, global access to hemophilia treatment, and the current and future strategies that are now under development. Finally, we summarize relevant new data on this topic presented at the ISTH 2021 virtual congress.

KEYWORDS
blood coagulation factors, emicizumab, factor IX, factor VIII, genetic therapy, hemophilia

Essentials
- Replacement therapy has been the standard of care for hemophilia since the late 1950s.
- Emicizumab, the first nonfactor therapy for hemophilia A, changed the hemophilia care scenario.
- Rebalancing agents and gene therapy are new options with ongoing studies and promising results.
- The main challenge remains the same: guarantee treatment for all.
1 | INTRODUCTION

Hemophilia A and B are congenital X-linked bleeding disorders caused by factor VIII (FVIII) or factor IX (FIX) deficiency, respectively. Clinical manifestation correlates with the residual endogenous clotting factor activity. People with FVIII or FIX plasma levels of <1 IU/dL are classified as having severe hemophilia and may have spontaneous bleeding events. In moderate (1-5 IU/dL) and mild (5 to ≤40 IU/dL) hemophilia, bleeding symptoms are usually associated with trauma or surgical procedures. In hemophilia, the most frequent bleeds occur in joints and muscles, resulting in chronic and progressive arthropathy with significant crippling morbidity. In addition, there is a risk of life-threatening bleeding, such as intracranial hemorrhage, particularly in people with severe phenotype, in the absence of adequate treatment.

Since the 1950s, several significant achievements have happened for hemophilia A and B, resulting in considerable improvements in hemophilia care and patient quality of life (Figure 1). Among these, the main improvement is the availability of safe options to replace the missing clotting factor and restore hemostasis. The so-called replacement therapy, using plasma-derived or recombinant products, has been considered the cornerstone for hemophilia treatment and ensures the adoption of prophylactic therapy. However, even with the progress achieved with bioengineered clotting factors, including extended half-life (EHL) FVIII or FIX products, replacement therapy is costly, requires burdensome frequent intravenous injections, and has the risk for development of inhibitors. These factors compromise adherence and access to adequate treatment for people with hemophilia worldwide.

New products have recently been developed. Nonreplacement therapies, including emicizumab and rebalancing products, are transforming the approach for hemophilia treatment. These products are administered subcutaneously and are effective prophylactic options, regardless of the presence of inhibitors. More recently, adeno-associated virus (AAV) vector-mediated gene therapy trials for hemophilia A and B have presented promising results.

Despite the several new options and strategies for hemophilia care, access to treatment still represents a critical challenge for most people with hemophilia worldwide and more needs to be done to guarantee adequate treatment for all people with hemophilia. In this article, we discuss the evolution of hemophilia care, the access to hemophilia treatment worldwide, and the current and future strategies that are now under development. Finally, we summarize relevant new data on this topic presented at the ISTH 2021 virtual congress.

2 | IMPROVING HEMOPHILIA CARE

Until the early 1960s, the only available treatment for hemophilia was based on whole blood or fresh plasma transfusion, which was insufficient to avoid most bleeding complications. Consequently, most people with severe hemophilia died in childhood and early adulthood. In 1958, Inga Nilsson and colleagues in Malmo, Sweden, were pioneers to use the regular prophylactic infusion of a local...
factor human fraction I-0 (antihemophilic factor concentrate) containing FVIII to convert the bleeding phenotype temporarily from severe to moderate, resulting in a significant decrease in bleeding episodes and reduction of the impact of arthropathy, particularly when started at an early age.5

Nevertheless, only after the advent of cryoprecipitate6 and, later, lyophilized plasma-derived FVIII and FIX concentrates, did home treatment become possible, a crucial step to guarantee the early treatment of the bleeding episodes and regular prophylaxis.7 However, the first plasma-derived clotting factor concentrates had no viral inactivation methods applied to their manufacturing process. It was only in 1985 that these methods were incorporated, reducing the risk of blood-borne infections drastically:3 Meanwhile, the cloning of FVIII and FIX genes (in 1982 and 1984, respectively) enabled the development of virus-free recombinant FVIII and FIX concentrates.8

The availability of safe FVIII and FIX products contributed to the evolution of prophylaxis as a feasible treatment modality. The benefits of prophylaxis have been recognized since the first publications. However, primary prophylaxis became the evidence-based standard of care for hemophilia after results from a randomized clinical trial conducted by Manco-Johnson et al.9 In fact, prophylaxis is preferable to episodic treatment, even when using a lower dose of factor concentrates. Long-term prophylaxis is recognized as the standard of care for all people with hemophilia with severe clinical phenotype10 and has been proven to be effective in preventing life-threatening bleeds and joint damage9 (Figure 1).

3 | NEW THERAPIES FOR HEMOPHILIA

3.1 | Replacement therapy: Extended half-life products

For many decades, hemophilia treatment was based on replacement therapy using plasma-derived clotting factors or recombinant products. However, advancing technology made it possible to develop unmodified recombinant products with a standard half-life (SHL), and clotting factor concentrates with EHL.3

EHL clotting factor concentrates are bioengineered molecules with increased half-life by at least 1.3 times over that of SHL FVIII or FIX concentrates.11 Different technologies were used for the development of EHL products. These include the conjugation with polyethylene glycol and fusion with other proteins, such as albumin or the fragment crystallizable (Fc) of IgG1. Table 1 summarizes the characteristics of the currently licensed EHL products.

EHL products were developed to lead to higher factor peaks and trough levels, decreasing the frequency of intravenous injections and reducing the burden of prophylaxis. These strategies to improve pharmacokinetic (PK) parameters resulted in a significant extension of FIX concentrates half-life, usually 3 to 5 times longer than SHL-FIX products.12-16 However, EHL-recombinant FVIII (rFVIII) products achieved only 1.5 to 1.8 times longer half-life than SHL-FVIII products17-24 This minor improvement in PK parameters is probably due to the function of von Willebrand factor (VWF). In the circulation, FVIII needs to be bound to VWF for stabilization. Therefore, the maximum half-life achieved by EHL-rFVIII products is the same as VWF’s half-life.25

More recently, a new EHL-rFVIII product has been under development to overcome this effect. BIVV001 (rFVIII-Fc-VWF-XTEN) is a novel fusion protein with two different technologies. A single recombinant B-domain deleted (BDD) FVIII protein is fused to the FVIII-binding D′D3 domain of VWF via IgG1 dimeric Fc domain and two XTEN polypeptides. The covalent link to the VWF D′D3 domain prevents binding between the rFVIII and endogenous VWF. This strategy confers to BIVV001 a fourfold longer half-life than SHL-FVIII products, a benefit similar to those from the EHL-rFIX products.26 Until later 2021, BIVV001 was on phase 3 clinical trial and not yet commercially available.

3.2 | Nonreplacement therapy

Although replacement therapy has been the standard therapeutic option to repair the hemostatic defect in hemophilia for several decades, it has limitations and challenges. That includes the burden from recurrent intravenous infusions, which compromises the adherence and, as a result, the efficacy of prophylactic treatments, even using EHL products. In this context, nonreplacement therapies could fulfill these unmet needs in hemophilia care.6

Nonreplacement therapy is a class of products developed using strategies beyond the concept of replacing the deficient clotting factor. These novel agents aim to either restore the hemostasis using mimetic products or establish the rebalance of the hemostasis, inhibiting the anticoagulant pathways. In addition, these nonreplacement products are administered subcutaneously, overcoming the burden associated with frequent intravenous administration. The subcutaneous route can be particularly exciting when caring for young children or patients with poor venous access. These agents also address the challenging scenario of managing patients with neutralizing anti-FVIII or anti-FIX antibodies (inhibitors), whose bleeding episodes are more frequent and difficult to control (Table 2).

3.2.1 | Factor VIII mimetics

In this setting, the approach to substitute rather than replace FVIII was highly successful. Emicizumab is the first nonreplacement therapy approved for prophylaxis in patients with hemophilia A with and without inhibitors. It is a humanized bispecific monoclonal antibody with binding sites to activated factor IX and factor X, mimicking FVIII in its cofactorial activity.27 It increases thrombin generation in patients with hemophilia A, regardless of their inhibitor status. Other advantages are its long half-life and good bioavailability, making it possible to achieve a stable hemostatic effect with subcutaneous dosing each 1, 2, or 4 weeks.26
| Product |
|---------|
| Factor VIII |
| Efmoroctocog alfa (Elocta, Eloctate) |
| Rurioctocog alfa pegol (Adynovi, Adynovate) |
| Damoctocog alfa pegol (BAY 94-9027, JIVI) |
| Turoctocog alfa pegol (N8-GP, Esperoct) |
| Factor IX |
| Eftrenonacog alfa (Alprolix) |
| Nonacog beta pegol (N9-GP, Refixia, Rebinyn) |
| Albutrepenonacog alfa (Idelvon) |

| Method to extend half-life | Dosing scheme | Half-life, h | Immunogenicity PTP | Immunogenicity PUP | References |
|---------------------------|----------------|-------------|---------------------|-------------------|------------|
| IgG1-Fc-fusion            | 25–65 IU/kg every 3-5 days | 19 (OSA) | No inhibitor | 31.1% inhibitor | 17,21,22,101 |
| Random PEGylation         | 45 ± 5 IU/kg twice per week | 14.3–16 (OSA) | No inhibitor | – | 18 |
| Site-specific PEGylation  | 30–60 IU/kg every 3-7 days | 19 (OSA) | No inhibitor | – | 20 |
| Site-specific glycoPEGylation | 50 IU/kg every 4 days 75 IU/kg per week | 19.9 (OSA) | 0.6% inhibitor | – | 19,23,24 |
| IgG1-Fc-fusion            | 50 IU/kg per week 100 IU/kg every 10 days | 82.1 | No inhibitor | 3% inhibitor | 12,13 |
| Site-specific glycoPEGylation | 40 IU/kg per week | 111 | No inhibitor | 6.1% inhibitor | 15,102,103 |
| Albumin fusion            | 35–50 IU/kg/wk 75 IU/kg every 10-14 days | 101.7 | No inhibitor | 1.6% hypersensitivity (n = 1) | 16 |

Abbreviations: Ab, antibody; CSA, chromogenic substrate assay; OSA, one-stage clotting assay; PTP, previously treated patients; PUP, previously untreated patients.

*Dosing scheme may vary between different countries.*
A series of emicizumab pivotal clinical trials (HAVEN 1-4) have confirmed efficacy and safety for children and adults with hemophilia A, with and without inhibitors. These trials have also reported a significant impact on the quality of life for both children and adults. Efficacy and safety in the pediatric population were also confirmed in the HOHOEMI trial, which included children as young as 4 months old. Reports from different hemophilia centers have confirmed the safety reported in the pivotal studies but observed a higher risk of breakthrough bleeds in older people.

Prophylaxis with emicizumab in previously untreated people is still under debate, while the incidence and consequences of anti-FVIII inhibitors are unknown in this scenario. Many trials are on the way to try to address this open issue. HAVEN 7 is now recruiting and will evaluate early prophylaxis with emicizumab in children under 12 months of age with hemophilia A without inhibitors. The Hemophilia Inhibitor Prevention Trial is a phase 3 randomized, open-label study that will compare inhibitor data from previously untreated people under prophylaxis with an EHL-rFVIII concentrate or emicizumab (NCT04303559). Another trial will also assess inhibitor data from primary prophylaxis with emicizumab and a concomitant low dose of simoctocog alfa, a recombinant FVIII concentrate (NCT04030052). Both trials are now recruiting.

Emicizumab also had a positive impact on the challenging scenario of periprocedural management. In the past few years, many reports with real-world data have shown the safety of managing minor and major procedures in people with and without inhibitors. Currently, we observe a learning curve for both patients and clinicians has advanced since emicizumab was licensed for people with hemophilia A. This disruptive approach to hemophilia also changed the way bleeds are identified and treated. Therefore, data on real-world use, including bleeding and periprocedural management, is essential to build knowledge outside clinical trials. Societies and groups of experts have also issued guidelines on bleeding treatment, perioperative management, and laboratory surveillance for emicizumab.

Other unanswered questions come from scenarios where emicizumab probably does not fully substitute FVIII. One of them is bone metabolism and health. The EmiMK study will evaluate bone health in patients under emicizumab (NCT04131036). Prophylaxis for sports is another intriguing scenario to be addressed in the STEP study, where emicizumab and FVIII concentrates will be compared for prophylaxis for provoked bleeding (NCT05022459).

Emicizumab use in other populations is also currently under evaluation in clinical trials. HAVEN 6 is designed to evaluate prophylaxis with emicizumab in people with mild and moderate hemophilia A without inhibitors (NCT04158648). Another ongoing trial will address the efficacy of prophylaxis with emicizumab in acquired hemophilia A (NCT04188639).

Since emicizumab was approved, new activated FVIII mimetic bispecific antibodies have been developed, such as BS-027125 (Bioverativ, Waltham, MA, USA) and Mim8 (Novo Nordisk, Bagsvaerd, Denmark). Preclinical analysis of Mim8, including in vitro
assays with FVIII deficient human plasma and in vivo assay using a hemophilia A mouse model, suggest that this new bispecific antibody could be more potent than emicizumab.\(^\text{40}\)

### 3.2.2 | Rebalancing therapies

This new class of products restores the hemostatic capacity in the blood, mainly inhibiting different natural anticoagulant pathways, establishing the hemostatic balance even in the absence of FVIII or FIX and the presence of inhibitors. At present, no rebalancing agents are licensed, and their use is limited to clinical trials (Table 2).

Tissue factor pathway inhibitor (TFPI) is an anticoagulant protein that can reversibly inhibit activated factor X, either by direct inhibition or by complexing with activated factor X and then inhibiting tissue factor and activated factor VII. Two monoclonal antibodies with anti-TFPI activity are now in phase 3 clinical studies: concizumab and marstacimab. The first agent may be administered subcutaneously, at 0.15 or 0.25 mg/kg, once daily. Marstacimab is evaluated with a single loading dose of 300 mg and then 150 mg or 300 mg weekly, depending on the person’s bleeding phenotype. Updates for both molecules were presented at ISTH 2021 congress (see ISTH 2021 Virtual Congress below).\(^\text{41-43}\)

Fitusiran, an antithrombin small interfering RNA, is now an ongoing phase 3 trial. In the randomized, open-label ATLAS-INH study (NCT03417102), 38 people with inhibitor (29 with hemophilia A and 9 with hemophilia B) with a mean age of 26.8 years (± 9.8) received once-monthly fixed doses of fitusiran 80 mg subcutaneously. A significant reduction of bleeding episodes was observed when compared to 19 people with inhibitor (16 with hemophilia A and 3 with hemophilia B) with a mean age of 28.4 years (± 11.1) receiving on-demand bypassing agents (BPAs), with 65.8% of patients with zero bleeding events in the fitusiran arm, compared to 5.3% in the on-demand BPA group. In addition, two people (5.3%) receiving fitusiran were reported with venous thrombosis events, and fitusiran was discontinued for one person.\(^\text{44}\) In the phase 3 ATLAS-A/B study (NCT03417245), people with severe hemophilia A or B without inhibitor, previously treated with on-demand factor concentrate, were randomly assigned to receive a fixed dose of fitusiran 80 mg once monthly or continuous with on-demand treatment. As expected, a significant reduction was observed in the fitusiran arm, but only 50.6% (40 of 79 people who completed 9 months of fitusiran prophylaxis) had zero treated bleeds during the study period.\(^\text{45}\)

SerpinPC is a highly specific activated protein C inhibitor, currently on phase 1/2a trial.\(^\text{46}\) In a press release in September 2021, Centessa Pharmaceuticals and its subsidiary ApcnteX Limited announced some results from its phase 2a proof-of-concept trial (AP-0101). Twenty-three people with severe hemophilia and on-demand therapy were enrolled in this trial (19 with hemophilia A and 4 with hemophilia B, both without inhibitors). Three doses were evaluated (0.3, 0.6, and 1.2 mg/kg) with subcutaneous administration every 4 weeks, and the primary outcome was assessed after 24 weeks of follow-up. Results presented a median 88% reduction in all bleeds (from 36.0 to 4.4) and a median 94% decrease in spontaneous joint bleeds (from 21.1 to 2.2) for the highest dose. No venous thromboembolism or other concerning adverse effects were observed. Two people developed anti-drug antibodies (ADAs) but displayed no apparent impacts on their bleeding phenotype.\(^\text{37}\)

### 3.3 | Gene therapy for hemophilia

Hemophilia has always been an attractive candidate for gene therapy. For decades, hemophilia gene therapy research groups have been dedicated to finding the ideal strategy for achieving lasting plasma factor levels with a one-time treatment. Current gene therapy clinical trials for hemophilia are using the same strategy. They are based on intravenous administration of the liver-directed delivery of FVIII or FIX transgene using recombinant nonintegrating AAV vectors (Figure 2A).

The first AAV liver-directed gene therapy clinical trial for hemophilia B was critical to establish the current successful strategies.\(^\text{48}\) This study used an AAV2 vector administered into the hepatic artery and revealed essential issues related to AAV vector immunogenicity. One was associated with the preexisting immunity due to the presence of neutralizing antibodies (NAbs) to the AAV vector capsid. NABs are AAV serotype specific and may impair the vector transduction efficacy. A plausible strategy to avoid this negative effect is to exclude people with anti-AAV vector antibodies. Another critical issue is related to the AAV capsid-mediated cellular immune response. Transduced cells may have a transient expression of AAV vector capsids peptides in its surface, which can induce a CD8 T-cell response and the destruction of transduced hepatocytes, clinically recognized by an increase in liver transaminase (alanine aminotransferase [ALT]) and/or decreasing the transgene expression.

In 2011, the first successful liver-directed AAV vector-based gene therapy for hemophilia B was reported, with two critical strategies for AAV immunogenicity management.\(^\text{49–50}\) The first was the exclusion of patients with anti-AAV NAbs. In addition, immunosuppression with corticosteroids was used in response to ALT elevation to control AAV capsid-mediated cellular immune response.

AAV vector-mediated gene therapy trials for hemophilia A and B have shown promising results. Some participants have presented meaningful expression of FVIII\(^\text{51–56}\) or FIX,\(^\text{57–63}\) shifting their phenotype from severe to mild or even achieving normal factor levels after a single vector injection. Table 3 shows the ongoing hemophilia A and B gene therapy clinical trials.

Despite the encouraging results of the recent hemophilia gene therapy clinical trials, several issues remain unclear and unresolved. The variability in FVIII and FIX expression levels among the clinical trial participants and the unpredictable responses are observed in both hemophilia A\(^\text{52,53,54,55,64}\) and B gene therapy trials.\(^\text{59,60,61,62,63,65,66}\) Furthermore, the cellular immune response is AAV vector dose dependent, and the ideal immunosuppressive therapy still needs to be determined. In addition, safety concerns due to integration and potential malignancy risk will need long-term
follow-up.\textsuperscript{57} However, long-term durability seems to be another bold challenge, particularly for hemophilia A gene therapy.\textsuperscript{52,67}

After initial phase 1 and 2 trials confirming safety and exploring efficacy outcomes, more extensive phase 3 clinical trials for hemophilia A and B have been conducted in many centers, even outside North America and Europe. Centers in Australia, Japan, Brazil, South Africa, Turkey, Taiwan, Saudi Arabia, and other countries also include participants in hemophilia gene therapy clinical trials (Figure 2B). This widespread participation of hemophilia centers worldwide may help achieve the target patient number faster, despite the difficulties associated with AAV-seroprevalence limitations. In addition, although it is early to predict, the involvement of several centers around the world in prelicensed clinical trials may help in the future to establish and improve infrastructure and knowledge for other potential market sites, increasing access to this possible therapeutic alternative.

Also, local gene therapy programs for hemophilia have been carried out in Japan, China, and India.\textsuperscript{68} Another engaging initiative has been organized by St. Jude Children's Research Hospital, a phase 2 feasibility trial of AAV-mediated hemophilia B gene therapy in low- and middle-income countries. This program has two stages. In stage 1, vector infusion will happen at St. Jude, in Memphis, Tennessee, and patients will be monitored at their local sites. In stage 2, infusion and monitoring will happen at the low and middle-income countries’ sites. This initiative expects to prove the principle of the feasibility to perform and give access to gene therapy for patients in resource-constrained settings.\textsuperscript{58,69}

4 | GLOBAL DISTRIBUTION OF PROCOAGULANT PRODUCTS

Despite the significant advances in hemophilia care in the last decades, access to adequate diagnosis and treatment is still considerably unbalanced.

FVIII usage per capita is a critical parameter that has been used to estimate access to clotting factor products for years. Although the consumption of new nonfactor products will require consideration when assessing this parameter, it remains a helpful tool to evaluate the local availability of hemophilia treatment. It is suggested that 1 IU of FVIII per capita is the minimum amount of FVIII concentrates to guarantee the long-term survival of people with hemophilia.\textsuperscript{70} On the other hand, according to the European consensus, the minimum to provide the standard-of-care treatment is 4 IU of FVIII, and 0.5 IU of FIX concentrates per capita.\textsuperscript{71} According to
TABLE 3 Ongoing AAV-base gene therapy clinical trials for hemophilia A and B

| Program (Sponsor) | Product | Dose (vg/kg) | Status | Reference |
|-------------------|---------|--------------|--------|-----------|
| **Hemophilia A trials** | | | | |
| BMN 270, GENE8-1 (Biomarin) | valoctocogene roxaparvec rAAV5-BDDFVIII | $6 \times 10^{13}$ | Phase 1/2: active ($n = 15$) | 51-54,67 |
| | | $6 \times 10^{13}$ | Phase 3: active ($n = 134$) | |
| SB-525, ALTA and AFFINE (Sangamo, Pfizer) | PF-07055480 giroctocogene fitelparvec rAAV2/6-hFVIII | $9 \times 10^{11}$ | Phase 1/2: active ($n = 11$) | 63,104 |
| | | $2 \times 10^{12}$ | Phase 3: recruiting | |
| GO8 (UCL) | AAV2/8-HLP-FVIII-V3 | $6 \times 10^{11}$ | Phase 1: recruiting | - |
| | | $2 \times 10^{12}$ | | |
| | | $6 \times 10^{12}$ | | |
| SPK8011 (Spark) | rAAV-SPK200-BDDFVIIIco | $5 \times 10^{11}$ | Phase 1/2: active ($n = 18$) | 55 |
| | | $1 \times 10^{11}$ | | |
| | | $1.5 \times 10^{12}$ | | |
| | | $2 \times 10^{12}$ | | |
| BAY 2599023, DTX201 (Bayer, Ultragenyx) | BAY 2599023 rAAVhu37-hFVIIIco | $5 \times 10^{12}$ | Phase 1/2: recruiting ($n = 8$) | 59 |
| | | $1 \times 10^{13}$ | | |
| | | $2 \times 10^{13}$ | | |
| GS001 (Institute of Hematology & Blood Diseases Hospital, China) | GS001 | $2 \times 10^{12}$ | Phase 1/2: recruiting | |
| | | $6 \times 10^{12}$ | | |
| | | $2 \times 10^{13}$ | | |

| **Hemophilia B trials** | | | | |
| BENEGENE-2 (Spark, Pfizer) | PF-06838435 fidanacogene elaparvec rAAV-SPK100-hFIX-Padua | $5 \times 10^{11}$ | Phase 2: active ($n = 15$) | 51 |
| | | | Phase 3: recruiting | |
| AMT 061, HOPE-B (uniQure) | etranacogene dezaparvec AAV5-Padua hFIX | $2 \times 10^{13}$ | Phase 2b: active ($n = 3$) | 61,62 |
| | | | Phase 3: active ($n = 54$) | |
| FLT-180a, B-AMAZE (UCL and Freeline) | verbrinacogene setparvec AAV2/S3-FRE1-Ti-FIXco1 | $3.84 \times 10^{11}$ | Phase 1/2: active ($n = 10$) | 105 |
| | | $6.4 \times 10^{11}$ | | |
| | | $8.32 \times 10^{11}$ | | |
| | | $1.28 \times 10^{12}$ | | |
| | | $7.7 \times 10^{12}$ | | |

Note: For trials with more than one dosing scheme, doses considered therapeutic are in bold.
Abbreviations: AAV, adeno-associated virus; UCL, University College London; vg/kg, vector genomes per kilogram of body weight.

*B-AMAZE study results suggested that the dose of $7.7 \times 10^{11}$ vg/kg was optimal, after evaluation of the four doses as mentioned.

the World Federation of Haemophilia (WFH) annual global survey, in 2020. 78% of FVIII products were consumed in countries from the Americas and Europe, which encompass 31% of the world’s hemophilia population. This disproportion is even more pronounced if we consider the gross national income, based upon World Bank economic ratings. Only 12.3% of the FVIII consumed in 2020 was destined to treat 63% of the world’s hemophilia population living in low- and lower-middle-income countries.72

4.1 | Novel therapies around the globe

Nowadays, the use of EHL products is increasing. In Ireland, this is the only FVIII and FIX concentrates used.72 The latest WFH Annual Global Survey reported the current scenario for hemophilia therapies, including EHL factor concentrates. In 2020, 23 countries purchased EHL-rFVIII concentrates for their patients, using 1 604 990 663 units of this product. EHL-rFIX products are available in 18 countries, with a total consumption of 565 695 806 units in 2020. The Humanitarian Aid Program has distributed 40 673 500 units of EHL FVIII and FIX products to 37 and 18 countries, respectively.72 Recently, Krumb et al73 published data on the adoption of emicizumab in Europe. Of the 144 contacted hemophilia treatment centers, 46 responded to the survey, representing 21 countries. Emicizumab data were available on 43 centers. This agent was available for people with inhibitors in all of them, but approval for people without inhibitors was more restricted (in 37 centers, 88.1%).

In the WFH 2020 Annual Global Survey, 62 countries reported that emicizumab was available for people with or without inhibitors.72 According to the manufacturer, emicizumab is now licensed for prophylaxis in people with hemophilia A with inhibitors in >100 countries and for people without inhibitors in >80 countries. As a result, >10 000 people worldwide are on prophylaxis with emicizumab.74 In countries where emicizumab is approved for people both with and without inhibitors, it has been widely prescribed and is now one of the leading agents in use. Few agents have been incorporated
into clinical practice at such a fast pace. According to Hermans and Makris, 25% to 35% of people with hemophilia A without inhibitors are under prophylaxis with emicizumab in Israel, the United Kingdom, and Belgium. In a real-world cost estimate in the United States, emicizumab seems to be economically favorable compared to other agents for prophylaxis in people with or without inhibitors.

Few data are available regarding the use of new therapies in Latin America. In Brazil, until 2021, there were no EHL products distributed through the Inherited Coagulopathies Program from the Ministry of Health. Prophylaxis with emicizumab has just become available for people with hemophilia A with inhibitors who failed the immune tolerance induction (ITI) protocol. Chile has a similar scenario, with no EHL products and emicizumab provided for people with inhibitors who failed ITI (Dr Verónica Soto, personal communication, June 2021). Both Argentina and Colombia have a few people on EHL products, and emicizumab is available for people with inhibitors and selected people without inhibitors (Drs Daniela Neme and Adriana Linares, personal communication, June 2021). In Mexico, EHL factor concentrates are expected to be available by 2022, and emicizumab is provided for people with inhibitors, particularly those who failed ITI (Dr Jaime García Chavez, personal communication, June 2021). According to the 2020 WFH Annual Global Survey, Ecuador has some people on EHL products; Bolivia and Venezuela are participants of the Humanitarian Aid Program, and Uruguay has only SHL factor concentrates.

4.2 WFH Humanitarian Aid Program

In 1996, WFH started the Humanitarian Aid Program, an important initiative to provide access to treatment for people with inherited bleeding disorders in resource-constrained countries. Nevertheless, until 2014, the donations of procoagulant products were limited and sporadic, and its use was restricted to emergencies since there were not enough factor concentrates to provide sustained on-demand treatment.

However, in 2014, pharmaceutical companies Sanofi (formerly Biogen and Bioleraverit) and SoBi announced the donation of 1 billion IU of EHL FVIII and FIX concentrates over 10 years. As a result of this expressive increase in donated products through the WFH Humanitarian Aid Program, it was possible to expand the goals and number of people and countries that benefit from this initiative. Since 2016, donated EHL products have provided low-dose prophylaxis for people from resource-constrained countries. In 2016, the number of people on prophylaxis with donated products rose from 0 to 852, with 458 people <10 years old. In 2020, the cumulative number of people on prophylaxis with donated products was 1804 (including 1145 people <10 years old). The Humanitarian Aid Program helped expand the use of EHL products worldwide for prophylaxis and surgical procedures.

Several pharmaceutical companies continue to help increase access to treatment through the Humanitarian Aid Program. Despite the challenges and logistical restrictions with the COVID-19 pandemic, the program was not interrupted and was crucial to guarantee access to treatment for people in 69 countries in 2020.

In 2019, Roche also became a contributor to the program, and in 2020, started to donate emicizumab. According to Dr Assad Haffar, the director of the Humanitarian Aid Program, in June 2021, 633 people with hemophilia A from 26 countries were on prophylaxis with donated emicizumab, including 356 (56.2%) people <12 years old. Among these patients on emicizumab, 227 (35.9%) are people with hemophilia A with inhibitors. The remaining 406 (64.1%) people with hemophilia A who do not have inhibitors but met the criteria to receive prophylaxis with emicizumab were frequent bleeders (annualized bleeding rate [ABR] ≥6) or history of life-threatening bleeding episodes (Dr Assad Haffar, personal communication, June 2021).

5 ISTH 2021 VIRTUAL CONGRESS

5.1 Nonreplacement therapy

Almost 4 years after its first approval, real-world data elucidated emicizumab’s performance outside controlled clinical trials. The ISTH 2021 virtual congress had reports from multiple cohorts, including people with and without inhibitors, with efficacy and safety results comparable to those from the original trials.

Alternative dosing regimens have also been proposed. Fischer et al reported the experience of prescribing only entire emicizumab vials for maintenance in people with severe hemophilia A, with adjusted intervals between doses. They reported that 79% of people had zero bleeds during follow-up.

Also, in the Netherlands, Bukkems et al reported an alternative simulated dosing regimen, with a target median emicizumab plasma concentration between 40 and 60 µg/mL at steady state. The authors reported that costs were saved in up to 60% of the virtual population and could reach a median of almost 60 000 euros.

Another interesting report of a small cohort (n = 3) with a reduced-dose regimen for emicizumab from Malaysia has shown satisfactory clinical outcomes. With a mean dose of 1.8 mg/kg every 4 weeks, only one bleeding event was reported after up to 132 weeks of follow-up. Patients reported improved quality of life and could dismiss walking aids.

Regarding the rebalancing agents, updates on efficacy and safety for concizumab and marstacimab were presented at the ISTH 2021 virtual congress.

Astermark et al presented data from concizumab in people with hemophilia A/B with inhibitors (explorer 4) and hemophilia A without inhibitors (explorer 5). Subjects in the explorer4 trial were randomly assigned to either on-demand recombinant activated factor VII followed by concizumab or straight to concizumab throughout the whole study period. A total of 61 subjects were enrolled, and 51 had completed the extension period when the data were presented. As for efficacy, the ABR was 6.4 for hemophilia A without inhibitors,
3.8 for hemophilia A with inhibitors, and 6.2 for hemophilia B with inhibitors. Joint ABR was 5.2, 2.7, and 3.8 for each group, respectively. Interestingly, 15 of the 61 subjects enrolled developed concizumab ADAs. ADAs were transient and presented with a low titer in most cases, with no clinical impact.

An open-label study with marstacimab prophylaxis was presented by Mahlangu et al.41 Twenty subjects with hemophilia A or B, with or without inhibitors, were enrolled. No ADAs or serious adverse events related to marstacimab were reported. As for efficacy, when comparing data from before and after prophylaxis with marstacimab, the ABR decreased from 20.2 to 1.5 in the higher-dose (300 mg weekly) arm, and 17.4 to 2.7 in the lower-dose (150 mg weekly) group.

5.2  |  Gene therapy

5.2.1  |  Gene therapy for hemophilia A

Results from the phase 1/2 trial with valoctocogene roxaparvovec (AAV5·hFVIII·SQ) were updated after participants completed a 5-year follow-up period. There was a sustained reduction in ABR for both dose cohorts (4 × 10^{13} vector genomes per kilogram of body weight [vg/kg] and 6 × 10^{13} vg/kg). All participants remained off FVIII prophylaxis 5 years after infusion. Mean FVIII activities measured by chromogenic assay were 5.6 and 11.6 IU/dL for the lower- and higher-vector dose, respectively.51

The GENEr8-1 phase 3 trial reported results after 134 people with severe hemophilia A were dosed with valoctocogene roxaparvovec and followed up for at least 52 weeks. All participants received a single infusion of 6 × 10^{13} vg/kg. At weeks 49 to 52, the mean FVIII activity for 132 participants was 42.9 IU/dL (chromogenic assay). Among 17 participants with a minimum follow-up of 2 years, the mean FVIII activity was 24.4 IU/dL at week 104. Regarding clinical outcomes, 80% of the participants reported zero treated bleeds 4 weeks after infusion. The ABR decreased by 83.8% from baseline after week 4, with statistical superiority over FVIII concentrate prophylaxis. ALT elevation was reported in 115 participants (85.8%), and 106 (79.1%) received corticosteroids due to this ALT elevation. The average duration of corticosteroid use was 33 months, and three serious adverse events related to corticosteroid use were reported.53

George et al.,55 shared updated results on a phase 1/2 trial for SPK-8011, after 18 participants with hemophilia A were treated with four different dosing protocols (5 × 10^{11} vg/kg, 1 × 10^{12} vg/kg, 2 × 10^{12} vg/kg, and 1.5 × 10^{12} vg/kg). Only 17 participants had completed 1 year of follow-up. Of this group, two participants completely lost the FVIII expression. The remaining 15 participants had a mean FVIII activity of 11 ± 6.8% of the normal value (one-stage FVIII assay) and had a 91.2% reduction in ABR. Transaminitis was also a significant adverse event in this trial. Corticosteroids were used prophylactically by 5 participants and on-demand by 10 participants.

Results from the phase 1/2 trial with BAY 2599023 (AAVhu37·hFVIIIco) were also presented. So far, eight people were infused with three different dosing schemes (0.5 × 10^{13} vg/kg, 1 × 10^{13} vg/kg, and 2 × 10^{13} vg/kg). Investigators have stated that BAY 2599023 delivered protective FVIII levels sustained over time. Follow-up duration ranged from 12 to 100 weeks. ALT elevation was observed in four people, all managed with corticosteroids. Two additional patients from the higher-dose cohort were prescribed prophylactic corticosteroids.59

5.2.2  |  Gene therapy for hemophilia B

Gene therapy for hemophilia B was also addressed at the ISTH 2021 virtual congress. UniQure reported results for three clinical trials. The first was a phase 1/2 trial for AMT-060 (AAV5·hFIX). Ten patients were equally divided between two different dosing cohorts (5 × 10^{12} vg/kg and 2 × 10^{13} vg/kg) and were followed for 5 years. Mean FIX activity was 5.2% in the lower-dose cohort and 7.2% in the second group. ABR and FIX consumption declined compared to baseline, and all participants remained prophylaxis free. No major safety concerns were observed, with no sustained liver enzyme elevation. An enhanced construct, with FIX Padua as the therapeutic transgene, was later developed and is now the product used in the ongoing phase 3 trial.60

Following AMT-060, the updates for a phase 2b trial with etranacogene dezaparvovec (AAV5·Padua hFIX, AMT-061) were presented, following completion of a 30-month evaluation period. Unlike other gene therapy trials, preexisting anti-AAV5 neutralizing antibodies were assessed but did not represent an exclusion criterion for this study. Mean FIX activity was 44.2% (range, 36%-52%) 2 years after infusion, and no relationship between response and anti-AAV5 neutralizing antibody status was observed.61

HOPE-B is a phase 3 study that evaluates the efficacy and safety of etranacogene dezaparvovec (AAV5·Padua hFIX, AMT-061). So far, 54 patients have been dosed and completed a minimum follow-up of 6 months. Mean FIX activity was 41.3 IU/dL at 26 weeks in participants without anti-AAV5 neutralizing antibodies (n = 31) and 32.7 IU/dL in those with positive anti-AAV5 neutralizing antibodies (n = 23). No significant correlation was found between anti-AAV5 neutralizing antibody titer and FIX activity, bleeding phenotype, or safety.62 Investigators also shared some data on the HOPE-B cohort participant diagnosed with hepatocellular carcinoma over a year after AMT-061 infusion. Tumor sample analysis showed that AAV integration was very infrequent, and after an extensive analysis, hepatocellular carcinoma occurrence was deemed unrelated to gene therapy.63

6  |  FUTURE DIRECTIONS

The following years promise to continue the revolution in the hemophilia care scenario, as many new therapies are under development or ongoing preclinical and early-phase clinical trials.
SIG-001 consists of 1.5 mm alginate spheres encapsulating genetically modified allogeneic cells engineered to express human FVIII. The capsules shield the cells from the host’s immune system, and the conjugation to alginate biomaterial avoids pericapsular fibrotic overgrowth. In preclinical trials, SIG-001 could produce functional FVIII in a dose-dependent manner and correct the bleeding phenotype of hemophilia A mice. After no concerns regarding safety or toxicology were observed in mice and nonhuman primates, a phase 1/2 clinical trial was announced. SIG-001 would be administered in the peritoneal cavity through laparoscopy in 18 patients. However, on July 9, 2021, the company announced that the study was put on clinical hold by the US Food and Drug Administration after a participant developed inhibitors to FVIII. Three participants had been dosed so far.

Currently, there are preclinical studies on subcutaneous agents for hemophilia care. The first is FVIII-ABD, a subcutaneous FVIII that has shown good availability (ranging from 15.3% to almost 50%, depending on the animal model studied). Another strategy is the coadministration of recombinant FVIII and recombinant VWF fragments containing the D3 domain (VWF-12 and VWF-13). This strategy resulted in a good availability (up to 18.5%) for subcutaneous human recombinant FVIII, with slow absorption and prolonged half-life.

As previously discussed, AAV vectors are widely used in gene transfer clinical trials for hemophilia A and B. Among its many benefits, AAV vectors are replication defective and target many different cells and tissues. Nevertheless, the unpredictable postinfusion response and the slow but progressive loss of expression over the years are still issues to be addressed. In this scenario, lentiviral vectors emerge as interesting candidates for gene transfer with long-term expression. Since lentivirus integrates into the host’s cell chromatin, its genome is maintained after each cellular duplication. Preclinical studies have shown multiyear transgene liver expression in mice and dogs with hemophilia A. More recently, some improvements in lentiviral vectors have been studied. Modifications in the vector surface to decrease T-cell–mediated immunogenicity and increased resistance to phagocytosis have led to higher FIX expression, reaching values up to 300% of normal.

Beyond viral vectors, lipid nanoparticles are also being studied for gene transfer. Chen et al. have described the use of FVIII-encoding mRNA, packaged into liver-directed lipid nanoparticles. After a single intravenous injection, hemophilia A mice have presented with a variable range of FVIII activity and maintained therapeutic FVIII levels up to 5 to 7 days after infusion.

**7 | SUMMARY**

Remarkable improvement in hemophilia care was achieved during the past decades. The availability of safe and effective clotting factor concentrates was crucial for prophylaxis feasibility, starting at a young age for people with severe hemophilia. New bioengineered clotting factors, such as EHL products, helped ameliorate the burden of frequent intravenous administration. However, the risk for inhibitors continues to be the major complication of hemophilia replacement therapy.

More recently, the development of nonreplacement therapies represents a unique alternative in the effective prevention of bleeding, regardless of the presence of inhibitors. Currently, emicizumab has become the preferable option for prophylaxis for people with hemophilia A with inhibitors, with several advantages for people with noninhibitor hemophilia A. These new therapies have also been used in resource-constrained countries through the WFH Humanitarian Aid Program and contributed to increasing the chance of elective surgeries and the number of people on prophylaxis in low-income countries.

In addition, gene therapy clinical trials for hemophilia have shown promising results, and this modality of treatment may become an attractive alternative for hemophilia management, even in resource-constrained countries.

Thus, in the near future, one of the recurrent challenges for hemophilia management will be to define the most appropriate treatment according to the needs of each person and local treatment availability.

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**AUTHOR CONTRIBUTIONS**

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