




Management of a High-Risk Surgery with Emicizumab and Factor VIII in a Child with a Severe Hemophilia A and Inhibitor

Charles R. Lefèvre^{1,2,3}  Anaïs Jaffré⁴ Adeline Pontis¹ Fabienne Nedelec-Gac^{1,2} Pierre Guéret^{1,2}
Isabelle Gouin-Thibault¹ Bernard Fraisse⁵ Sophie Bayart² Benoit Guillet^{2,3}

¹ Laboratoire d'Hémostase Bioclinique, CHU de Rennes, Bretagne, France

² Centre Régional de Traitement de l'Hémophilie, CHU de Rennes, Bretagne, France

³ Univ Rennes, CHU Rennes, Inserm, EHESP, Irset (Institut de recherche en santé, environnement et travail) - UMR_S 1085, Rennes, France

⁴ Service de Réanimation Pédiatrique, CHU de Rennes, Bretagne, France

⁵ Service de Chirurgie Pédiatrique, CHU de Rennes, Bretagne, France

Address for correspondence Benoit Guillet, MD, PhD, Centre Régional de Traitement de l'Hémophilie – CHU de Rennes, Bretagne, France (e-mail: Benoit.GUILLET@chu-rennes.fr).

TH Open 2021;00:e163–e165.

The recent development of a humanized, bi-specific, and monoclonal antibody mimicking the function of activated factor VIII was a revolution in the management of patients suffering from severe hemophilia A with inhibitors.¹ The phase III randomized studies have shown a more efficient prophylaxis of this subcutaneous administered drug in these patients compared with recombinant FVIIa (rFVIIa) and activated prothrombin complex concentrates (aPCC).^{2,3} Nonetheless, there are “real life” matters that need to be explored in this new era of managing hemophilia patients, such as surgery management under emicizumab, especially in children.^{4–6} Here, we report the first case, to our knowledge, of major orthopedic surgery managed with factor VIII infusions in a child with inhibitor receiving emicizumab.

Our patient is a 9-year-old boy with severe hemophilia A. He presented an inhibitor after the 10th exposed day with a factor VIII (FVIII) concentrate (historic peak: 412 BU/mL). Thereafter, he presented an important intracranial hemorrhage (subdural and intraparenchymal hematomas). The first treatment with rFVIIa (Novoseven, 90 µg/kg/2 h) and tranexamic acid was inefficient. It was therefore switched with aPCC (FEIBA 80 IU/kg/8 h) that was rapidly effective. He also presented some knees and left hip hemarthrosis responsible for arthropathies. Several attempts of immune tolerance induction through Port-a-Cath failed whatever the FVIII concentrate used, recombinant FVIII or plasma-derived FVIII. Because of an insufficient efficacy

of rFVIIa for totally stopping some bleeding events, a prophylaxis with aPCC was performed with daily infusion of 80 IU/kg. Given the cumbersome nature of this treatment, emicizumab (Hemlibra) was introduced instead of aPCC in December 2018. Emicizumab was started with 3 mg/kg/week for 4 weeks and then adapted to 1.5 mg/kg/week. The boy had no more bleeding including clinically evident hemarthrosis. However, the left hip arthropathy worsened with aggravation of limping and pain leading to the use of a wheelchair. Hip X-ray showed an osteonecrosis of the left femoral head requiring a femoral varization osteotomy that was planned in June 2020. Because the inhibitor titer was low at 2 BU/mL, rFVIII-Fc (Elocta) was administered to normalize the coagulation during the surgery. A rFVIII-Fc bolus with 6,000 IU (150 IU/kg) was initially administered, followed by a continuous infusion 12 IU/kg/h (→ **Fig. 1A**). The emicizumab treatment was continued (1.5 mg/kg/week).

This complex surgery performed on rFVIII-Fc required a close monitoring with frequent FVIII:C measures to adjust rFVIII-Fc administrations. All FVIII:C levels were measured with a chromogenic method (STA-R Max3 analyzer ; Trini-CHROM FVIII:C reagent – Stago) by using bovine coagulation factors X and IXa that do not interfere with emicizumab.^{7–9} In same plasma samples, thrombin generation assays (TGA) were performed by using low concentration of tissue factor (1 pM) as suggested for the TGA in hemophilia A patients.¹⁰

received

November 15, 2020

accepted after revision

January 25, 2021

DOI <https://doi.org/>

10.1055/s-0041-1728667.

ISSN 2512-9465.

© 2021. The Author(s).

This is an open access article published by Thieme under the terms of the Creative Commons Attribution License, permitting unrestricted use, distribution, and reproduction so long as the original work is properly cited. (<https://creativecommons.org/licenses/by/4.0/>)

Georg Thieme Verlag KG, Rüdigerstraße 14, 70469 Stuttgart, Germany

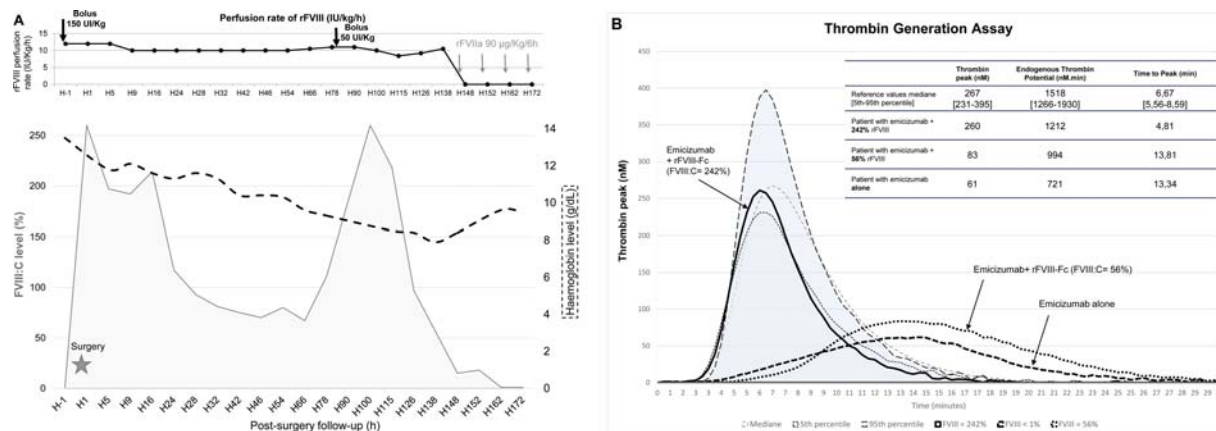


Fig. 1 (A) Evolution of FVIII:C and haemoglobin levels. (B) Evolution of thrombin generation assays. Reference values were established in 32 healthy patients.

TGA were performed before the surgery without rFVIII-Fc and throughout rFVIII-Fc infusions (►Fig. 1B).

The FVIII:C level measured 30 minutes after bolus was high at 242% and maintained above 200% during the first 24 hours, following postsurgery without change of the rFVIII-Fc infusion rate (►Fig. 1A). The orthopedic surgery included a proximal femoral derotation osteotomy and then fixed by osteosynthesis material (►Fig. 2). No bleeding (or other complication) occurred during and after surgery with low per-operative blood loss (200 mL). His immediate postsurgery hemoglobin level was 11.8 g/dL. After the first postsurgery day, the FVIII:C levels progressively decreased until 100% at H28. At the 78th hour, a 50 IU/kg rFVIII-Fc bolus was performed for drains ablation. After a new peak, the FVIII:C level further decreased despite maintaining the continuous infusion at the dose of 10 IU/kg/h because of the inhibitor resurgence at H162 with a titer of 1.8 BU/mL which rose to 33 BU/mL at H200. TGA's parameters evolved within normal ranges during first postsurgical 24 hours and then progressively decreased together with FVIII:C levels (►Fig. 1B). The 6th postsurgery day, rFVIII-Fc was replaced with rFVIIa boluses at 90 µg/kg that were stopped

after 1 day. During his stay, the patient did not bleed and therefore did not require any red blood cell concentrate transfusion; the lowest hemoglobin he presented was 7.9 g/dL and was only corrected with a single iron sucrose injection (Venofer). Furthermore, no markers of coagulation activation were detected during rFVIII-Fc and rFVIIa treatments. Always receiving a prophylaxis with emicizumab alone, the patient was able to walk within 2 months after surgery and presented no bleeding.

To date, most of the surgery reports performed with emicizumab in cases with inhibitor, concerned adults. rFVIIa was most used to prevent bleeding during and after surgical procedures. In pediatrics, only minor procedures were described.^{4-6,11-13} Our case is so, to our knowledge, the first description of a major orthopaedic surgery managed with FVIII concentrates in a child with severe hemophilia A with inhibitor while receiving a prophylaxis with emicizumab. The choice we made of using rFVIII-Fc for bleeding prevention during and after the surgery was foremost driven by the presence of a low titer inhibitor. Furthermore, we prohibited aPCC because of its thrombotic risk in association with emicizumab.¹⁴ Finally,

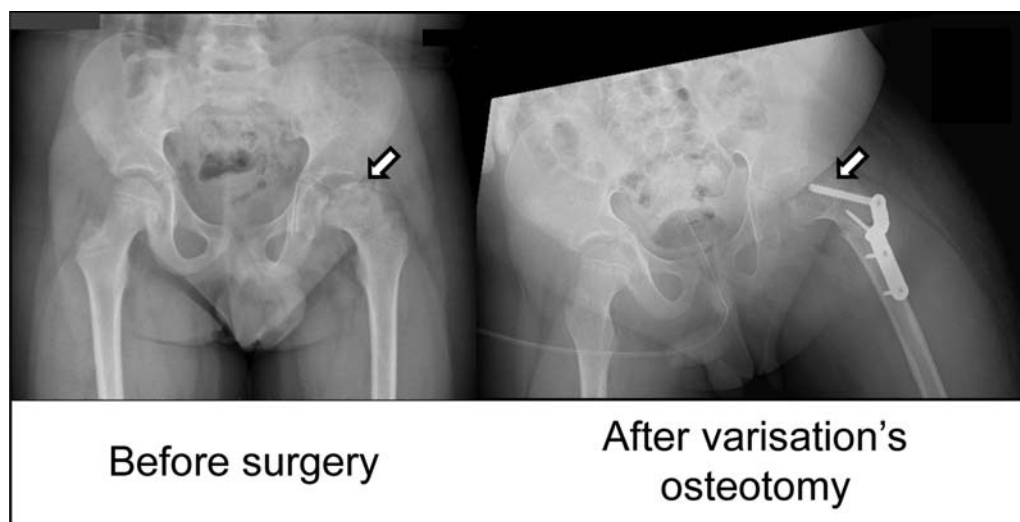


Fig. 2 The arrow on the left shows the severe arthropathy of the left hip with a coxa plana and an alteration of cartilageneous and osseous structures. The arrow on the right shows the osteosynthesis material inserted after the varisation osteotomy of the femoral neck.

rFVIIa was ruled out because it was often only partially effective for this patient. We show here that, as for some adults reported to date, major surgeries can be safely managed with FVIII concentrates in children with severe hemophilia A with inhibitor while receiving emicizumab. This dual treatment could also be applicable without an inhibitor at the doses usually administered.¹⁵ In our patient, the FVIII:C chromogenic assay appeared to be a reliable tool for peri-surgical monitoring in children despite concomitant treatment with emicizumab as it was proposed.^{8,9} TGA with low TF concentration performed in parallel to FVIII:C measures could be helpful.⁹ However, our results did not show a perfect correlation between TGA's parameters and normalized FVIII levels. The TGA with these conditions is so not enough reliable for the monitoring of emicizumab treatments and need further adjustments. Finally, it was rapidly observed that maintaining the prophylaxis with emicizumab reduces the duration of post-surgical treatment in major orthopaedic surgeries.¹³ Indeed, for our patient, as already reported for adults receiving emicizumab, FVIII or rFVIIa injections were only necessary until the 7th day post-surgery. Thereafter, prevention with emicizumab alone was sufficient to protect against late bleeding during rehabilitation.

Authors' Contributions

All authors made substantial contributions to the conception or design of the work, the acquisition, analysis, interpretation of data, as well as drafting or revising the work critically and giving final approval of the version to be published.

Funding

None.

Conflict of Interest

None declared.

Acknowledgments

Authors would like to thank Mrs. Mélanie Delanoe for her technical support.

References

- Shima M, Hanabusa H, Taki M, et al. Factor VIII-mimetic function of humanized bispecific antibody in hemophilia A. *N Engl J Med* 2016;374(21):2044–2053
- Oldenburg J, Mahlangu JN, Kim B, et al. Emicizumab prophylaxis in hemophilia A with inhibitors. *N Engl J Med* 2017;377(09):809–818
- Young G, Liesner R, Chang T, et al. A multicenter, open-label phase 3 study of emicizumab prophylaxis in children with hemophilia A with inhibitors. *Blood* 2019;134(24):2127–2138
- Barg AA, Livnat T, Budnik I, et al. Emicizumab treatment and monitoring in a paediatric cohort: real-world data. *Br J Haematol* 2020;191(02):282–290
- Seaman CD, Ragni MV. Emicizumab use in major orthopedic surgery. *Blood Adv* 2019;3(11):1722–1724
- Evans MS, Davis C, Eyster ME. Total knee replacement with and without emicizumab: a unique comparison of perioperative management. *Blood Adv* 2020;4(05):855–857
- Nougier C, Jeanpierre E, Ternisien C, et al; With contributions from the Working Group on Biology of Haemorrhagic Disorders (BIMHO, Under the Aegis of GFHT) Emicizumab treatment: impact on coagulation tests and biological monitoring of haemostasis according to clinical situations (BIMHO group proposals). *Eur J Haematol* 2020;105(06):675–681
- Bowyer AE, Lowe AE, Tiefenbacher S. Laboratory issues in gene therapy and emicizumab. *Haemophilia* 2020
- Unterberger M, Wagner L, Binder NB. Use of chromogenic Factor VIII activity determination in Haemophilia A plasma of patients under emicizumab treatment. :1. Accessed 2019 at: https://www.technoclone.com/sites/default/files/pb1299_technochrom_fviii_c_emicizumab_isth2019.pdf
- Dargaud Y, Béguin S, Lienhart A, et al. Evaluation of thrombin generating capacity in plasma from patients with haemophilia A and B. *Thromb Haemost* 2005;93(03):475–480
- Kizilcak H, Yukhtman CL, Marquez-Casas E, Lee J, Donkin J, Young G. Management of perioperative hemostasis in a severe hemophilia A patient with inhibitors on emicizumab using global hemostasis assays. *Ther Adv Hematol* 2019;10:2040620719860025
- Santagostino E, Mancuso ME, Novembrino C, Solimeno LP, Tripodi A, Peyvandi F. Rescue factor VIII replacement to secure hemostasis in a patient with hemophilia A and inhibitors on emicizumab prophylaxis undergoing hip replacement. *Haematologica* 2019;104(08):e380–e382
- Biron-Andreani C, Diaz-Cau I, Ranc A, et al. Major surgery management in patients with haemophilia A and inhibitors on emicizumab prophylaxis without global coagulation monitoring. *Br J Haematol* 2020;189(03):e100–e103
- Makris M, Iorio A, Lenting PJ. Emicizumab and thrombosis: The story so far. *J Thromb Haemost* 2019;17(08):1269–1272
- Hermans C, Apte S, Santagostino E. Invasive procedures in patients with haemophilia: review of low-dose protocols and experience with extended half-life FVIII and FIX concentrates and non-replacement therapies. *Haemophilia* 2020;hae.13978