

Langer-Giedion Syndrome

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Recent Publications on Langer-Giedion Syndrome:

- [Further case of microdeletion of 8q24 with phenotype overlapping Langer-Giedion without TRPS1 deletion.](#)
*Langer-Giedion syndrome results from a microdeletion at 8q24.1...*17th June, 2008
Our Lady's Children's Hospital, National Centre for Medical Genetics,- *Am J Med Genet A.* 2008 Jun 15;146A(12):1587-92. ([DOI Direct Link](#))
- [A novel model surgery technique for LeFort III advancement.](#)
*Current techniques for model surgery and occlusal splint fabrication lack...*7th December, 2007
Division of Dentistry, Children's Hospital Los Angeles, USC School of - *J Craniofac Surg.* 2007 Sep;18(5):1230-5. ([DOI Direct Link](#))
- [Trichorhinophalangeal syndrome type II without the chromosome 8 deletion that resembled metachondromatosis.](#)
*We describe a 5-year-old girl with features resembling...*14th November, 2007
Department of Orthopaedic Surgery, Yamagata University School of Medicine,- *Congenit Anom (Kyoto).* 2007 Sep;47(3):105-7. ([DOI Direct Link](#))
- [Clinical and intraoral findings of a patient with tricho-rhino-phalangeal syndrome type I.](#)
*Tricho-rhino-phalangeal syndrome (TRPS) is a rare and an autosomal...*11th September, 2007
Department of Orthodontics, Gulhane Military Medical Academy, Ankara,- *J Indian Soc Pedod Prev Dent.* 2007 Mar;25(1):43-5.
- [Trichorhinophalangeal syndrome type III in Turkish siblings.](#)
*Tricho-rhino-phalangeal syndrome (TRPS) is a rare and an autosomal...*23rd February, 2007
- *Acta Paediatr.* 2006 Nov;95(11):1511-3. ([DOI Direct Link](#))
- [Ectrodactyly with aplasia of long bones \(OMIM; 119100\) in a large inbred Arab family with an apparent autosomal dominant inheritance and reduced penetrance: clinical and genetic analysis.](#)
*Ectrodactyly with aplasia of long bones syndrome is one of the most...*9th December, 2006
Center for Arab Genomic Studies (CAGS), Dubai, United Arab Emirates.- *Am J Med Genet A.* 2006 Jul 1;140(13):1440-6. ([DOI Direct Link](#))
- [Postmortem findings at 36 weeks gestation in trichorhinophalangeal syndrome: a case report.](#)
*Ectrodactyly with aplasia of long bones syndrome is one of the most...*9th June, 2006
- *Pathology.* 2006 Apr;38(2):170-2. ([DOI Direct Link](#))
- [Trichorhinophalangeal syndrome.](#)
*Langer-Giedion syndrome results from a genetic deletion on chromosome 8....*24th May, 2006
Institute of Dermatology, University of Messina, Messina, Italy.- *J Am Acad Dermatol.* 2005 Nov;53(5):858-60. ([DOI Direct Link](#))

- [Langer-Giedion Syndrome.](#)
Ectrodactyly with aplasia of long bones syndrome is one of the most... 19th May, 2006
Department of Pediatrics, Advanced Pediatric Center, PGIMER, Chandigarh- Indian Pediatr. 2006 Feb;43(2):174-5.
- [Carpal tunnel syndrome in a patient with trichorhinophalangeal syndrome.](#)
Ectrodactyly with aplasia of long bones syndrome is one of the most... 13th April, 2006
Department of Plastic and Reconstructive Surgery, St. James's University- Plast Reconstr Surg. 2006 Mar;117(3):36e-39e. ([DOI Direct Link](#))
- [Langer-giedion syndrome with renal cyst.](#)
Here it is reported a 4-year-old boy with Langer-Giedion syndrome... 6th April, 2006
Department of Pediatrics, JJMMC, Davangere, Karnataka, India.- Indian J Pediatr. 2005 Feb;72(2):181.
- [Lower eyelid entropion in Langer-Giedion syndrome.](#)
Langer-Giedion syndrome results from a genetic deletion on chromosome 8... 29th March, 2006
Cullen Eye Institute, Baylor College of Medicine, Department of- Ophthal Plast Reconstr Surg. 2006 Jan-Feb;22(1):59-60.
- [V-shaped, longitudinal nail dystrophies in trichorhinophalangeal syndrome type I.](#)
Trichorhinophalangeal syndrome (TRPS) is a rare genodermatosis with growth... 13th December, 2005
Department of Dermatology, University Hospital Basel, Basel, Switzerland.- Dermatology. 2005;211(2):162-4. ([DOI Direct Link](#))
- [The gene associated with trichorhinophalangeal syndrome in humans is overexpressed in breast cancer.](#)
A comprehensive differential gene expression screen on a panel of 54... 21st September, 2005
Sanofi Pasteur, Toronto,- Proc Natl Acad Sci U S A. 2005 Aug 2;102(31):11005-10. Epub 2005 Jul 25. ([DOI Direct Link](#))
- [A young girl with abnormal hands, symptoms of short stature. Trichorhinophalangeal syndrome.](#)
Here it is reported a 4-year-old boy with Langer-Giedion syndrome... 20th April, 2005
Feinberg School of Medicine, Northwestern University, Chicago, IL, USA.- Pediatr Ann. 2005 Jan;34(1):20, 22.

Langer-Giedion Syndrome Patents:



- 7368527- [HADDE71 polypeptides](#)
- 5888740- [Detection of aneuploidy and gene deletion by PCR-based gene- dose co-amplification of chromosome specific sequences with synthetic sequences with synthetic internal controls](#)
- 5976790- [Comparative Genomic Hybridization \(CGH\)](#)
- 6093699- [Method for gene therapy involving suppression of an immune response](#)
- 6251601- [Simultaneous measurement of gene expression and genomic abnormalities using nucleic acid microarrays](#)
- 6287802- [EXT2 gene](#)
- 6303294- [Methods of detecting genetic deletions and mutations associated with Digeorge syndrome, Velocardiofacial syndrome, CHARGE association, conotruncal cardiac defect, and cleft palate and probes useful therefore](#)
- 6335167- [Comparative genomic hybridization \(CGH\)](#)
- 6602693- [Gene encoding hyaluronan synthase](#)

- 6919431- [Polypeptide-human SNARE protein 25 and a polynucleotide encoding the same](#)
- 7035739- [Computer systems and methods for identifying genes and determining pathways associated with traits](#)
- 7094534- [Detection of chromosomal abnormalities associated with breast cancer](#)
- 7273724- [Polypeptide-human actin-binding protein 54 and a polynucleotide encoding the same](#)
- 7291461- [Methods for identifying small molecules that modulate premature translation termination and nonsense mRNA decay](#)
- 7335470- [Compilations of nucleic acids and arrays and methods of using them](#)
- 7351529- [Methods for detecting genetic mosaicisms using arrays](#)
- 5576178- [Method of detecting genetic deletions and mutations associated with DiGeorge syndrome, Velocardiofacial syndrome, charge association, conotruncal cardiac defect, and cleft palate and probes useful therefor](#)

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