

Coagulation factor V deficiency

Citation for published version (APA):

Nuzzo, F. (2016). *Coagulation factor V deficiency: from molecular diagnosis to molecular therapy*. Uitgeverij BOXPRESS. <https://doi.org/10.26481/dis.20160114fn>

Document status and date:

Published: 01/01/2016

DOI:

[10.26481/dis.20160114fn](https://doi.org/10.26481/dis.20160114fn)

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 rights.

- 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.

Stellingen

behorende bij het proefschrift

Coagulation factor V deficiency: From molecular diagnosis to molecular therapy

1. The current *F5* mutational spectrum reflects more the tools that are used to identify mutations than the actual distribution of mutation types (this thesis).
2. A synonymous mutation is not always neutral (this thesis).
3. The complexity of the splicing code offers many chances for mutations to strike (this thesis).
4. Factor V-deficient patients with homozygous splicing mutations have a particularly high incidence of intracranial haemorrhages (this thesis).
5. Pharmacological inhibition of tissue factor pathway inhibitor (TFPI) protects FV-deficient patients from severe bleeding (Duckers *et al.* Blood 2008).
6. There is no disease so rare that it does not deserve attention (Orphanet, the portal for rare diseases and orphan drugs).
7. Bovine thrombin is a universal and highly effective PCR enhancer (Zhang *et al.* BioTechniques 2014).
8. The personalized nature of mutation-specific antisense molecules is one of the main obstacles to their therapeutic development.
9. North and South are relative concepts.
10. Facts are not science as a dictionary is not literature (Martin H. Fischer).
11. Anything you try to fix will take longer and cost more than you thought (Bryson's Law of Repairs).