

ADAMTS13%0A

Download PDF Ebook and Read OnlineAdamts13%0A. Get **Adamts13%0A ADAMTS13 Wikipedia** ADAMTS13 (a disintegrin and metalloproteinase with a thrombospondin type 1 motif, member 13) also known as von Willebrand factor-cleaving protease (VWFCP) is a zinc-containing metalloprotease enzyme that cleaves von Willebrand factor (vWf), a large protein involved in blood clotting.

<http://www.thunder-girls.de/ADAMTS13-Wikipedia.pdf>

ADAMTS13 Activity with Reflex to Inhibitor Quest Diagnostics

ADAMTS-13 is a zinc metalloprotease that cleaves ultra large vWF multimers. Studies have shown that low levels of ADAMTS-13 activity are associated with thrombotic thrombocytopenic purpura (TTP), a life-threatening hematological condition characterized by a low platelet count, microvascular thrombi, red cell fragmentation, and renal complications.

http://www.thunder-girls.de/ADAMTS13_Activity_with_Reflex_to_Inhibitor-Quest_Diagnostics.pdf

OMIM Entry 604134 A DISINTEGRIN LIKE AND

Motto et al. (2005) generated Adamts13-deficient mice, which were viable and exhibited normal survival. Introduction of a genetic background from a mouse strain with elevated plasma vWF resulted in the appearance of spontaneous thrombocytopenia in a subset of Adamts13-deficient mice and significantly decreased survival.

http://www.thunder-girls.de/OMIM_Entry-__604134-A_DISINTEGRIN_LIKE_AND-_.pdf

Pathology Outlines Case of the week 403

5 October 2016 - Case of the Week #403 All cases are archived on our website. To view them sorted by case number, diagnosis or category, visit our main Case of the Week page.

http://www.thunder-girls.de/Pathology_Outlines-Case_of_the_week__403.pdf

Thrombotic thrombocytopenic purpura Wikipedia

Thrombotic thrombocytopenic purpura (TTP) is a blood disorder that results in blood clots forming in small blood vessels throughout the body. This results in a low platelet count, low red blood cells due to their breakdown, and often kidneys, heart, and brain dysfunction. Symptoms may include large bruises, fever, weakness, shortness of breath

http://www.thunder-girls.de/Thrombotic_thrombocytopenic_purpura-Wikipedia.pdf

ADAMTS13

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Trombotische trombocytopenische purpura Wikipedia

Trombotische trombocytopenische purpura (TTP) of ziekte van Moschcowitz is een zeldzame aandoening van de bloedstolling. In de meeste gevallen komt de ziekte voort uit een tekort aan of remming van een enzym, ADAMTS13, dat verantwoordelijk is voor het "knippen" van grote multimeren van de von willebrandfactor.

http://www.thunder-girls.de/Trombotische_trombocytopenische_purpura-Wikipedia.pdf

Blood Bank Guy's Glossary of Blood Banking Terms

The Blood Bank Glossary is a great place to find the definition of any weird term that you heard a blood banker use in a lecture (or everyday conversation!)

http://www.thunder-girls.de/Blood_Bank_Guy's_Glossary_of_Blood_Banking_Terms.pdf

P r p u r a t r o m b o c i t o p n i c a t r o m b t i c a W i k i p e d i a l a

La p r p u r a t r o m b o c i t o p n i c a t r o m b t i c a es una enfermedad hemorr gica microangiop tica poco frecuente. [1] El primer caso fue registrado en 1924 por Moschcowitz.

http://www.thunder-girls.de/P__rpura_trombocitop__nica_tromb__tica-Wikipedia__la-_.pdf

Practical Haemostasis com A Practical Guide and Resource

Practical-Haemostasis.com is designed to teach you laboratory haemostasis. Furthermore it provides a series of data-interpretation type questions [with answers] that you can work through.

http://www.thunder-girls.de/Practical_Haemostasis_com-A_Practical_Guide_and_Resource-_.pdf

UP Health System Marquette Laboratory Reference Lab

The UPHS Marquette Reference Lab is a fully integrated, comprehensive provider of laboratory services to inpatients, outpatients, outreach clients and other community health care organizations across Michigan's Upper Peninsula.

http://www.thunder-girls.de/UP_Health_System-__Marquette_Laboratory-Reference_Lab-_.pdf

OMIM Entry 613160 VON WILLEBRAND FACTOR VWF

The VWF gene encodes von Willebrand factor (VWF), a large multimeric glycoprotein that plays a central role in the blood coagulation system, serving both as a major mediator of platelet-vessel wall interaction and platelet adhesion, and as a carrier for coagulation factor VIII (F8; 300841).

http://www.thunder-girls.de/OMIM_Entry-__613160-VON_WILLEBRAND_FACTOR__VWF.pdf

Hematology Essentials A Foundation for Accurate Smear Reviews

Christine Hinz, MS, MLS(ASCP)CM Hematology Essentials: A Foundation for Accurate Smear Reviews

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Porpora trombotica trombocitopenica Wikipedia

La porpora trombotica trombocitopenica (PTT o sindrome di Moschcowitz) una malattia rara caratterizzata da anemia emolitica, piastrinopenia da consumo, sintomi neurologici. una anemia emolitica microangiopatica, Coombs negativa, associata a piastrinopenia con esordio acuto o subdolo e decorso spesso fulminante.

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(: Thrombocytopenia,) .

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Home Blood Journal

A weekly snapshot of the hottest studies from Blood, the official journal of the American Society of Hematology (ASH), hand-picked from each issue by the editors, Bob L wenberg, MD, PhD and Nancy Berliner, MD.

http://www.thunder-girls.de/Home-Blood_Journal.pdf

Fibrinogen Assays Practical Haemostasis com

Fibrinogen consists of three pairs of polypeptide chains: two A , two B and two . These are linked together by 29 disulphide bonds in such a way that N-terminal regions of the 6-polypeptide chains meet to form a central E-domain.

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Terminal Complement Inhibitor Eculizumab in Atypical

Atypical hemolytic uremic syndrome is a genetic, life-threatening, chronic disease of complement-mediated thrombotic microangiopathy. Plasma exchange or infusion may transiently maintain normal
http://www.thunder-girls.de/Terminal_Complement_Inhibitor_Eculizumab_in_Atypical-_.pdf

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