Jacqueline Neubauer

List of Publications by Year in Descending Order

12 papers	141 citations	5 h-index	11 g-index
13 ext. papers	181 ext. citations	3 avg, IF	2.89 L-index

#	Paper	IF	Citations
12	Genetic variants in eleven central and peripheral chemoreceptor genes in sudden infant death syndrome.. *Pediatric Research*, **2022**,	3.2	1
11	Benefits and outcomes of a new multidisciplinary approach for the management and financing of sudden unexplained death cases in a forensic setting in Switzerland.. *Forensic Science International*, **2022**, 334, 111240	2.6	0
10	A collaborative exercise on DNA methylation-based age prediction and body fluid typing.. *Forensic Science International: Genetics*, **2021**, 57, 102656	4.3	1
9	Re-evaluation of single nucleotide variants and identification of structural variants in a cohort of 45 sudden unexplained death cases. *International Journal of Legal Medicine*, **2021**, 135, 1341-1349	3.1	4
8	Forensische DNA-Methylierungsanalyse. *Rechtsmedizin*, **2021**, 31, 202-216	0.6	1
7	Forensische DNA-Methylierungsanalyse. *Rechtsmedizin*, **2021**, 31, 192-201	0.6	2
6	Forensic transcriptome analysis using massively parallel sequencing. *Forensic Science International: Genetics*, **2021**, 52, 102486	4.3	9
5	Functional characterization of a novel SCN5A variant associated with long QT syndrome and sudden cardiac death. *International Journal of Legal Medicine*, **2019**, 133, 1733-1742	3.1	2
4	Exome analysis in 34 sudden unexplained death (SUD) victims mainly identified variants in channelopathy-associated genes. *International Journal of Legal Medicine*, **2018**, 132, 1057-1065	3.1	30
3	Sex-dependent differences in the in vivo respiratory phenotype of the TASK-1 potassium channel knockout mouse. *Respiratory Physiology and Neurobiology*, **2017**, 245, 13-28	2.8	6
2	Post-mortem whole-exome analysis in a large sudden infant death syndrome cohort with a focus on cardiovascular and metabolic genetic diseases. *European Journal of Human Genetics*, **2017**, 25, 404-409	5.3	66
1	Post-mortem whole-exome sequencing (WES) with a focus on cardiac disease-associated genes in five young sudden unexplained death (SUD) cases. *International Journal of Legal Medicine*, **2016**, 130, 1011-1021	3.1	18