Agnieszka Rzdzynska-Swiatkowska

List of Publications by Citations

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

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#	Paper	IF	Citations
16	Thyroid hormone resistance syndrome due to mutations in the thyroid hormone receptor Igene (THRA). <i>Journal of Medical Genetics</i> , 2015 , 52, 312-6	5.8	66
15	Polish 2012 growth references for preschool children. European Journal of Pediatrics, 2013, 172, 753-61	4.1	28
14	Mucopolysaccharidosis type II in females and response to enzyme replacement therapy. <i>American Journal of Medical Genetics, Part A</i> , 2012 , 158A, 450-4	2.5	22
13	Growth patterns in children with mucopolysaccharidosis I and II. World Journal of Pediatrics, 2015 , 11, 226-31	4.6	19
12	Mucopolysaccharidosis type VI in Russia, Kazakhstan, and Central and Eastern Europe. <i>Pediatrics International</i> , 2014 , 56, 520-5	1.2	19
11	Enzyme replacement therapy in an attenuated case of mucopolysaccharidosis type I (Scheie syndrome): a 6.5-year detailed follow-up. <i>Pediatric Neurology</i> , 2012 , 47, 461-5	2.9	16
10	Restricted joint range of motion in patients with MPS II: correlation with height, age and functional status. <i>Acta Paediatrica, International Journal of Paediatrics</i> , 2012 , 101, e183-8	3.1	13
9	The effect of recombinant human iduronate-2-sulfatase (Idursulfase) on growth in young patients with mucopolysaccharidosis type II. <i>PLoS ONE</i> , 2014 , 9, e85074	3.7	11
8	Can Macrosomia or Large for Gestational Age Be Predictive of Mucopolysaccharidosis Type I, II and VI?. <i>Pediatrics and Neonatology</i> , 2016 , 57, 181-7	1.8	7
7	Bioimpedance Analysis as a Method to Evaluate the Proportion of Fatty and Muscle Tissues in Progressive Myopathy in Pompe Disease. <i>JIMD Reports</i> , 2016 , 26, 45-51	1.9	5
6	Novel data on growth phenotype and causative genotypes in 29 patients with Morquio (Morquio-Brailsford) syndrome from Central-Eastern Europe. <i>Journal of Applied Genetics</i> , 2019 , 60, 163-	1774	4
5	Ultrasonographic Features of Hip Joints in Mucopolysaccharidoses Type I and II. <i>PLoS ONE</i> , 2015 , 10, e0123792	3.7	3
4	Mild phenotype of glutaric aciduria type 1 in polish patients - novel data from a group of 13 cases. <i>Metabolic Brain Disease</i> , 2019 , 34, 641-649	3.9	2
3	Bone metabolism in patients with mucopolysaccharidosis type II. <i>Reumatologia</i> , 2014 , 6, 354-361	1.7	1
2	Musculoskeletal manifestations of mucopolysaccharidosis type VI and effects of enzyme replacement therapy. <i>Open Medicine (Poland)</i> , 2012 , 7, 154-162	2.2	1
1	The importance of anthropological methods in the diagnosis of rare diseases. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2019 , 32, 311-320	1.6	