

## Genetic Inborn

[View the current Genetic Inborn InDepth page on BioPortfolio.com \(PDF\)](#)

([http://www.bioportfolio.com/indepth/Genetic\\_Inborn.html](http://www.bioportfolio.com/indepth/Genetic_Inborn.html)) - Regularly Updated.



### Recent Publications on Genetic Inborn:

- [\[Importance of studies about the etiopathogenesis of childhood diseases for clinical practice. Introduction.\]](#)  
*Genetic factors have long been recognized as contributors to variantion in...* 18th November, 2009  
- Med Wieku Rozwoj. 2009 Apr-Jun;13(2):79-80.
- [A new way to classify genetic skin disease.](#)  
*Genetic disorders with skin manifestations often affect other organs as...* 13th November, 2009  
Department of Dermatology, Birmingham Children's Hospital, Birmingham, UK.- J Invest Dermatol. 2009 Nov;129(11):2543-5. ([DOI Direct Link](#))
- [CONSANGUINITY IN QATAR: KNOWLEDGE, ATTITUDE AND PRACTICE IN A POPULATION BORN BETWEEN 1946 AND 1991.](#)  
*SummaryFrom March 2007 to March 2008 a cross-sectional study was conducted...* 10th November, 2009  
\*Shafallah Medical Genetics Center, Doha, Qatar.- J Biosoc Sci. 2009 Nov 9:1-24. ([DOI Direct Link](#))
- [\[Between theory and interpretation of the hereditary transmission process of mental disease. The introduction of Mendelism in German and North American psychiatry, 1911-1930\]](#)  
*1911 saw the beginning of decisive developments in psychiatric genetic...* 10th November, 2009  
Institut fur Geschichte der Medizin der Universitat Giessen, Giessen.- NTM. 2009;17(1):35-54.
- [\[Cerebral creatine deficiency: First Spanish patients harbouring mutations in GAMT gene.\]](#)  
*BACKGROUND AND OBJETIVE: Brain creatine (Cr) deficiencies are a group of...* 7th November, 2009  
Neuropediatrics, Hospital Sant Joan de Deu y Centro de Investigacion- Med Clin (Barc). 2009 Nov 21;133(19):745-9. Epub 2009 Nov 5. ([DOI Direct Link](#))
- [Old and new ideas about genes and behaviour.](#)  
*Genetic factors have long been recognized as contributors to variantion in...* 7th November, 2009  
Department of Molecular Biology, Umea University, Umea, Sweden.- Hereditas. 2009 Oct;146(5):198-203. ([DOI Direct Link](#))
- [RAS and CSF3R mutations in severe congenital neutropenia.](#)  
*Genetic factors have long been recognized as contributors to variantion in...* 7th November, 2009  
- Blood. 2009 Oct 15;114(16):3504-5. ([DOI Direct Link](#))
- [Clinical diagnostics in human genetics with semantic similarity searches in ontologies.](#)  
*The differential diagnostic process attempts to identify candidate...* 29th October, 2009  
Institute for Medical Genetics, Charite-Universitatsmedizin Berlin,- Am J Hum Genet. 2009 Oct;85(4):457-64. ([DOI Direct Link](#))
- [When should children be tested for genetic diseases?](#)

- OBJECTIVES: Hemoglobinopathies are the most common hereditary disorders in...* 24th October, 2009  
Division of Genetics, Department of Pediatrics, University of Texas- Pediatrics. 2009  
Oct;124(4):e807-8. Epub 2009 Sep 21. ([DOI Direct Link](#))
- [Finding the missing heritability of complex diseases.](#)  
*Genome-wide association studies have identified hundreds of genetic...* 21st October, 2009  
National Human Genome Research Institute, Building 31, Room 4B09, 31- Nature. 2009 Oct  
8;461(7265):747-53. ([DOI Direct Link](#))
  - [Doctors explore research potential of blood samples from newborns.](#)  
*Sulfite oxidase is a mitochondrial enzyme encoded by the SUOX gene and...* 20th October, 2009  
- BMJ. 2009 Sep 28;339:b3984. doi: 10.1136/bmj.b3984.
  - [Functional deficiencies of sulfite oxidase: Differential diagnoses in neonates presenting with intractable seizures and cystic encephalomalacia.](#)  
*Sulfite oxidase is a mitochondrial enzyme encoded by the SUOX gene and...* 2nd October, 2009  
Labor fur Klinische Biochemie und Stoffwechsel, Zentrum fur Kinder- und- Brain Dev. 2009 Sep 28. ([DOI Direct Link](#))
  - [Contiguous gene deletion syndrome in a female with ornithine transcarbamylase deficiency.](#)  
*OTC deficiency, a partially dominant X-linked trait, is the most frequent...* 29th September, 2009  
Genetic Metabolic Disorders Service, The Children's Hospital at Westmead,- Mol Genet Metab. 2009  
Aug 26. ([DOI Direct Link](#))
  - [Xp22.3 Genomic deletions involving the CDKL5 gene in girls with early onset epileptic encephalopathy.](#)  
*Summary Purpose: Mutations of the X-linked gene cyclin-dependent...* 29th September, 2009  
Pediatric Neurology Unit and Laboratories, Children's Hospital A. Meyer -- Epilepsia. 2009 Sep 22. ([DOI Direct Link](#))
  - [\[Diagnosis of hemoglobinopathies in newborns in Venezuela hospitals.\]](#)  
*OBJECTIVES: Hemoglobinopathies are the most common hereditary disorders in...* 25th September, 2009  
Laboratorio de Investigacion de Hemoglobinas Anormales (LIHA), Hospital- An Pediatr (Barc). 2009  
Oct;71(4):314-8. Epub 2009 Sep 22. ([DOI Direct Link](#))

## BioNews Results for Genetic Inborn

- [Genetic Testing! The Solution to All Your Parenting Woes!](#)  
*GenomeWeb: Nov 20 2009 7:25PM Matching: genetic inborn*
- [Vaccination Safe for Kids With Genetic Disorders](#)  
*eMedicine Health: Nov 3 2009 12:58PM Matching: genetic inborn*

\*\*\*

Resources from the [NCBI](#) used in this document, [NCBI's standard disclaimer applies](#).

Nothing in this document should be used in place of personal medical advice from your own qualified medical practitioner. See BioPortfolio.com [User Agreement](#)

Send comments and feedback to:

Peter Barfoot Managing Director, BioPortfolio Ltd.

UK Tel: (+44) 1300 321501

USA Voicemail and Fax: (+1) 415 680 2472

[Peter Barfoot peter.barfoot@bioportfolio.com](mailto:peter.barfoot@bioportfolio.com)

All rights reserved. All other trademarks recognized.

BioPortfolio Limited is registered in England & Wales at Stafford House, 10 Prince of Wales Road, Dorchester, Dorset, DT1 1PW, UK. No.3312883 VAT No. GB 744 6483 10

Copyright 1997-2009 - BioPortfolio Limited.

