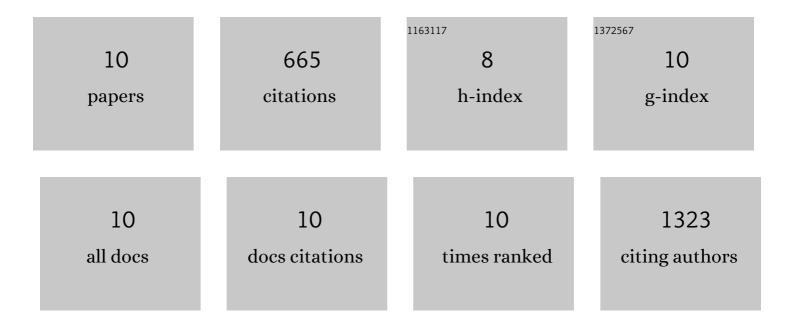
Rachel Morissette

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/3157373/publications.pdf Version: 2024-02-01



#	Article	IF	CITATIONS
1	sFRP2 in the aged microenvironment drives melanoma metastasis and therapy resistance. Nature, 2016, 532, 250-254.	27.8	290
2	Revisiting the prevalence of nonclassic congenital adrenal hyperplasia in US Ashkenazi Jews and Caucasians. Genetics in Medicine, 2017, 19, 1276-1279.	2.4	90
3	Clinical and biochemical profiles suggest fibromuscular dysplasia is a systemic disease with altered TGFâ€Î² expression and connective tissue features. FASEB Journal, 2014, 28, 3313-3324.	0.5	68
4	Tenascin-X Haploinsufficiency Associated with Ehlers-Danlos Syndrome in Patients with Congenital Adrenal Hyperplasia. Journal of Clinical Endocrinology and Metabolism, 2013, 98, E379-E387.	3.6	59
5	Broadening the Spectrum of Ehlers Danlos Syndrome in Patients With Congenital Adrenal Hyperplasia. Journal of Clinical Endocrinology and Metabolism, 2015, 100, E1143-E1152.	3.6	51
6	Transforming Growth Factor-β and Inflammation in Vascular (Type IV) Ehlers–Danlos Syndrome. Circulation: Cardiovascular Genetics, 2014, 7, 80-88.	5.1	45
7	Ehlers-Danlos Syndrome Caused by Biallelic <i>TNXB</i> Variants in Patients with Congenital Adrenal Hyperplasia. Human Mutation, 2016, 37, 893-897.	2.5	36
8	Transforming growth factor-β (TGF-β) pathway abnormalities in tenascin-X deficiency associated with CAH-X syndrome. European Journal of Medical Genetics, 2014, 57, 95-102.	1.3	16
9	The soluble domains of Gpi8 and Gaa1, two subunits of glycosylphosphatidylinositol transamidase (GPI-T), assemble into a complex. Archives of Biochemistry and Biophysics, 2017, 633, 58-67.	3.0	7
10	Defining the boundaries of species specificity for the Saccharomyces cerevisiae glycosylphosphatidylinositol transamidase using a quantitative inÂvivo assay. Bioscience Reports, 2012, 32, 577-586.	2.4	3