

Yuanyuan Liu

List of Publications by Year in descending order

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12
papers

626
citations

1040056

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h-index

1281871

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17
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17
docs citations

17
times ranked

1006
citing authors

| # | ARTICLE | IF | CITATIONS |
|----|--|------|-----------|
| 1 | Impaired Action Potential Initiation in GABAergic Interneurons Causes Hyperexcitable Networks in an Epileptic Mouse Model Carrying a Human Na ^v 1.1 Mutation. <i>Journal of Neuroscience</i> , 2014, 34, 14874-14889. | 3.6 | 138 |
| 2 | Neuronal mechanisms of mutations in <i>SCN8A</i> causing epilepsy or intellectual disability. <i>Brain</i> , 2019, 142, 376-390. | 7.6 | 92 |
| 3 | CAPS Facilitates Filling of the Rapidly Releasable Pool of Large Dense-Core Vesicles. <i>Journal of Neuroscience</i> , 2008, 28, 5594-5601. | 3.6 | 75 |
| 4 | Genotype-phenotype correlations in <i>SCN8A</i> -related disorders reveal prognostic and therapeutic implications. <i>Brain</i> , 2022, 145, 2991-3009. | 7.6 | 69 |
| 5 | Synaptobrevin2 is the v-SNARE required for cytotoxic T-lymphocyte lytic granule fusion. <i>Nature Communications</i> , 2013, 4, 1439. | 12.8 | 65 |
| 6 | Two distinct secretory vesicle priming steps in adrenal chromaffin cells. <i>Journal of Cell Biology</i> , 2010, 190, 1067-1077. | 5.2 | 58 |
| 7 | The Coffin-Lowry syndrome-associated protein RSK2 is implicated in calcium-regulated exocytosis through the regulation of PLD1. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2008, 105, 8434-8439. | 7.1 | 50 |
| 8 | Relationship of electrophysiological dysfunction and clinical severity in <i>SCN2A</i> -related epilepsies. <i>Human Mutation</i> , 2018, 39, 1942-1956. | 2.5 | 29 |
| 9 | Activity of NaV1.2 promotes neurodegeneration in an animal model of multiple sclerosis. <i>JCI Insight</i> , 2016, 1, e89810. | 5.0 | 22 |
| 10 | Therapeutic Potential of Sodium Channel Blockers as a Targeted Therapy Approach in KCNA1-Associated Episodic Ataxia and a Comprehensive Review of the Literature. <i>Frontiers in Neurology</i> , 2021, 12, 703970. | 2.4 | 15 |
| 11 | A <i>SCN8A</i> variant associated with severe early onset epilepsy and developmental delay: Loss- or gain-of-function?. <i>Epilepsy Research</i> , 2021, 178, 106824. | 1.6 | 7 |
| 12 | OUP accepted manuscript. <i>Brain</i> , 2022, , . | 7.6 | 0 |