

Factor X Deficiency

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Recent Publications on Factor X Deficiency:



- [Menorrhagia and reproductive health in rare bleeding disorders: a study from the Indian subcontinent.](#)
*At this centre, 130 women with rare bleeding disorders (RBD) were...*23rd September, 2008
National Institute of Immunohaematology (ICMR), KEM Hospital, Parel,- Haemophilia. 2008 Sep 18. ([DOI Direct Link](#))
- [Phenotype and genotype report on homozygous and heterozygous patients with congenital factor X deficiency.](#)
*Factor X deficiency is a severe rare hemorrhagic condition inherited as an...*16th September, 2008
Hematology and Thrombosis Unit, Hematology Research Centre, Shiraz- Haematologica. 2008 Jun;93(6):934-8. Epub 2008 Apr 9. ([DOI Direct Link](#))
- [Different genotypes are responsible for the normal Russell viper venom assays seen in some cases of congenital factor X deficiency.](#)
*At this centre, 130 women with rare bleeding disorders (RBD) were...*13th September, 2008
University of Padua Medical School, Department of Medical and Surgical- Am J Hematol. 2008 Aug 21. ([DOI Direct Link](#))
- [Congenital factor X deficiencies with a defect only or predominantly in the extrinsic or in the intrinsic system: a critical evaluation.](#)
*Congenital Factor X deficiency is commonly classified as type I, in which...*9th September, 2008
Department of Medical and Surgical Sciences, University of Padua Medical- Am J Hematol. 2008 Aug;83(8):668-71. ([DOI Direct Link](#))
- [\[Secondary amyloidosis connected with neoplasma of urinary bladder and acquired disorders of blood coagulation. Case report\]](#)
*Amyloidosis is a general term encompassing a group of disorders...*30th August, 2008
Katedra i Klinika Nefrologii Akademii Medycznej w Lublinie im. prof.- Pol Arch Med Wewn. 2006 Sep;116(3):868-72.
- [Intracranial haemorrhage in patients with congenital haemostatic defects.](#)
*We investigated 52 of 457 patients with congenital factor deficiencies...*22nd July, 2008
Department of Hematology, All India Institute of Medical Sciences, New- Haemophilia. 2008 Jul 14. ([DOI Direct Link](#))
- [Plasma exchange in a case of severe factor X deficiency in pregnancy: critical review of the literature.](#)
*Hereditary factor X deficiency represents an uncommon challenge in...*27th June, 2008
Department of Obstetrics and Gynecology, Division of Maternal and Fetal- Am J Perinatol. 2008 Mar;25(3):189-92. ([DOI Direct Link](#))
- [Congenital FX deficiency combined with other clotting defects or with other abnormalities: a critical](#)

[evaluation of the literature.](#)

- The presence of more than one congenital clotting defect in a given...* 13th June, 2008
Department of Medical and Surgical Sciences, University of Padua Medical- Haemophilia. 2008 Mar;14(2):323-8. Epub 2007 Dec 10. ([DOI Direct Link](#))
- [Recurrent ovarian haemorrhage in a girl with congenital factor X deficiency.](#)
Congenital Factor X deficiency is commonly classified as type I, in which... 28th May, 2008
Department of Obstetrics & Gynaecology, All India Institute of Medical- Haemophilia. 2008 May 12. ([DOI Direct Link](#))
 - [Perivascular tissue factor is down-regulated following cutaneous wounding: implications for bleeding in hemophilia.](#)
Healing of skin wounds is delayed in hemophilia B (HB) mice. HB mice do... 22nd May, 2008
Duke University, Durham, NC, USA.- Blood. 2008 Feb 15;111(4):2046-8. Epub 2007 Nov 30. ([DOI Direct Link](#))
 - [Congenital bleeding disorders of the vitamin K-dependent clotting factors.](#)
Congenital bleeding disorders of the vitamin K-dependent coagulation... 25th April, 2008
Departement of Medical and Surgical Sciences, University of Padua Medical- Vitam Horm. 2008;78:281-374. ([DOI Direct Link](#))
 - [Factor X Debrecen: Gly204Arg mutation in factor X causes the synthesis of a non-secretable protein and severe factor X deficiency.](#)
Due to a homozygous Gly204Arg mutation in the factor X (FX) gene no... 2nd April, 2008
Clinical Research Center, Thrombosis and Vascular Biology Research Group- Haematologica. 2008 Feb;93(2):299-302. ([DOI Direct Link](#))
 - [A nonstop mutation in the factor \(F\)X gene of a severely haemorrhagic patient with complete absence of coagulation FX.](#)
We identified a previously unknown mutation by sequencing the factor (F)X... 27th February, 2008
Department of Pediatrics, Division of Hematology and Oncology, Medical- Thromb Haemost. 2007 Dec;98(6):1165-9.
 - [Report on a disease-adapted treatment in a patient with severe factor X deficiency resulting from a homozygous factor X gene mutation.](#)
Due to a homozygous Gly204Arg mutation in the factor X (FX) gene no... 22nd February, 2008
- Thromb Haemost. 2008 Jan;99(1):238-9. ([DOI Direct Link](#))
 - [Spontaneous rupture of the spleen as immediate complication in autologous transplantation for primary systemic amyloidosis.](#)
Although splenic rupture is a recognized complication of primary... 6th February, 2008
Department of Hematology, Institute of Hematology and Oncology, Hospital- Eur J Haematol. 2008 Feb;80(2):182-4. Epub 2008 Jan 1. ([DOI Direct Link](#))

BioNews Results for Factor X Deficiency

- [Merck Serono Announces One-Month Results of Pharmacogenomics Study PREDICT at 47th Annual Meeting of the Europ](#)
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- 7427396- [AAV vectors for gene delivery to the lung](#)
- 6093392- [Methods and compositions for use in gene therapy for treatment of hemophilia](#)
- 6080575- [Nucleic acid construct for expressing active substances which can be activated by proteases, and preparation and use](#)
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- 5837679- [Agents affecting thrombosis and hemostasis](#)
- 5830655- [Oligonucleotide sizing using cleavable primers](#)
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- 5700642- [Oligonucleotide sizing using immobilized cleavable primers](#)
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- 5635481- [Recombinant agents affecting thrombosis](#)
- 5597799- [Recombinant agents affecting thrombosis](#)
- 5583107- [Agents affecting thrombosis and hemostasis](#)
- 5415758- [Method and apparatus for electro-elution of biological molecules](#)
- 4725538- [Method of assaying the presence of cancer cells](#)
- 4501731- [Treatment of disparate bleeding disorders with factor X zymogen](#)
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- 6818439- [Methods for administration of recombinant gene delivery vehicles for treatment of hemophilia and other disorders](#)
- 6790838- [Combinations and methods for promoting in vivo liver cell proliferation and enhancing in vivo liver-directed gene transduction](#)
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- 6620310- [Electrochemical coagulation assay and device](#)
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