

Mental Retardation

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Recent Publications on Mental Retardation:



- [Axenfeld-Rieger ocular anomaly and retinoblastoma caused by constitutional chromosome 13q deletion.](#)
*Axenfeld-Rieger (AR) ocular anomaly might be due to deletions of different...*21st November, 2009
Department of Child Neurology, Hospital Sant Joan de Deu, Barcelona,- *Pediatr Blood Cancer*. 2009 Nov 19. ([DOI Direct Link](#))
- [Examination of FMR1 transcript and protein levels among 74 premutation carriers.](#)
*Fragile X-associated disorders are caused by a CGG trinucleotide repeat...*21st November, 2009
Department of Human Genetics, Emory University School of Medicine,- *J Hum Genet*. 2009 Nov 20. ([DOI Direct Link](#))
- [Sudden death in a child with Carpenter Syndrome. Case report and literature review.](#)
*Carpenter syndrome (Acrocephalopolysyndactyly type II) is a rare disorder...*20th November, 2009
Department of Pathology and Laboratory Medicine, University of Kentucky- *Forensic Sci Med Pathol*. 2009 Nov 19. ([DOI Direct Link](#))
- [The M-current inhibitor XE991 decreases the stimulation threshold for long-term synaptic plasticity in healthy mice and in models of cognitive disease.](#)
*Aging, mental retardation, number of psychiatric and neurological...*19th November, 2009
Division de Neurociencias, Universidad Pablo de Olavide de Sevilla,- *Hippocampus*. 2009 Nov 17. ([DOI Direct Link](#))
- [Prenatal diagnosis of two different unbalanced forms of an inherited \(Y;12\) translocation.](#)
*The identification of an unexpected structural chromosome rearrangement at...*19th November, 2009
Servei de Bioquímica i Genètica Molecular, Hospital Clinic, Barcelona,- *Am J Med Genet A*. 2009 Nov 16. ([DOI Direct Link](#))
- [Autosomal dominant syndrome of mental retardation, hypotelorism, and cleft palate resembling Schilbach-Rott syndrome.](#)
*We present a family segregating for an autosomal dominant syndrome of...*19th November, 2009
The Raphael Recanati Genetics Institute, Rabin Medical Center, Beilinson- *Am J Med Genet A*. 2009 Nov 16. ([DOI Direct Link](#))
- [Characterization of double ring chromosome 4 mosaicism associated with bilateral hip dislocation, cortical dysgenesis, and epilepsy.](#)
*We present the clinical and molecular findings in a Turkish child with a...*19th November, 2009
Afyon Kocatepe University Faculty of Medicine, Department of Medical- *Am J Med Genet A*. 2009 Nov 16. ([DOI Direct Link](#))
- [Mandibulofacial syndrome with growth and mental retardation, microcephaly, ear anomalies with skin](#)

[tags, and cleft palate in a mother and her son: Autosomal dominant or X-linked syndrome?](#)

We report on a Brazilian mother and her son affected with mandibulofacial... 19th November, 2009
Department of Clinical Genetics, Hospital of Rehabilitation of- Am J Med Genet A. 2009 Nov 16. ([DOI Direct Link](#))

- [Novel variants identified in methyl-CpG-binding domain genes in autistic individuals.](#)
Misregulation of the methyl-CpG-binding protein 2 (MECP2) gene has been... 19th November, 2009
John P. Hussman Institute for Human Genomics, University of Miami, 1501 NW- Neurogenetics. 2009 Nov 18. ([DOI Direct Link](#))
- [Identification of ANKRD11 and ZNF778 as candidate genes for autism and variable cognitive impairment in the novel 16q24.3 microdeletion syndrome.](#)
The clinical use of array comparative genomic hybridization in the... 19th November, 2009
Department of Human Genetics, Radboud University Nijmegen Medical Centre,- Eur J Hum Genet. 2009 Nov 18. ([DOI Direct Link](#))
- [Heparin binding epidermal growth factor-like growth factor reduces ethanol-induced apoptosis and differentiation in human embryonic stem cells.](#)
Alcohol affects approximately 1% (40,000) of new born infants each year... 19th November, 2009
Department of Anesthesiology, Emory University School of Medicine,- Growth Factors. 2009 Dec;27(6):362-369.
- [Longitudinal course of cognitive, adaptive, and behavioral characteristics in Costello syndrome.](#)
Costello syndrome is a rare rasopathy caused by germline mutations in the... 18th November, 2009
Psychology Service, Department of Pediatrics, Texas Children's Hospital,- Am J Med Genet A. 2009 Nov 16. ([DOI Direct Link](#))
- [Sjogren-Larsson syndrome in two brothers: a case report.](#)
Sjogren-Larsson syndrome is a rare autosomal recessive disorder that was... 18th November, 2009
- Cases J. 2009 Sep 9;2:8434. ([DOI Direct Link](#))
- [MPEP Reduces Seizure Severity in Fmr-1 KO mice over Expressing Human Abeta.](#)
Metabotropic glutamate receptor 5 (mGluR(5)) regulates the translation of... 18th November, 2009
Department of Pathology & Laboratory Medicine and Waisman Center for- Int J Clin Exp Pathol. 2009 Oct 10;3(1):56-68.
- [Frequency and phenotype of SPG11 and SPG15 in complicated hereditary spastic paraplegia.](#)
BACKGROUND: Hereditary spastic paraplegias (HSP) are clinically and... 18th November, 2009
Department of Neurology, University of Tübingen, Tübingen, Germany.- J Neurol Neurosurg Psychiatry. 2009 Dec;80(12):1402-4. ([DOI Direct Link](#))

BioNews Results for Mental Retardation

- [Researchers at Signature Genomics Diagnose Rare Syndrome in Children with Unexplained Mental Retardation and B](#)
PharmiWeb: Nov 19 2009 1:41AM Matching: mental retardation
- [Researchers at Signature Genomics Diagnose Rare Syndrome in Children with Unexplained Mental Retardation and B](#)
NewsBlaze: Nov 18 2009 2:17AM Matching: mental retardation
- [Researchers at Signature Genomics Diagnose Rare Syndrome in Children with Unexplained Mental Retardation and Birth Defects](#)
Centre Daily Times: Nov 18 2009 12:11AM Matching: mental retardation
- [Mental Health](#)
NOLA38: Nov 18 2009 10:33AM Matching: mental retardation
- [Researchers at Signature Genomics Diagnose Rare Syndrome in Children with Unexplained Mental Retardation and B](#)

- Reuters: Nov 17 2009 11:43PM Matching: mental retardation*
- [Researchers at Signature Genomics Diagnose Rare Syndrome in Children with Unexplained Mental Retardation and B](#)
- Genetic Engineering News: Nov 17 2009 11:27PM Matching: mental retardation*
- [Researchers at Signature Genomics Diagnose Rare Syndrome in Children with Unexplained Mental Retardation and B](#)
- Street Insider: Nov 17 2009 10:42PM Matching: mental retardation*
- [Researchers at Signature Genomics Diagnose Rare Syndrome in Children with Unexplained Mental Retardation and Birth](#)
- PR inside: Nov 17 2009 10:25PM Matching: mental retardation*
- [Researchers at Signature Genomics Diagnose Rare Syndrome in Children with Unexplained Mental Retardation and B](#)
- Business Wire via Yahoo!: Nov 17 2009 10:22PM Matching: mental retardation*
- [Researchers at Signature Genomics Diagnose Rare Syndrome in Children with Unexplained Mental Retardation and B](#)
- RBC Dain Rauscher: Nov 17 2009 10:14PM Matching: mental retardation*
- [Researchers at Signature Genomics Diagnose Rare Syndrome in Children with Unexplained Mental Retardation and Birth Defects](#)
- Business Wire UK: Nov 17 2009 10:11PM Matching: mental retardation*
- [Researchers at Signature Genomics Diagnose Rare Syndrome in Children with Unexplained Mental Retardation and B](#)
- Business Wire: Nov 17 2009 10:10PM Matching: mental retardation*
- [Researchers at Signature Genomics Diagnose Rare Syndrome in Children with Unexplained Mental Retardation and B](#)
- Yahoo! Canada: Nov 17 2009 10:19PM Matching: mental retardation*
- [Researchers at Signature Genomics Diagnose Rare Syndrome in Children with Unexplained Mental Retardation and B](#)
- CNBC: Nov 17 2009 10:13PM Matching: mental retardation*
- [City caps mental health aid at \\$100,000](#)
- El Paso Times: Nov 12 2009 2:59AM Matching: mental retardation*

Mental Retardation Patents:



- 7566714- [Methods and compositions for the treatment of metabolic disorders](#)
- 7585629- [Compositions and methods for spinocerebellar ataxia](#)
- 7585634- [Method of predicting risk of lung cancer recurrence, and a composition, kit and microarray for the same](#)
- 7585881- [Additional heteropolycyclic compounds and their use as metabotropic glutamate receptor antagonists](#)
- 7585949- [IL-1 zeta splice variant polypeptide](#)
- 7585996- [Acyloxyalkyl carbamate prodrugs, methods of synthesis and use](#)
- 7592010- [Albumin fusion proteins](#)
- 7592321- [Methods of treating lysosomal storage related diseases by gene therapy](#)
- 7592438- [Human gene](#)

- 7595159- [Prediction of Parkinson's disease using gene expression levels of peripheral blood samples](#)
- 7595315- [Morpholine derivatives as norepinephrine reuptake inhibitors](#)
- 7585626- [Methods for nucleic acid amplification](#)
- 7582734- [Functional antibodies that specifically bind to the Chemerin receptor](#)
- 7569384- [Albumin fusion proteins](#)
- 7572442- [Selected antibody compositions for binding to aminophospholipids](#)
- 7572448- [Combined cancer treatment methods using selected antibodies to aminophospholipids](#)
- 7572580- [Promoter variants of the alpha-7 nicotinic acetylcholine receptor](#)
- 7572621- [Detection, characterization and treatment of viral infection and methods thereof](#)
- 7575745- [Ligand for herpes simplex virus entry mediator and methods of use](#)
- 7576077- [Fused heterocyclic compounds and their use as metabotropic glutamate receptor antagonists](#)
- 7576101- [Composition for and treatment of demyelinating diseases and paralysis by administration of remyelinating agents](#)
- 7580798- [Method for predicting treatment classes using animal behavior informatics](#)
- 7582416- [Ligand for G-protein coupled receptor FPRL2 and uses thereof](#)
- 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](#)
- 7612226- [Amino acid derivatives](#)
- 7613343- [Nosologic system of diagnosis](#)
- 7615219- [Inhibition of PACAP signalling for the prevention and treatment of thrombocytopenia](#)
- 7615223- [Selected immunoconjugates for binding to aminophospholipids](#)
- 7615224- [Multiplex-bead complex for determination of lysosomal storage disorders](#)
- 7618621- [Mammalian multipotent neural stem cells and compositions, methods of preparation and methods of administration thereof](#)
- 7618948- [Devices, systems and methods for improving and/or cognitive function through brain delivery of siRNA](#)
- 7619005- [Methods for treating cognitive impairment in humans with Multiple Sclerosis](#)
- 7619007- [Method and composition for administering an NMDA receptor antagonist to a subject](#)
- 7619096- [3-Aminopyrrolidines as inhibitors of monoamine uptake](#)
- 7612107- [Treatment and inhibition of disease conditions using flexible heteroarotinoids](#)
- 7612038- [Biglycan and related therapeutics and methods of use](#)
- 7601127- [Therapeutic stimulation of genital tissue or reproductive organ of an infertility or impotence diagnosed patient](#)
- 7604948- [Biomarkers for diagnosing an autism spectrum disorder](#)
- 7604958- [Synthetic mammalian .alpha.-N-acetylglucosaminidase and genetic sequences encoding same](#)
- 7605241- [Synthesis of inhibitors of p90Rsk](#)
- 7608685- [Tumor antigen based on products of the tumor suppressor gene WT1](#)
- 7608704- [Secreted proteins](#)
- 7611704- [Compositions and methods for treating viral infections using antibodies and immunoconjugates to aminophospholipids](#)
- 7611718- [Attenuation of cytomegalovirus virulence](#)
- 7611839- [Methods for diagnosing RCC and other solid tumors](#)
- 7611869- [Multiplexed methylation detection methods](#)
- 7619110- [Amino acid derived prodrugs of propofol, compositions, uses and crystalline forms thereof](#)

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