Christina Lampe

List of Publications by Year
in descending order

14 papers	474 citations	840776 11 h-index	996975 15 g-index
15 all docs	15 docs citations	15 times ranked	435 citing authors

#	Article	IF	Citations
1	Spinal involvement in mucopolysaccharidosis IVA (Morquioâ€Brailsford or Morquio A syndrome): presentation, diagnosis and management. Journal of Inherited Metabolic Disease, 2013, 36, 339-355.	3.6	100
2	Natural history and galsulfase treatment in mucopolysaccharidosis VI (MPS VI, Maroteauxâ€“Lamy) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 American Journal of Medical Genetics, Part A, 2014, 164, 1953-1964.	1.2	74
3	Design, baseline characteristics, and early findings of the MPS VI (mucopolysaccharidosis VI) Clinical Surveillance Program (CSP). Journal of Inherited Metabolic Disease, 2013, 36, 373-384.	3.6	57
4	Clinical characteristics of adults with slowly progressing mucopolysaccharidosis VI: a case series. Journal of Inherited Metabolic Disease, 2012, 35, 1071-1079.	3.6	46
5	A multinational, multidisciplinary consensus for the diagnosis and management of spinal cord compression among patients with mucopolysaccharidosis VI. Molecular Genetics and Metabolism, 2012, 107, 15-24.	1.1	39
6	Cervical cord compression in mucopolysaccharidosis VI (MPS VI): Findings from the MPS VI Clinical Surveillance Program (CSP). Molecular Genetics and Metabolism, 2016, 118, 310-318.	1.1	38
7	Development of a Scoring System to Evaluate the Severity of Craniocervical Spinal Cord Compression in Patients with Mucopolysaccharidosis IVA (Morquio A Syndrome). JIMD Reports, 2013, 11, 65-72.	1.5	27
8	Surgical management of neurological manifestations of mucopolysaccharidosis disorders. Molecular Genetics and Metabolism, 2017, 122, 41-48.	1.1	27
9	Craniocervical decompression in patients with mucopolysaccharidosis VI: development of a scoring system to determine indication and outcome of surgery. Journal of Inherited Metabolic Disease, 2013, 36, 1005-1013.	3.6	22
10	Long-Term Galsulfase Treatment Associated With Improved Survival of Patients With Mucopolysaccharidosis VI (Maroteaux-Lamy Syndrome). FIRE Forum for International Research in Education, 2018, 6, 232640981875580.	0.7	13
11	Stand-alone craniocervical decompression is feasible in children with mucopolysaccharidosis type I, IVA, and VI. Spine Journal, 2018, 18, 1455-1459.	1.3	12
12	Critical clinical situations in adult patients with Mucopolysaccharidoses (MPS). Orphanet Journal of Rare Diseases, 2020, 15, 114.	2.7	8
13	Understanding the Early Presentation of Mucopolysaccharidoses Disorders. FIRE Forum for International Research in Education, 2018, 6, 232640981880034.	0.7	7
14	Mucopolysaccharidoses and Orthopedic Management (Focused also on Craniocervical Junction). Journal of Child Science, 2018, 08, e128-e137.	0.2	2