

Gaucher Disease

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Recent Publications on Gaucher Disease:



- [Psychological aspects of patients with Fabry disease.](#)
*The Minnesota Multiphasic Personality Inventory (MMPI-2) is widely used in...*20th November, 2009
Pacific Graduate School of Psychology at Palo Alto University, 1791- J Inherit Metab Dis. 2009 Oct 31. ([DOI Direct Link](#))
- [Effective desensitization to imiglucerase in a patient with type I Gaucher disease.](#)
*We describe a child who had anaphylactic hypersensitivity to imiglucerase...*17th November, 2009
Department of Pediatrics, University of Verona, Verona, Italy.- J Pediatr. 2009 Dec;155(6):940-1. ([DOI Direct Link](#))
- [Substrate deprivation therapy in juvenile Sandhoff disease.](#)
*Substrate deprivation therapy has been successfully applied in a number of...*10th November, 2009
Department of Paediatrics, Radboud University Nijmegen Medical Centre,- J Inherit Metab Dis. 2009 Nov 4. ([DOI Direct Link](#))
- [ACOG Committee Opinion No. 442: Preconception and prenatal carrier screening for genetic diseases in individuals of Eastern European Jewish descent.](#)
*Certain autosomal recessive disease conditions are more prevalent in...*6th November, 2009
- Obstet Gynecol. 2009 Oct;114(4):950-3. ([DOI Direct Link](#))
- [Introduction to Brazilian Guidelines to Diagnosis, Treatment, and Monitoring for Gaucher Disease, Fabry Disease, Mucopolysaccharidosis I, and Pompe Disease.](#)
*Chaperone therapy is a newly developed molecular approach to lysosomal...*23rd October, 2009
Universidade Federal de Sao Paulo, Sao Paulo, Brazil- J Pediatr. 2009 Oct;155(4 Suppl):S9.
- [Chaperone Activity of Bicyclic Nojirimycin Analogues for Gaucher Mutations in Comparison with N-\(n-nonyl\)Deoxynojirimycin.](#)
*Gaucher disease (GD), the most prevalent lysosomal storage disorder, is...*16th October, 2009
Division of Child Neurology, Institute of Neurological Sciences, Faculty- Chembiochem. 2009 Oct 14;10(17):2780-2792. ([DOI Direct Link](#))
- [Effort bruising disclosing Gaucher disease in a 55-year-old non-Jewish woman.](#)
*Gaucher disease (GD), the most prevalent lysosomal storage disorder, is...*13th October, 2009
Hematology Center Karolinska, Karolinska University Hospital Huddinge,- J Inherit Metab Dis. 2009 Oct 11. ([DOI Direct Link](#))
- [Gaucher Disease with Communicating Hydrocephalus and Cardiac Involvement.](#)
*A 14-year-old female with Gaucher disease presented with hydrocephalus,...*10th October, 2009
Division of Pediatric Cardiology, Baskent University Hospital, Ankara,- Clin Cardiol. 2009 Oct 8. ([DOI Direct Link](#))

- [Development of genomic DNA reference materials for genetic testing of disorders common in people of ashkenazi jewish descent.](#)
*Many recessive genetic disorders are found at a higher incidence in people...*10th October, 2009
Laboratory Practice Evaluation and Genomics Branch, National Center for- J Mol Diagn. 2009 Nov;11(6):530-6. Epub 2009 Oct 8. ([DOI Direct Link](#))
- [Chaperone therapy for neuronopathic lysosomal diseases: competitive inhibitors as chemical chaperones for enhancement of mutant enzyme activities.](#)
*Chaperone therapy is a newly developed molecular approach to lysosomal...*9th October, 2009
International University of Health and Welfare Graduate School, Kita- Perspect Medicin Chem. 2009 May 26;3:7-19.
- [In vivo and ex vivo evaluation of L-type calcium channel blockers on acid beta-glucosidase in Gaucher disease mouse models.](#)
*Gaucher disease is a lysosomal storage disease caused by mutations in acid...*8th October, 2009
The Division of Human Genetics, Cincinnati Children's Hospital Medical- PLoS One. 2009 Oct 7;4(10):e7320. ([DOI Direct Link](#))
- [Plasma chitotriosidase and CCL18 as surrogate markers for granulomatous macrophages in sarcoidosis.](#)
*BACKGROUND: Accumulation of macrophages in multiple organs is a common...*8th October, 2009
Department of Medical Biochemistry, Academic Medical Center, University of- Clin Chim Acta. 2009 Oct 4. ([DOI Direct Link](#))
- [Force Majeure: Therapeutic measures in response to restricted supply of imiglucerase \(Cerezyme\) for patients with Gaucher disease.](#)
*Gaucher disease is the first lysosomal disorder for which clinically...*7th October, 2009
Department of Endocrinology and Metabolism, Academic Medical Centre,- Blood Cells Mol Dis. 2009 Oct 3. ([DOI Direct Link](#))
- [The clinical course of untreated Gaucher disease in 22 patients over 10 years: hematological and skeletal manifestations.](#)
*Gaucher disease (GD) is a lysosomal storage disorder characterized by...*2nd October, 2009
University of Toronto, Faculty of Medicine, Mount Sinai Hospital, Toronto,- Blood Cells Mol Dis. 2009 Nov-Dec;43(3):289-93. Epub 2009 Sep 30. ([DOI Direct Link](#))
- [Vitamin D Receptor \(VDR\) Polymorphisms in the Cardiac Variant of Gaucher Disease.](#)
*Type 3c Gaucher disease, an ultra-rare cardiac variant with progressive...*29th September, 2009
Gaucher Clinic, Shaare Zedek Medical Center, Jerusalem, 91031, Israel.- Pediatr Cardiol. 2009 Sep 26. ([DOI Direct Link](#))

BioNews Results for Gaucher Disease

- [Gaucher Disease Gene Therapy: New Gaucher disease gene therapy findings from Yale University, Department of Pediatrics published](#)
NewsRX: Nov 16 2009 5:43AM Matching: gaucher disease
- [FDA Grants Priority Review for Shire's velaglucerase alfa for Type 1 Gaucher Disease](#)
BioPharm International: Nov 11 2009 6:49PM Matching: gaucher disease
- [FDA Grants Priority Review for Shire's velaglucerase alfa for Type 1 Gaucher Disease](#)
PR Newswire: Nov 8 2009 5:57PM Matching: gaucher disease
- [FDA Grants Priority Review for Shire's velaglucerase alfa for Type 1 Gaucher Disease](#)
PharmiWeb: Nov 6 2009 5:28AM Matching: gaucher disease
- [SHIRE PLC - FDA Grants Priority Review for Shire's velaglucerase](#)
Individual.com: Nov 5 2009 7:32AM Matching: gaucher disease
- [SHIRE PLC - FDA Grants Priority Review for Shire's velaglucerase](#)

- Pharmacy Choice: Nov 5 2009 3:06AM Matching: gaucher disease*
- [FDA Grants Priority Review for Shire's velaglucerase alfa for Type 1 Gaucher Disease](#)
Pharma Live: Nov 5 2009 1:30PM Matching: gaucher disease
- [FDA Grants Priority Review for Shire's velaglucerase alfa for Type 1 Gaucher Disease](#)
Pharmacy Choice: Nov 5 2009 3:06AM Matching: gaucher disease
- [First Patient Enrolled in Phase 3 Trials of Genzyme Oral Capsule for Gaucher Disease](#)
Pharmacy Choice: Nov 5 2009 3:43PM Matching: gaucher disease
- [FDA Grants Priority Review for Shire's velaglucerase alfa for Type 1 Gaucher Disease](#)
Montana's News Station: Nov 4 2009 7:21PM Matching: gaucher disease
- [FDA Grants Priority Review for Shire's velaglucerase alfa for Type 1 Gaucher Disease](#)
Genetic Engineering News: Nov 4 2009 1:33PM Matching: gaucher disease
- [FDA Grants Priority Review for Shire's velaglucerase alfa for Type 1 Gaucher Disease](#)
Yahoo! Canada: Nov 4 2009 12:25PM Matching: gaucher disease
- [SHIRE PLC - FDA Grants Priority Review for Shire's velaglucerase](#)
Therapeutics Daily: Nov 4 2009 3:30PM Matching: gaucher disease
- [FDA Grants Priority Review for Shire's velaglucerase alfa for Type 1 Gaucher Disease](#)
KOTA Territory News: Nov 4 2009 3:33PM Matching: gaucher disease
- [FDA Grants Priority Review for Shire's velaglucerase alfa for Type 1 Gaucher Disease](#)
Therapeutics Daily: Nov 4 2009 3:31PM Matching: gaucher disease

Gaucher Disease Patents:



- 7557190- [Optimized proteins that target Ep-CAM](#)
- 7572628- [Eukaryotic layered vector initiation systems](#)
- 7575924- [Methods and compositions relating to improved lentiviral vectors and their applications](#)
- 7576101- [Composition for and treatment of demyelinating diseases and paralysis by administration of remyelinating agents](#)
- 7576191- [Tumor suppressor Killin](#)
- 7578999- [Antibodies specific for sclerostin](#)
- 7579166- [Glycoprotein and process for producing the same](#)
- 7579466- [Pyrimidinyl sulfonamide compounds which inhibit leukocyte adhesion mediated by VLA-4](#)
- 7582292- [Adipose tissue derived stromal cells for the treatment of neurological disorders](#)
- 7582734- [Functional antibodies that specifically bind to the Chemerin receptor](#)
- 7584058- [Genetic diagnosis using multiple sequence variant analysis](#)
- 7572618- [Polynucleotides encoding novel PCSK9 variants](#)
- 7569544- [Methods of increasing delivery of active agents to brain comprising administering receptor associated protein \(RAP\) fragments conjugated to active agents](#)
- 7560424- [Targeted therapeutic proteins](#)
- 7560431- [Methods of increasing delivery of active agents to brain comprising administering receptor associated protein \(RAP\) fragments conjugated to active agents](#)
- 7560449- [Methods and therapeutic combinations for the treatment of demyelination](#)
- 7560538- [Porcine isogloboside 3 synthase protein, cDNA, genomic organization, and regulatory region](#)
- 7563459- [Methods and compositions for regenerating tissue](#)
- 7563591- [Method for assaying the activity of lysosomal enzymes](#)

- 7563617- [Hybrid adenovirus/adeno-associated virus vectors and methods of use thereof](#)
- 7569217- [Porcine adenovirus E1 and E4 regions](#)
- 7569384- [Albumin fusion proteins](#)
- 7569385- [Multipotent amniotic fetal stem cells](#)
- 7585840- [Use of osteoprotegerin for the treatment and/or prevention of fibrotic disease](#)
- 7592010- [Albumin fusion proteins](#)
- 7592321- [Methods of treating lysosomal storage related diseases by gene therapy](#)
- 7611879- [27877, 18080, 14081, 32140, 50352, 16658, 14223, 16002, 50566, 65552 and 65577 molecules and uses therefor](#)
- 7612050- [Drug for the treatment of osteonecrosis and for the management of patients at risk of developing osteonecrosis](#)
- 7612080- [GTPase inhibitors and use thereof for controlling platelet hyperactivity](#)
- 7612093- [Compositions of treating hepatitis virus infections with N-substituted-1,5-dideoxy-1,5-imino-D-glucitol compounds in combination therapy](#)
- 7612107- [Treatment and inhibition of disease conditions using flexible heteroarotinoids](#)
- 7612169- [Osteoprotegerin variant proteins](#)
- 7612178- [Anti-IGF-1R antibodies and uses thereof](#)
- 7615217- [Artificial proteins with reduced immunogenicity](#)
- 7615224- [Multiplex-bead complex for determination of lysosomal storage disorders](#)
- 7615573- [Synthesis of UDP-glucose: N-acylsphingosine glucosyltransferase inhibitors](#)
- 7608422- [Simian immunodeficiency virus \(SIV\) molecular clone encoding mutant gag gene lacking inhibitory/instability regions](#)
- 7605249- [Treatment of neurodegenerative disease through intracranial delivery of siRNA](#)
- 7592429- [Sclerostin-binding antibody](#)
- 7593818- [Genetic diagnosis using multiple sequence variant analysis](#)
- 7595038- [Methods of treatment using antibodies to follistatin-3](#)
- 7595318- [Polyethylene glycol conjugates of heterocycloalkyl carboxamido propanoic acids](#)
- 7598031- [Method for the detection of gene transcripts in blood and uses thereof](#)
- 7598060- [Rapid analysis of variations in a genome](#)
- 7598348- [Adiponectin receptor variant, AdipoR2v2](#)
- 7604960- [Transient protein expression methods](#)
- 7604990- [Methods of regulating differentiation in stem cells](#)
- 7605239- [Skeletal-targeted radiation to treat bone-associated pathologies](#)
- 7618948- [Devices, systems and methods for improving and/or cognitive function through brain delivery of siRNA](#)

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