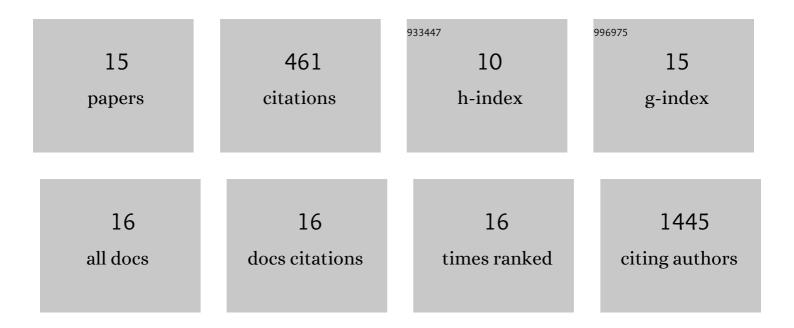
Juha W Koskenvuo

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/3499629/publications.pdf Version: 2024-02-01



#	Article	IF	CITATIONS
1	Genetics and genotype–phenotype correlations in Finnish patients with dilated cardiomyopathy. European Heart Journal, 2015, 36, 2327-2337.	2.2	130
2	Genetic Basis of Severe Childhood-OnsetÂCardiomyopathies. Journal of the American College of Cardiology, 2018, 72, 2324-2338.	2.8	97
3	Suppression of endothelial CD39/ENTPD1 is associated with pulmonary vascular remodeling in pulmonary arterial hypertension. American Journal of Physiology - Lung Cellular and Molecular Physiology, 2015, 308, L1046-L1057.	2.9	43
4	Loss of PPARÎ ³ in endothelial cells leads to impaired angiogenesis. Journal of Cell Science, 2016, 129, 693-705.	2.0	32
5	Relevance of Titin Missense and Non-Frameshifting Insertions/Deletions Variants in Dilated Cardiomyopathy. Scientific Reports, 2019, 9, 4093.	3.3	30
6	Clinical disease presentation and ECG characteristics of <i>LMNA</i> mutation carriers. Open Heart, 2017, 4, e000474.	2.3	26
7	Intramyocardial injection of SERCA2aâ€expressing lentivirus improves myocardial function in doxorubicinâ€induced heart failure. Journal of Gene Medicine, 2016, 18, 124-133.	2.8	18
8	Extracellular ATP protects endothelial cells against DNA damage. Purinergic Signalling, 2016, 12, 575-581.	2.2	18
9	Reference Values for Echocardiography in Middleâ€Aged Population: The Cardiovascular Risk in Young Finns Study. Echocardiography, 2016, 33, 193-206.	0.9	17
10	Case reports of two pedigrees with recessive arrhythmogenic right ventricular cardiomyopathy associated with homozygous Thr335Ala variant in DSG2. BMC Medical Genetics, 2017, 18, 86.	2.1	15
11	Pegylated and liposomal doxorubicin is associated with high mortality and causes limited cardiotoxicity in mice. BMC Research Notes, 2018, 11, 148.	1.4	9
12	Systemic Dosing of Thymosin Beta 4 before and after Ischemia Does Not Attenuate Global Myocardial Ischemia-Reperfusion Injury in Pigs. Frontiers in Pharmacology, 2016, 7, 115.	3.5	8
13	Biallelic loss-of-function in NRAP is a cause of recessive dilated cardiomyopathy. PLoS ONE, 2021, 16, e0245681.	2.5	8
14	Accurate genetic diagnosis of Finnish pulmonary arterial hypertension patients using oligonucleotideâ€selective sequencing. Molecular Genetics & Genomic Medicine, 2015, 3, 354-362.	1.2	5
15	Systemic Hypoxia Increases Circulating Concentration of Apelin in Humans. High Altitude Medicine and Biology, 2017, 18, 292-295.	0.9	5