

Hemochromatosis

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Recent Publications on Hemochromatosis:

- [Genetic mechanisms and modifying factors in hereditary hemochromatosis.](#)
*Primary iron overload is one of the most common inherited diseases...*18th November, 2009
Department of Internal Medicine I, Clinical Immunology and Infectious- Nat Rev Gastroenterol Hepatol. 2009 Nov 17. ([DOI Direct Link](#))
- [Iron overload. Subtle signs, early treatment.](#)
*Hepcidin is the central regulator of systemic iron homeostasis....*18th November, 2009
- Mayo Clin Health Lett. 2009 Oct;27(10):7.
- [HLA-A3-B14 and the origin of the hemochromatosis C282Y mutation. Founder effects and recombination events during 12 generations in a Scandinavian family with major iron overload.](#)
*Background: The hemochromatosis (HH) mutation C282Y occurred once in a...*17th November, 2009
Department of Medicine, Sahlgrenska University Hospital, Goteborg, Sweden.- Eur J Haematol. 2009 Nov 12. ([DOI Direct Link](#))
- [Deferasirox \(Exjade\) for the treatment of iron overload.](#)
*Deferasirox is a once-daily oral iron chelator with established...*13th November, 2009
Fondazione Ospedale Maggiore Policlinico, Istituto di Ricovero e Cura a- Acta Haematol. 2009;122(2-3):165-73. Epub 2009 Nov 10. ([DOI Direct Link](#))
- [Rare types of genetic hemochromatosis.](#)
*Most types of genetic hemochromatosis are due to mutations in the HFE...*13th November, 2009
Vita-Salute University and San Raffaele Scientific Institute, Milan,- Acta Haematol. 2009;122(2-3):140-5. Epub 2009 Nov 10. ([DOI Direct Link](#))
- [The natural history of untreated HFE-related hemochromatosis.](#)
*Hemochromatosis has generally been considered to be a genetic disease in...*13th November, 2009
University- Acta Haematol. 2009;122(2-3):134-9. Epub 2009 Nov 10. ([DOI Direct Link](#))
- [The role of hepcidin in iron metabolism.](#)
*Hepcidin is the central regulator of systemic iron homeostasis....*13th November, 2009
David Geffen School of Medicine, University of California, Los Angeles,- Acta Haematol. 2009;122(2-3):78-86. Epub 2009 Nov 10. ([DOI Direct Link](#))
- [BMP-6 and mesenchymal stem cell differentiation.](#)
*Bone morphogenetic protein-6 (BMP-6) is produced by bone...*11th November, 2009
Laboratory of Mineralized Tissues, Center for Translational Research,- Cytokine Growth Factor Rev. 2009 Nov 7. ([DOI Direct Link](#))
- [Clinical application of cardiac CMR.](#)
*Cardiovascular magnetic resonance (CMR) imaging is an important clinical...*10th November, 2009

Wayne State University School of Medicine, Detroit, MI, USA.- Rev Cardiovasc Med. 2009 Summer;10(3):134-41.

- [Fiber evanescent wave spectroscopy using the mid-infrared provides useful fingerprints for metabolic profiling in humans.](#)
*Fiber evanescent wave spectroscopy (FEWS) explores the mid-infrared...*10th November, 2009
UMR-CNRS 6226, Campus de Beaulieu/Equipe Verres et Ceramiques, University- J Biomed Opt. 2009 Sep-Oct;14(5):054033. ([DOI Direct Link](#))
- [Screening for iron overload: Lessons from the HEMochromatosis and IRon Overload Screening \(HEIRS\) Study.](#)
*BACKGROUND: The HEMochromatosis and IRon Overload Screening (HEIRS) Study...*7th November, 2009
London Health Sciences Centre, London, Canada.- Can J Gastroenterol. 2009 Nov;23(11):769-72.
- [RELATIONSHIP BETWEEN GENE EXPRESSION OF DUODENAL IRON TRANSPORTERS AND IRON STORES IN HEMOCHROMATOSIS SUBJECTS.](#)
*To test the hypothesis that differences in duodenal iron absorption may...*7th November, 2009
1Virginia Mason Medical Center.- Am J Physiol Gastrointest Liver Physiol. 2009 Nov 5. ([DOI Direct Link](#))
- [Neonatal hemochromatosis and Martinez-Frias syndrome of intestinal atresia and diabetes mellitus in a consanguineous newborn.](#)
*Neonatal hemochromatosis is a heterogeneous disorder of iron metabolism...*6th November, 2009
Neonatal Intensive Care Unit, Erasme Hospital, Universite Libre de- Eur J Med Genet. 2009 Nov 1. ([DOI Direct Link](#))
- [Lack of association of C282Y and H63D mutations in the hemochromatosis \(HFE\) gene with diabetes mellitus type 2 in a case-control study of women in Brazil.](#)
*Hereditary hemochromatosis is one of the most common autosomal recessive...*31st October, 2009
Faculdade de Farmacia, Universidade Federal de Minas Gerais, Belo- Genet Mol Res. 2009 Oct 27;8(4):1285-91. ([DOI Direct Link](#))
- [Potential Nonresponse Bias in a Clinical Examination After Initial Screening Using Iron Phenotyping and HFE Genotyping in the Hemochromatosis and Iron Overload Screening Study.](#)
*Background: Little is known about the factors affecting participation in...*29th October, 2009
1 Department of Epidemiology & Biostatistics, University of Western- Genet Test Mol Biomarkers. 2009 Oct 27. ([DOI Direct Link](#))

BioNews Results for Hemochromatosis

- [Hemochromatosis mutations linked to hypertriglyceridemia](#)
MedWire News: Oct 15 2009 5:34PM Matching: hemochromatosis

Hemochromatosis Patents:



- 7510881- [Method and kit for the quantitative and/or qualitative detection of components in a sample](#)

- 7531563- [Thiazoline acid derivatives](#)
- 7534764- [Competitive regulation of hepcidin mRNA by soluble and cell-associated hemojuvelin](#)
- 7541434- [Polypeptide for a TRP channel family member, TRP-PLIK2, and splice variants](#)
- 7547551- [Transfection of eukaryotic cells with linear polynucleotides by electroporation](#)
- 7547698- [Bicyclic heterocyclic derivatives and their use as inhibitors of stearyl-coadesaturase \(SCD\)](#)
- 7547766- [Anti-tango294 antibodies and uses thereof](#)
- 7553627- [Process for high throughput DNA methylation analysis](#)
- 7553855- [Compositions of N-\[2,4-bis\(1,1-dimethylethyl\)-5-hydroxyphenyl\]-1,4-dihydro-4-oxoquinoline-3-carboxamide](#)
- 7560424- [Targeted therapeutic proteins](#)
- 7566533- [Clinically intelligent diagnostic devices and methods](#)
- 7531310- [Methods of diagnosing Crohn's disease by measuring expression level of RNA encoding human G-protein coupled receptor, HGPRBMY11](#)
- 7527962- [Human diacylglycerol acyltransferase 2 \(DGAT2\) family members and uses therefor](#)
- 7527928- [Reactions on a solid surface](#)
- 7511018- [Juvenile hemochromatosis gene \(HFE2A\) cleavage products and uses thereof](#)
- 7511811- [Pulsed-multiline excitation for color-blind fluorescence detection](#)
- 7514419- [Phosphorus-containing thymomimetics](#)
- 7514436- [Pyridazine derivatives and their use as therapeutic agents](#)
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- 7517890- [GTPase inhibitors and methods of use](#)
- 7521055- [Ferroportin1 antibodies and methods](#)
- 7521446- [Haloaryl substituted aminopurines, compositions thereof, and methods of treatment therewith](#)
- 7522757- [Method and apparatus for computer controlled cell based diagnosis](#)
- 7527811- [Topical depigmenting formulations comprising an extract of Bellis perennis](#)
- 7566567- [Immortalized hepatocytes](#)
- 7572596- [Modulation of the neuroendocrine system as a therapy for motor neuron disease](#)
- 7601335- [Pretreatment of a biological sample from an autoimmune disease subject](#)
- 7601359- [Compositions and methods for the prevention and treatment of primary and metastatic neoplastic diseases and infectious diseases with heat shock/stress proteins](#)
- 7601493- [Methods and apparatus for screening and detecting multiple genetic mutations](#)
- 7601684- [Diagnosis and treatment of disorders of iron metabolism](#)
- 7605161- [Pyridyl derivatives and their use as therapeutic agents](#)
- 7608401- [Mutations in the ferroportin 1 gene associated with hereditary haemochromatosis](#)
- 7608406- [Diagnostic markers of stroke and cerebral injury and methods of use thereof](#)
- 7609369- [Methods of detection of iron deficiency and hemochromatosis](#)
- 7611691- [Compounds for treatment of copper overload](#)
- 7611879- [27877, 18080, 14081, 32140, 50352, 16658, 14223, 16002, 50566, 65552 and 65577 molecules and uses therefor](#)
- 7598412- [Modulators of ATP-binding cassette transporters](#)
- 7598348- [Adiponectin receptor variant, AdipoR2v2](#)
- 7598060- [Rapid analysis of variations in a genome](#)
- 7572618- [Polynucleotides encoding novel PCSK9 variants](#)
- 7576131- [Use of \(-\) \(3-trihalomethylphenoxy\) \(4-halophenyl\) acetic acid derivatives for treatment of insulin resistance, type 2 diabetes, hyperlipidemia and hyperuricemia](#)
- 7579169- [Hereditary hemochromatosis gene](#)
- 7582421- [Methods for determination of single nucleic acid polymorphisms using a bioelectronic microchip](#)
- 7582470- [Device for amplifying and detecting a target nucleic acid](#)
- 7585840- [Use of osteoprotegerin for the treatment and/or prevention of fibrotic disease](#)
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- 7592143- [Methods of treating diseases responsive to induction of apoptosis and screening assays](#)
- 7592343- [Pyridazine-piperazine compounds and their use as stearyl-CoA desaturase inhibitors](#)

- 7595385- [Polymorphisms in the region of the human hemochromatosis gene](#)
- 7612107- [Treatment and inhibition of disease conditions using flexible heteroarotinoids](#)

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