

## ADAMTS13: Biology And Disease

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ADAMTS13: Biology and Disease Sep 14, 2015. by George M. Rodgers. Hardcover. \$151.92 \$159.00. Available for Pre-order. This item will be released on September 14, 2015. [http://www.amazon.com/s?ie=UTF8&page=1&rh=n%3A283155%2Ck%3Aclinical%20hematology%2Cp\\_n\\_publication\\_date%3A1250228011](http://www.amazon.com/s?ie=UTF8&page=1&rh=n%3A283155%2Ck%3Aclinical%20hematology%2Cp_n_publication_date%3A1250228011)

The term thrombotic thrombocytopenic purpura-hemolytic uremic syndrome typically associated with severely deficient ADAMTS13 (See 'Evolution of disease <http://www.uptodate.com/contents/causes-of-thrombotic-thrombocytopenic-purpura-hemolytic-uremic-syndrome-in-adults>

Localization of ADAMTS13 to the stellate cells the deficiency of plasma ADAMTS13 activity in liver diseases may Hemostasis, Thrombosis, and Vascular Biology; <http://www.bloodjournal.org/content/106/3/922?sso-checked=true>

View Joshua Muia's professional profile on LinkedIn. The structural and functional studies of the Wilson disease-causing Most ADAMTS13 assays use non <https://www.linkedin.com/pub/joshua-muia/15/991/b0b>

Dec 01, 2010 This review focuses on the kidney disease caused by Shiga toxin and TTP is initiated by deficient ADAMTS13 all of the changes in biology and <http://www.ncbi.nlm.nih.gov/pmc/articles/PMC3032420/>

Von Willebrand disease (VWD) is the most common inherited bleeding disorder, affecting up to 1 percent of the population as assessed by random laboratory screening <http://www.uptodate.com/contents/clinical-presentation-and-diagnosis-of-von-willebrand-disease>

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von Willebrand factor biochemistry and cell biology; von Willebrand disease diagnostic criteria; Molecular genetic pathology of type 3 vWD; ADAMTS13 metalloprotease;

[http://hstalks.com/main/view\\_talk.php?t=61](http://hstalks.com/main/view_talk.php?t=61)

Von Willebrand factor (vWF) is a blood glycoprotein involved in hemostasis. It is deficient or defective in von Willebrand disease and is involved in a large number

[http://en.wikipedia.org/wiki/Von\\_Willebrand\\_factor](http://en.wikipedia.org/wiki/Von_Willebrand_factor)

ADAMTS13: biology and disease, chapter Garnier, A., Bienaime, F., Dragon-Durey, M.A., Ngo, S. et al. Genetics and outcome of atypical hemolytic uremic syndrome:

[http://www.tmreviews.com/article/S0887-7963\(14\)00070-4/fulltext](http://www.tmreviews.com/article/S0887-7963(14)00070-4/fulltext)

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<http://www.springer.com/life+sciences/cell+biology?SGWID=4-10032-66-653429-0&sba=EXCLUDE&originalID=315111&resultStart=41>

Nov 04, 2014 Most sporadic cases of TTP appear to be associated with severe deficiency of ADAMTS13 although TTP is thought to be a rare disease. Biology and

<http://emedicine.medscape.com/article/206598-overview>

Insights on ADAMTS proteases and ADAMTS-like proteins in an allergic airways disease model : Adamts13 / of their biology was the apparent

<http://www.sciencedirect.com/science/article/pii/S0945053X15000529>

ADAMTS-13 Biology and Disease. av George Comprehensive in its exploration of the ADAMTS13 protease in disease, ADAMTS13: Biology and Disease is a

<http://www.bokus.com/bok/9783319087160/adamts-13/>

New Concepts of Thrombotic Thrombocytopenic Purpura and a Strategy to Prevent its Relapse, Han-Mou Tsai Home In G. Rodgers (ed): ADAMTS13: Biology and Disease,

<http://www.esciencecentral.org/journals/new-concepts-of-thrombotic-thrombocytopenic-purpura-and-a-strategy-to-prevent-its-relapse-2329-8790.1000157.php?aid=31831>

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Some of the topics currently pursued by the laboratory include development of extra-embryonic tissues and vascular biology ed, ADAMTS13: Biology and Disease

<http://www.lerner.ccf.org/bme/apte/>

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Molecular mechanisms of chronic alcohol and obesity induced fatty liver disease : and biology of ADAMTS13 metalloprotease: ADAMTS13, von Willebrand factor,

<http://services.medicine.uab.edu/facultydirectory/FacultyListingType.asp?FacultyTypeID=BSB>

Books: Medical Sciences: Internal Medicine: Hematology. Results 1 - 100: George M. Rodgers (2015) ADAMTS13: Biology and Disease; Springer; 3319087169.

<http://www.researchbooks.org/Books/Medical-Sciences/Internal-Medicine/Hematology/>

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Leber's congenital amaurosis (LCA) is a group of inherited blinding diseases with onset during childhood. One form of the disease, LCA2, is caused by mutations in the

<http://www.nejm.org/doi/full/10.1056/NEJMoa0802315>

In the century since the discovery of the ABO blood groups, numerous associations between ABO groups and disease have been noted. However, the selection pressures

<http://www.bloodjournal.org/content/110/7/2250?sso-checked=true>

Nov 04, 2014 TTP can affect any organ system, but involvement of the peripheral blood, the central nervous system, and the kidneys causes the clinical manifestations.

<http://emedicine.medscape.com/article/206598-workup>