

# An overview in acquired hemophilia A

Citation for published version (APA):

Pasca, S. (2023). An overview in acquired hemophilia A: a rare but complicated disease. [Doctoral Thesis, Maastricht University]. Maastricht University. https://doi.org/10.26481/dis.20231009sp

#### **Document status and date:**

Published: 01/01/2023

DOI:

10.26481/dis.20231009sp

#### **Document Version:**

Publisher's PDF, also known as Version of record

#### Please check the document version of this publication:

- A submitted manuscript is the version of the article upon submission and before peer-review. There can be important differences between the submitted version and the official published version of record. People interested in the research are advised to contact the author for the final version of the publication, or visit the DOI to the publisher's website.
- The final author version and the galley proof are versions of the publication after peer review.
- The final published version features the final layout of the paper including the volume, issue and page numbers.

Link to publication

#### General rights

Copyright and moral rights for the publications made accessible in the public portal are retained by the authors and/or other copyright owners and it is a condition of accessing publications that users recognise and abide by the legal requirements associated with these

- Users may download and print one copy of any publication from the public portal for the purpose of private study or research.
- You may not further distribute the material or use it for any profit-making activity or commercial gain
  You may freely distribute the URL identifying the publication in the public portal.

If the publication is distributed under the terms of Article 25fa of the Dutch Copyright Act, indicated by the "Taverne" license above, please follow below link for the End User Agreement:

www.umlib.nl/taverne-license

### Take down policy

If you believe that this document breaches copyright please contact us at:

repository@maastrichtuniversity.nl

providing details and we will investigate your claim.

Download date: 17 May. 2024

# Propositions belonging to this thesis

# An overview on acquired hemophilia A: a rare but complicated disease

# By Samantha Pasca

- 1. Acquired hemophilia A may occur in patients without congenital bleeding disorders, especially in older adults with some underlying comorbidities (This thesis)
- 2. The diagnosis of acquired hemophilia A is often delayed, due to the presence of confounding factors such as taking anticoagulants or other clinical conditions of the patients (This thesis)
- 3. Although acquired hemophilia is a rare disease, the costs, considered too high, of bypassing agents and susoctocog alfa sometimes make these drugs unavailable, especially in small hospitals, thus making timely treatment of acute bleeding impossible. (This thesis)
- 4. Plasma-derived concentrates still can represent a therapeutic option when other drugs are not available and/or in developing countries (This thesis)
- 5. Approximately one fifth of patients present a relapse within a month of resolution of the acute event, despite this and despite several reports having demonstrated the effectiveness of short or long-term prophylaxis with aPCC and now also with emicizumab, there are still no guidelines that recommend its use. (This thesis)
- 6. Accurate information and training of physicians is desirable to ensure the correct management of patients affected by acquired hemophilia A (Impact paragraph)
- 7. In recent years, the treatments of rare bleeding disorders have notably changed. Extended-half-life coagulation factor concentrates, subcutaneous non-substitutive drugs, and ultimately gene therapy have significantly improved the quality of life of patients (my discipline)
- 8. A multidisciplinary approach is needed to manage patients with rare bleeding disorders in the 21<sup>st</sup> century (my discipline)
- 9. New drugs and new therapeutic approaches require a continuous and constant updating of clinicians dealing with rare bleeding disorders (my discipline)
- 10. Diagnosing and managing patients with bleeding disorders' is a competency that requires higher-level cognitive processes and is dependent on one's ability to collect and understand the necessary knowledge, and then apply, analyze, and synthesize that knowledge for a specific patient within a particular context, and finally, evaluate this information for a given purpose (Khalife et al., J Thromb Haemost 2022)