Caroline Stanasiuk

List of Publications by Year
in descending order

8 papers	197 citations	1307594 7 h-index	1720034 7 g-index
8 all docs	8 docs citations	8 times ranked	235 citing authors

#	Article	IF	Citations
1	Restrictive Cardiomyopathy is Caused by a Novel Homozygous Desmin (DES) Mutation p.Y122H Leading to a Severe Filament Assembly Defect. Genes, 2019, 10, 918.	2.4	47
2	The Novel Desmin Variant p.Leu115Ile Is Associated With a Unique Form of Biventricular Arrhythmogenic Cardiomyopathy. Canadian Journal of Cardiology, 2021, 37, 857-866.	1.7	28
3	A homozygous DSC2 deletion associated with arrhythmogenic cardiomyopathy is caused by uniparental isodisomy. Journal of Molecular and Cellular Cardiology, 2020, 141, 17-29.	1.9	27
4	Noncompaction cardiomyopathy is caused by a novel inâ€frame desmin (DES) deletion mutation within the 1A coiledâ€coil rod segment leading to a severe filament assembly defect. Human Mutation, 2019, 40, 734-741.	2.5	26
5	The Desmin (DES) Mutation p.A337P Is Associated with Left-Ventricular Non-Compaction Cardiomyopathy. Genes, 2021, 12, 121.	2.4	26
6	Desminopathy: Novel Desmin Variants, a New Cardiac Phenotype, and Further Evidence for Secondary Mitochondrial Dysfunction. Journal of Clinical Medicine, 2020, 9, 937.	2.4	24
7	Hemi- and Homozygous Loss-of-Function Mutations in DSG2 (Desmoglein-2) Cause Recessive Arrhythmogenic Cardiomyopathy with an Early Onset. International Journal of Molecular Sciences, 2021, 22, 3786.	4.1	19
8	Back Cover, Volume 40, Issue 6. Human Mutation, 2019, 40, ii.	2.5	0