Michael Watson

List of Publications by Year
in descending order

9 papers	1,779 citations	1307594 7 h-index	1474206 9 g-index
9 all docs	9 docs citations	9 times ranked	5102 citing authors

#	Article	IF	Citations
1	Maximizing the Benefit of Life-Saving Treatments for Pompe Disease, Spinal Muscular Atrophy, and Duchenne Muscular Dystrophy Through Newborn Screening. JAMA Neurology, 2019, 76, 978.	9.0	14
2	Current conditions in medical genetics practice. Genetics in Medicine, 2019, 21, 1874-1877.	2.4	103
3	Duchenne Muscular Dystrophy Newborn Screening, a Case Study for Examining Ethical and Legal Issues for Pilots for Emerging Disorders: Considerations and Recommendations. International Journal of Neonatal Screening, 2018, 4, 6.	3.2	6
4	ClinGen Variant Curation Expert Panel experiences and standardized processes for disease and geneâ€level specification of the ACMG/AMP guidelines for sequence variant interpretation. Human Mutation, 2018, 39, 1614-1622.	2.5	132
5	Unique aspects of sequence variant interpretation for inborn errors of metabolism (IEM): The ClinGen IEM Working Group and the Phenylalanine Hydroxylase Gene. Human Mutation, 2018, 39, 1569-1580.	2.5	50
6	Recommendations for reporting of secondary findings in clinical exome and genome sequencing, 2016 update (ACMG SF v2.0): a policy statement of the American College of Medical Genetics and Genomics. Genetics in Medicine, 2017, 19, 249-255.	2.4	1,398
7	The Human Variome Project. Human Mutation, 2016, 37, 505-507.	2.5	16
8	Determining barriers to effective data sharing in cancer genomic sequencing initiatives: A Global Alliance for Genomics and Health (GA4GH) survey.. Journal of Clinical Oncology, 2016, 34, 11502-11502.	1.6	3
9	Reporting genomic secondary findings: ACMG members weigh in. Genetics in Medicine, 2015, 17, 27-35.	2.4	57