

# CITATION REPORT

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**Nosology and classification of genetic skeletal disorders: 2006 revision**

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**American Journal of Medical Genetics, Part A, 2007, 143A, 1-18.**

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#	Paper	IF	Citations
273	The Shwachman-Bodian-Diamond syndrome gene mutations cause a neonatal form of spondylometaphyseal dysplasia (SMD) resembling SMD Sedaghatian type. <i>Journal of Medical Genetics</i> , <b>2007</b> , 44, e73	5.8	13
272	The genetics of osteoarthritis. <b>2007</b> , 3, 503-16		2
271	Visualising A Skeletal Dysplasia Knowledgebase. <b>2007</b> ,		
270	Significant ocular findings are a feature of heritable bone dysplasias resulting from defects in type II collagen. <b>2007</b> , 91, 1148-51		18
269	New gene associations in osteoarthritis: what do they provide, and where are we going?. <b>2007</b> , 19, 429-34		57
268	Ost�chondrodysplasies l�ales. <b>2007</b> , 2, 1-6		1
267	Osteogenesis imperfecta and holoprosencephaly. <b>2007</b> , 16, 189-191		
266	Chondrodysplasies responsables d'insuffisance staturale. <b>2007</b> , 2, 1-11		
265	Type XXVII collagen at the transition of cartilage to bone during skeletogenesis. <b>2007</b> , 41, 535-42		56
264	Achondroplasia. <b>2007</b> , 370, 162-172		366
263	Mutations in FAM20C are associated with lethal osteosclerotic bone dysplasia (Raine syndrome), highlighting a crucial molecule in bone development. <i>American Journal of Human Genetics</i> , <b>2007</b> , 81, 906-12	11	149
262	Angulated femurs and the skeletal dysplasias: experience of the International Skeletal Dysplasia Registry (1988-2006). <i>American Journal of Medical Genetics, Part A</i> , <b>2007</b> , 143A, 1159-68	2.5	30
261	A new case of spondyloenchondrodysplasia with immune dysregulation confirms the pleiotropic nature of the disorder: comment on "A syndrome of immunodeficiency, autoimmunity, and spondylometaphyseal dysplasia" by M.L. Kulkarni, K. Baskar, and P.M. Kulkarni [2006]. <i>American Journal of Medical Genetics, Part A</i> , <b>2007</b> , 143A, 1394-5	2.5	6
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258	Osseous dysplasia with severe short stature, multiple dislocations, and delayed bone age: report on a second Lebanese patient. <i>American Journal of Medical Genetics, Part A</i> , <b>2007</b> , 143A, 1782-7	2.5	4
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255	Acanthosis nigricans in a child with mild osteochondrodysplasia and K650Q mutation in the FGFR3 gene. <i>American Journal of Medical Genetics, Part A</i> , <b>2007</b> , 143A, 3144-9	2.5	18
254	An unclassifiable short rib-polydactyly syndrome with acromesomelic hypomineralization and campomelia in sibilings. <i>American Journal of Medical Genetics, Part A</i> , <b>2007</b> , 143A, 2607-11	2.5	10
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