Yingmin Zhu

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/11749749/publications.pdf

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516710 642732 1,594 25 16 23 h-index citations g-index papers 26 26 26 2399 times ranked docs citations citing authors all docs

| # | Article | IF | CITATIONS |
|----|---|--------------|-----------|
| 1 | Abstract 930: Production of functionally active recombinant FOXA1: The first step towards targeted drug discovery., 2021,,. | | O |
| 2 | Conformational Changes of $ROR\hat{I}^3$ During Response Element Recognition and Coregulator Engagement. Journal of Molecular Biology, 2021, 433, 167258. | 4.2 | 4 |
| 3 | Acetylation of histone H3K27 signals the transcriptional elongation for estrogen receptor alpha. Communications Biology, 2020, 3, 165. | 4.4 | 26 |
| 4 | Refolding Proteins from Inclusion Bodies using Differential Scanning Fluorimetry Guided (DGR) Protein Refolding and MeltTraceur Web. Current Protocols in Molecular Biology, 2019, 125, e78. | 2.9 | 6 |
| 5 | Pyrazolo[1,5-a]pyrimidine TRPC6 antagonists for the treatment of gastric cancer. Cancer Letters, 2018, 432, 47-55. | 7.2 | 45 |
| 6 | Pyrazolopyrimidines as Potent Stimulators for Transient Receptor Potential Canonical 3/6/7 Channels. Journal of Medicinal Chemistry, 2017, 60, 4680-4692. | 6.4 | 44 |
| 7 | Identification of novel 2-(benzo[d]isoxazol-3-yl)-2-oxo-N-phenylacetohydrazonoyl cyanide analoguesas potent EPAC antagonists. European Journal of Medicinal Chemistry, 2017, 134, 62-71. | 5. 5 | 8 |
| 8 | Structure-activity relationships of 2-substituted phenyl- N -phenyl-2-oxoacetohydrazonoyl cyanides as novel antagonists of exchange proteins directly activated by cAMP (EPACs). Bioorganic and Medicinal Chemistry Letters, 2017, 27, 5163-5166. | 2.2 | 9 |
| 9 | Structure–Activity Relationship Studies with Tetrahydroquinoline Analogs as EPAC Inhibitors. ACS Medicinal Chemistry Letters, 2017, 8, 1183-1187. | 2.8 | 19 |
| 10 | A cell-based, quantitative and isoform-specific assay for exchange proteins directly activated by cAMP. Scientific Reports, 2017, 7, 6200. | 3.3 | 3 |
| 11 | Critical role for Epac1 in inflammatory pain controlled by GRK2-mediated phosphorylation of Epac1. Proceedings of the National Academy of Sciences of the United States of America, 2016, 113, 3036-3041. | 7.1 | 104 |
| 12 | Functionalized <i>N</i> , <i>N</i> -Diphenylamines as Potent and Selective EPAC2 Inhibitors. ACS Medicinal Chemistry Letters, 2016, 7, 460-464. | 2.8 | 17 |
| 13 | Organelle-specific Subunit Interactions of the Vertebrate Two-pore Channel Family. Journal of Biological Chemistry, 2015, 290, 1086-1095. | 3.4 | 24 |
| 14 | Biochemical and Pharmacological Characterizations of ESI-09 Based EPAC Inhibitors: Defining the ESI-09 "Therapeutic Window― Scientific Reports, 2015, 5, 9344. | 3.3 | 90 |
| 15 | Structure–Activity Relationship Studies of Substituted 2-(Isoxazol-3-yl)-2-oxo- <i>N</i> à€²-phenyl-acetohydrazonoyl Cyanide Analogues: Identification of Potent Exchange Proteins Directly Activated by cAMP (EPAC) Antagonists. Journal of Medicinal Chemistry, 2015, 58, 6033-6047. | 6.4 | 38 |
| 16 | Identification and optimization of 2â€aminobenzimidazole derivatives as novel inhibitors of <scp>TRPC</scp> 4 and <scp>TRPC</scp> 5 channels. British Journal of Pharmacology, 2015, 172, 3495-3509. | 5 . 4 | 38 |
| 17 | Rp-cAMPS Prodrugs Reveal the cAMP Dependence of First-Phase Glucose-Stimulated Insulin Secretion. Molecular Endocrinology, 2015, 29, 988-1005. | 3.7 | 32 |
| 18 | Biochemical and Pharmacological Characterizations of ESIâ€09 based EPAC inhibitors. FASEB Journal, 2015, 29, 1022.4. | 0.5 | 0 |

YINGMIN ZHU

| # | Article | IF | CITATION |
|----|--|------|----------|
| 19 | Dual depolarization responses generated within the same lateral septal neurons by TRPC4-containing channels. Pflugers Archiv European Journal of Physiology, 2014, 466, 1301-1316. | 2.8 | 21 |
| 20 | Depolarization potentials in mouse lateral septal nucleus neurons mediated by TRPC4â€like channels. FASEB Journal, 2012, 26, 1048.17. | 0.5 | 0 |
| 21 | Identification of ML204, a Novel Potent Antagonist That Selectively Modulates Native TRPC4/C5 Ion Channels. Journal of Biological Chemistry, 2011, 286, 33436-33446. | 3.4 | 171 |
| 22 | Cyclic Adenosine Diphosphate Ribose Activates Ryanodine Receptors, whereas NAADP Activates Two-pore Domain Channels. Journal of Biological Chemistry, 2011, 286, 9136-9140. | 3.4 | 78 |
| 23 | Activation of TRPA1 channels by fenamate nonsteroidal anti-inflammatory drugs. Pflugers Archiv European Journal of Physiology, 2010, 459, 579-592. | 2.8 | 110 |
| 24 | NAADP mobilizes calcium from acidic organelles through two-pore channels. Nature, 2009, 459, 596-600. | 27.8 | 687 |
| 25 | Expression of penicillin G acylase from the cloned pac gene of Escherichia coli ATCC11105. FEBS Journal, 2001, 268, 1298-1303. | 0.2 | 17 |