

Cafe-Au-Lait Spots

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Recent Publications on Cafe-Au-Lait Spots:

- [Combined retinal hamartomas leading to the diagnosis of neurofibromatosis type 2.](#)
*PURPOSE: To report two cases of neurofibromatosis type 2 (NF2) initially...*26th September, 2008
School of Medicine, Oregon Health & Science University, Portland, Oregon- Ophthalmic Genet. 2008 Sep;29(3):133-8. ([DOI Direct Link](#))
- [Ocular manifestations of neurofibromatosis 1--m. Recklinghausen.](#)
*Neurofibromatosis type 1 (NF 1)--morbus von Recklinghausen is an autosomal...*26th September, 2008
Department of Paediatric Ophtalmology, Comenius University, Children- Bratisl Lek Listy. 2008;109(6):259.
- [Leopard syndrome.](#)
*The L.E.O.P.A.R.D. syndrome is an autosomal, dominant disorder with...*25th September, 2008
University of Rome La Sapienza, Department of Dermatology, Policlinico- Dermatol Online J. 2008 Mar 15;14(3):7.
- [Constitutional mismatch repair-deficiency syndrome: have we so far seen only the tip of an iceberg?](#)
*Heterozygous mutations in one of the mismatch repair (MMR) genes MLH1,...*24th September, 2008
Department of Medical Genetics, Medical University of Vienna,- Hum Genet. 2008 Sep;124(2):105-22. Epub 2008 Aug 18. ([DOI Direct Link](#))
- [Neurofibromatosis type 1 with intracranial hemorrhage and horseshoe kidney.](#)
*A 12-year-old boy presented with a history of sudden-onset vomiting,...*23rd September, 2008
Department of Pediatrics, Advanced Pediatric Center, Postgraduate- Pediatr Neurol. 2008 Oct;39(4):295-7. ([DOI Direct Link](#))
- [LEOPARD syndrome: what are cafe noir spots?](#)
*Lentigines, electrocardiographic abnormalities, ocular hypertelorism,...*16th September, 2008
Service of Dermatology, Universitary Hospital Reina Sofia, Cordoba, Spain.- Pediatr Dermatol. 2008 Jul-Aug;25(4):444-8. ([DOI Direct Link](#))
- [\[Neurofibromatosis type 1 in children. Experiences of the Gdansk Paediatric Oncohaematology Centre. Preliminary results\]](#)
*INTRODUCTION: Neurofibromatosis type 1 (NF1) is a frequent genetic...*30th August, 2008
Klinika Pediatrii, Hematologii, Onkologii i Endokrynologii, Akademia- Med Wieku Rozwoj. 2007 Jul-Sep;11(3 Pt 2):307-12.
- [Ocular manifestations of neurofibromatosis 1 - m. Recklinghausen.](#)
*Neurofibromatosis type 1 (NF 1) - morbus von Recklinghausen is an...*30th August, 2008
Department of Paediatric Ophtalmology, Comenius University, Children- Bratisl Lek Listy. 2008;109(5):246-7.

- [Cutaneous features associated with microcephalic osteodysplastic primordial dwarfism type II.](#)
*We present an 18-month-old Pakistani girl who had short stature,...*30th August, 2008
Department of Dermatology, Barts and the London NHS Trust, London, UK.- *Pediatr Dermatol.* 2008 May-Jun;25(3):401-2. ([DOI Direct Link](#))
- [Autoimmune diseases associated with neurofibromatosis type 1.](#)
*Associations of autoimmune diseases with neurofibromatosis type 1 have...*30th August, 2008
Pediatric Dermatology Unit, As'ad Al-Hamad Dermatology Center, Al-Sabah- *Pediatr Dermatol.* 2008 May-Jun;25(3):392-3. ([DOI Direct Link](#))
- [Segmental neurofibromatosis associated with renal angiomyolipomas.](#)
*Segmental neurofibromatosis (SN) is a rare disorder characterized by...*21st August, 2008
Division of Dermatology, David Geffen School of Medicine, University of- *Cutis.* 2008 Jul;82(1):65-8.
- [Bilateral nevus of Ito and nevus spilus in the same patient.](#)
*INTRODUCTION: Neurofibromatosis type 1 (NF1) is a frequent genetic...*14th August, 2008
- *J Am Acad Dermatol.* 2008 Aug;59(2 Suppl 1):S51-2. ([DOI Direct Link](#))
- [Giant plexiform neurofibroma and suboccipital meningocele manifesting as segmental neurofibromatosis.](#)
*A 34-year-old woman presented with segmental neurofibromatosis manifesting...*14th August, 2008
Department of Neurosurgery, University of Toyama, Toyama, Japan.- *Neurol Med Chir (Tokyo).* 2008 Jun;48(6):271-4.
- [Zebrafish with mutations in mismatch repair genes develop neurofibromas and other tumors.](#)
*Defective mismatch repair (MMR) in humans causes hereditary nonpolyposis...*6th August, 2008
Hubrecht Institute for Developmental Biology and Stem Cell Research,- *Cancer Res.* 2008 Jul 1;68(13):5059-66. ([DOI Direct Link](#))
- [Androgen receptor overexpression in Becker nevus: histopathologic and immunohistochemical analysis.](#)
*Background: Becker nevus (BN) is a unilateral, hairy, pigmented cutaneous...*12th July, 2008
Department of Dermatology, Ajou University School of Medicine, Suwon,- *J Cutan Pathol.* 2008 Jun 18. ([DOI Direct Link](#))

Cafe-Au-Lait Spots Clinical Trials:



- [Effects of Letrozole on Precocious Puberty Due to McCune Albright Syndrome](#)
McCune Albright Syndrome; Polyostotic Fibrous Dysplasia; Precocious Puberty

Cafe-Au-Lait Spots Patents:



- 7416724- [Diagnosis and treatment of diseases arising from defects in the tuberous sclerosis pathway](#)
- 5391575- [Method for treating neurofibromatosis](#)
- 5496731- [Broad-spectrum tumor suppressor genes, gene products and methods for tumor suppressor](#)

[gene therapy](#)

- 5576013- [Treating vascular and neoplastic tissues](#)
- 5691460- [Epidermal surface antigen gene](#)
- 5859195- [Neurofibromatosis gene](#)
- 5912236- [Broad-spectrum tumor suppressor genes gene products and methods for tumor suppressor gene therapy](#)
- 6090794- [Inhibition of neurofibrosarcoma growth and angiogenesis](#)
- 6104959- [Method and apparatus for treating subcutaneous histological features](#)
- 6238861- [Neurofibromatosis gene](#)
- 6362392- [Nude mouse model for the growth and treatment of human neurally-derived tumors](#)
- 6602856- [Antagonists of alpha-melanocyte stimulating hormone and methods based thereon](#)
- 6670122- [Method for detecting transcription templates](#)
- 6759525- [Protein kinase A and carney complex](#)
- 7169594- [Diagnosis and treatment of diseases arising from defects in the tuberous sclerosis pathway](#)
- 7335469- [Methods for diagnosing Noonan syndrome](#)
- 7335470- [Compilations of nucleic acids and arrays and methods of using them](#)
- 7359748- [Apparatus for total immersion photography](#)
- 4582787- [Method of testing a patient for a predisposition to lung cancer, certain other cancers, neurofibromatosis and certain other hereditary disorders](#)

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