Toshiaki Okada

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/1985902/publications.pdf

Version: 2024-02-01

1307594 1720034 8 249 7 7 citations g-index h-index papers 8 8 8 296 docs citations times ranked citing authors all docs

| # | Article | IF | CITATIONS |
|---|--|------|-----------|
| 1 | Cell Volume-Activated and Volume-Correlated Anion Channels in Mammalian Cells: Their Biophysical, Molecular, and Pharmacological Properties. Pharmacological Reviews, 2019, 71, 49-88. | 16.0 | 61 |
| 2 | The organic anion transporter <scp>SLCO</scp> 2A1 constitutes the core component of the Maxiâ€Cl channel. EMBO Journal, 2017, 36, 3309-3324. | 7.8 | 46 |
| 3 | Maxi-anion channel and pannexin 1 hemichannel constitute separate pathways for swelling-induced ATP release in murine L929 fibrosarcoma cells. American Journal of Physiology - Cell Physiology, 2012, 303, C924-C935. | 4.6 | 38 |
| 4 | The properties, functions, and pathophysiology of maxi-anion channels. Pflugers Archiv European Journal of Physiology, 2016, 468, 405-420. | 2.8 | 38 |
| 5 | Specific and essential but not sufficient roles of LRRC8A in the activity of volume-sensitive outwardly rectifying anion channel (VSOR). Channels, 2017, 11, 109-120. | 2.8 | 30 |
| 6 | Molecular Identities and ATP Release Activities of Two Types of Volume-Regulatory Anion Channels, VSOR and Maxi-Cl. Current Topics in Membranes, 2018, 81, 125-176. | 0.9 | 27 |
| 7 | The ATP-Releasing Maxi-Cl Channel: Its Identity, Molecular Partners, and Physiological/Pathophysiological Implications. Life, 2021, 11, 509. | 2.4 | 9 |
| 8 | Current Ion Channel-targeted Drugs and Potential of Venom-derived Peptides as a Therapeutic New Modality. Venoms and Toxins, 2022, 2, . | 0.3 | 0 |