Sylvie Odent

List of Publications by Year in Descending Order

10 papers	1,334 citations	9 h-index	11 g-index
11 ext. papers	1,507 ext. citations	13.2 avg, IF	2.53 L-index

#	Paper	IF	Citations
10	Incidence of cardiovascular events and risk markers in a prospective study of children diagnosed with Marfan syndrome. *Archives of Cardiovascular Diseases*, **2020**, 113, 40-49	2.7	5
9	International Registry of Patients Carrying TGFBR1 or TGFBR2 Mutations: Results of the MAC (Montalcino Aortic Consortium). *Circulation: Cardiovascular Genetics*, **2016**, 9, 548-558		105
8	Marfan Sartan: a randomized, double-blind, placebo-controlled trial. *European Heart Journal*, **2015**, 36, 2160-6	9.5	134
7	Orthopedics management of acromicric dysplasia: follow up of nine patients. *American Journal of Medical Genetics, Part A*, **2014**, 164A, 331-7	2.5	9
6	Mutations in the TGF□binding-protein-like domain 5 of FBN1 are responsible for acromicric and geleophysic dysplasias. *American Journal of Human Genetics*, **2011**, 89, 7-14	11	171
5	Rationale and design of a randomized clinical trial (Marfan Sartan) of angiotensin II receptor blocker therapy versus placebo in individuals with Marfan syndrome. *Archives of Cardiovascular Diseases*, **2010**, 103, 317-25	2.7	59
4	Comparison of clinical presentations and outcomes between patients with TGFBR2 and FBN1 mutations in Marfan syndrome and related disorders. *Circulation*, **2009**, 120, 2541-9	16.7	160
3	Mutations in the beta-tubulin gene TUBB2B result in asymmetrical polymicrogyria. *Nature Genetics*, **2009**, 41, 746-52	36.3	290
2	Large spectrum of lissencephaly and pachygyria phenotypes resulting from de novo missense mutations in tubulin alpha 1A (TUBA1A). *Human Mutation*, **2007**, 28, 1055-64	4.7	186
1	BBS10 encodes a vertebrate-specific chaperonin-like protein and is a major BBS locus. *Nature Genetics*, **2006**, 38, 521-4	36.3	214