RÃ³bert Sepp

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/9557733/publications.pdf

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19	3,656	933447 10	794594 19
papers	citations	h-index	g-index

19	19	19	5324
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	2014 ESC Guidelines on diagnosis and management of hypertrophic cardiomyopathy. European Heart Journal, 2014, 35, 2733-2779.	2.2	3,469
2	Different patterns of left ventricular rotational mechanics in cardiac amyloidosis-results from the three-dimensional speckle-tracking echocardiographic MAGYAR-Path Study. Quantitative Imaging in Medicine and Surgery, 2015, 5, 853-7.	2.0	24
3	Systemic antibiotic prophylaxis does not affect infectious complications in pediatric burn injury: A meta-analysis. PLoS ONE, 2019, 14, e0223063.	2.5	22
4	Direct Anticoagulants and Risk of Myocardial Infarction, a Multiple Treatment Network Meta-Analysis. Angiology, 2020, 71, 27-37.	1.8	17
5	Timothy syndrome 1 genotype without syndactyly and major extracardiac manifestations. American Journal of Medical Genetics, Part A, 2017, 173, 784-789.	1.2	15
6	Short-term beat-to-beat variability of the QT interval is increased and correlates with parameters of left ventricular hypertrophy in patients with hypertrophic cardiomyopathy. Canadian Journal of Physiology and Pharmacology, 2015, 93, 765-772.	1.4	14
7	A novel â€˜splice siteâ€™ HCN4 Gene mutation, c.1737 + 1 G > T, causes familial bradycardia, reduced heart rate response, impaired chronotropic competence and increased short-term heart rate variability. International Journal of Cardiology, 2017, 241, 364-372.	1.7	12
8	Double Stenting for Malignant Biliary and Duodenal Obstruction: A Systematic Review and Meta-Analysis. Clinical and Translational Gastroenterology, 2020, 11, e00161.	2.5	12
9	Identification of Two Novel LAMP2 Gene Mutations in Danon Disease. Canadian Journal of Cardiology, 2016, 32, 1355.e23-1355.e30.	1.7	11
10	Left atrial dysfunction in light-chain cardiac amyloidosis and hypertrophic cardiomyopathy â€“ A comparative three-dimensional speckle-tracking echocardiographic analysis from the MAGYAR-Path Study. Revista Portuguesa De Cardiologia, 2017, 36, 905-913.	0.5	11
11	Identification of a Novel GLA Gene Mutation, p.Ile239Met, in Fabry Disease With a Predominant Cardiac Phenotype. International Heart Journal, 2017, 58, 454-458.	1.0	11
12	Sandblasting reduces dental implant failure rate but not marginal bone level loss: A systematic review and meta-analysis. PLoS ONE, 2019, 14, e0216428.	2.5	8
13	Noninvasive ventilation improves the outcome in patients with pneumonia-associated respiratory failure: Systematic review and meta-analysis. Journal of Infection and Public Health, 2022, 15, 349-359.	4.1	8
14	Prognostic Value of Reduced Heart Rate Reserve during Exercise in Hypertrophic Cardiomyopathy. Journal of Clinical Medicine, 2021, 10, 1347.	2.4	6
15	Variant Transthyretin Amyloidosis (ATTRv) in Hungary: First Data on Epidemiology and Clinical Features. Genes, 2021, 12, 1152.	2.4	4
16	The Genetic Architecture of Hypertrophic Cardiomyopathy in Hungary: Analysis of 242 Patients with a Panel of 98 Genes. Diagnostics, 2022, 12, 1132.	2.6	4
17	Termination of Persistent Perimitral Atrial Flutter by Selective Contrast Injection IntoÂthe Vein of Marshall. JACC: Clinical Electrophysiology, 2015, 1, 596-597.	3.2	3
18	Identification and functional characterisation of a novel KCNJ2 mutation, Val302del, causing Andersenâ€“Tawil syndrome. Canadian Journal of Physiology and Pharmacology, 2015, 93, 569-575.	1.4	3

#	Article	IF	Citations
19	Impaired cytoplasmic domain interactions cause co-assembly defect and loss of function in the p.Glu293Lys KNCJ2 variant isolated from an Andersenâ€“Tawil syndrome patient. Cardiovascular Research, 2020, 117, 1923-1934.	3.8	2