

# Juan Diego RamÃ- rez

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/9242112/publications.pdf>

Version: 2024-02-01

11  
papers

1,177  
citations

933447

10  
h-index

1281871

11  
g-index

11  
all docs

11  
docs citations

11  
times ranked

1803  
citing authors

#	ARTICLE	IF	CITATIONS
1	A Gain-of-Function Mutation in TRPA1 Causes Familial Episodic Pain Syndrome. <i>Neuron</i> , 2010, 66, 671-680.	8.1	376
2	The Pain in Neuropathy Study (PiNS). <i>Pain</i> , 2016, 157, 1132-1145.	4.2	230
3	The clinical approach to small fibre neuropathy and painful channelopathy. <i>Practical Neurology</i> , 2014, 14, 368-379.	1.1	122
4	Rare Nav1.7 variants associated with painful diabetic peripheral neuropathy. <i>Pain</i> , 2018, 159, 469-480.	4.2	116
5	TRPA1 Gene Deletion Affects Somatosensory Neuron Function and Delays Mechanical Hypersensitivity in Response to Peripheral Nerve Damage. <i>Journal of Neuroscience</i> , 2013, 33, 16412-16426.	3.6	105
6	Sensory, psychological, and metabolic dysfunction in HIV-associated peripheral neuropathy: A cross-sectional deep profiling study. <i>Pain</i> , 2014, 155, 1846-1860.	4.2	87
7	Novel Mutations Mapping to the Fourth Sodium Channel Domain of Nav1.7 Result in Variable Clinical Manifestations of Primary Erythromelalgia. <i>NeuroMolecular Medicine</i> , 2013, 15, 265-278.	3.4	56
8	A Fovea for Pain at the Fingertips. <i>Current Biology</i> , 2013, 23, 496-500.	3.9	33
9	Imaging the neural correlates of neuropathic pain and pleasurable relief associated with inherited erythromelalgia in a single subject with quantitative arterial spin labelling. <i>Pain</i> , 2012, 153, 1122-1127.	4.2	29
10	Intermediate Charcot-Marie-Tooth disease due to a novel Trp101Stop myelin protein zero mutation associated with debilitating neuropathic pain. <i>Pain</i> , 2012, 153, 1763-1768.	4.2	16
11	Null mutation in <i>SCN9A</i> in which noxious stimuli can be detected in the absence of pain. <i>Neurology</i> , 2014, 83, 1577-1580.	1.1	7