Tingyou Li

List of Publications by Year in descending order

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1125717 933410 14 263 10 13 citations h-index g-index papers 14 14 14 275 docs citations times ranked citing authors all docs

| # | Article | IF | CITATIONS |
|----|--|-----|-----------|
| 1 | Nitric oxide-donating and reactive oxygen species-responsive prochelators based on 8-hydroxyquinoline as anticancer agents. European Journal of Medicinal Chemistry, 2021, 212, 113153. | 5.5 | 13 |
| 2 | Neuroprotective Effect of <i>N</i> -Cyclohexylethyl-[A/G]-[D/E]-X-V Peptides on Ischemic Stroke by Blocking nNOS–CAPON Interaction. ACS Chemical Neuroscience, 2021, 12, 244-255. | 3.5 | 6 |
| 3 | Novel µ opioid antagonists derived from the µ opioid agonists endomorphin and [Dmt 1]DALDA (Hâ€Dmtâ€Dâ€Argâ€Pheâ€Lysâ€NH 2). Chemical Biology and Drug Design, 2020, 96, 1305-1314. | 3.2 | O |
| 4 | Intracerebroventricular administration of CYX-6, a potent $\hat{l}\frac{1}{4}$ -opioid receptor agonist, a \hat{l} - and \hat{l}° -opioid receptor antagonist and a biased ligand at $\hat{l}\frac{1}{4}$, \hat{l} & amp; \hat{l}° -opioid receptors, evokes antinociception with minimal constipation and respiratory depression in rats in contrast to morphine. European Journal of Pharmacology, 2020, 871, 172918. | 3.5 | 12 |
| 5 | Synthesis and Biological Evaluation of Fentanyl Analogues Modified at Phenyl Groups with Alkyls. ACS Chemical Neuroscience, 2019, 10, 201-208. | 3.5 | 8 |
| 6 | [Dmt1]DALDA analogues modified with tyrosine analogues at position 1. Bioorganic and Medicinal Chemistry Letters, 2016, 26, 3629-3631. | 2.2 | 6 |
| 7 | Endomorphin analogues with mixed \hat{l} /4-opioid (MOP) receptor agonism \hat{l} -opioid (DOP) receptor antagonism and lacking \hat{l}^2 -arrestin2 recruitment activity. Bioorganic and Medicinal Chemistry, 2014, 22, 2208-2219. | 3.0 | 12 |
| 8 | [Dmt1]DALDA analogues with enhanced \hat{l} 4 opioid agonist potency and with a mixed \hat{l} 4 \hat{l} 2 opioid activity profile. Bioorganic and Medicinal Chemistry, 2014, 22, 2333-2338. | 3.0 | 16 |
| 9 | Bifunctional [2â€~,6â€~-Dimethyl-l-tyrosine1]endomorphin-2 Analogues Substituted at Position 3 with Alkylated Phenylalanine Derivatives Yield Potent Mixed μ-Agonist/l´-Antagonist and Dual μ-Agonist/l´-Agonist Opioid Ligands. Journal of Medicinal Chemistry, 2007, 50, 2753-2766. | 6.4 | 39 |
| 10 | Transformation of $\hat{l}\frac{1}{4}$ -opioid receptor agonists into biologically potent $\hat{l}\frac{1}{4}$ -opioid receptor antagonists. Bioorganic and Medicinal Chemistry, 2007, 15, 1237-1251. | 3.0 | 18 |
| 11 | Enantioselective Synthesis of a Phenylalanine Library Containing Alkyl Groups on the Aromatic Moiety: Confirmation of Stereostructure by X-Ray Analysis. Chemical and Pharmaceutical Bulletin, 2006, 54, 873-877. | 1.3 | 16 |
| 12 | Potent in vivo antinociception and opioid receptor preference of the novel analogue [Dmt1]endomorphin-1. Pharmacology Biochemistry and Behavior, 2006, 84, 252-258. | 2.9 | 18 |
| 13 | Development of Potent ν-Opioid Receptor Ligands Using Unique Tyrosine Analogues of Endomorphin-2. Journal of Medicinal Chemistry, 2005, 48, 586-592. | 6.4 | 45 |
| 14 | Structural studies of $[2\hat{a}\in^2,6\hat{a}\in^2$ -dimethyl-l-tyrosine1]endomorphin-2 analogues: enhanced activity and cis orientation of the Dmt-Pro amide bond. Bioorganic and Medicinal Chemistry, 2003, 11, 1983-1994. | 3.0 | 54 |