Yan Shi

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/4352517/publications.pdf

Version: 2024-02-01

623734 642732 23 748 14 23 citations h-index g-index papers 23 23 23 794 docs citations citing authors all docs times ranked

#	Article	lF	Citations
1	Î ³ -AApeptides: Design, Structure, and Applications. Accounts of Chemical Research, 2016, 49, 428-441.	15.6	126
2	Helical Sulfono- \hat{l}^3 -AApeptides with Aggregation-Induced Emission and Circularly Polarized Luminescence. Journal of the American Chemical Society, 2019, 141, 12697-12706.	13.7	106
3	Stabilization of IncRNA GAS5 by a Small Molecule and Its Implications in Diabetic Adipocytes. Cell Chemical Biology, 2019, 26, 319-330.e6.	5.2	80
4	Inhibition of β-catenin/B cell lymphoma 9 proteinâ^'protein interaction using α-helix–mimicking sulfono-γ-AApeptide inhibitors. Proceedings of the National Academy of Sciences of the United States of America, 2019, 116, 10757-10762.	7.1	78
5	γâ€AApeptides as a New Class of Peptidomimetics. Chemistry - A European Journal, 2016, 22, 5458-5466.	3.3	52
6	Sulfono- \hat{I}^3 -AApeptides as Helical Mimetics: Crystal Structures and Applications. Accounts of Chemical Research, 2020, 53, 2425-2442.	15.6	51
7	α-Helix-Mimicking Sulfono-γ-AApeptide Inhibitors for p53–MDM2/MDMX Protein–Protein Interactions. Journal of Medicinal Chemistry, 2020, 63, 975-986.	6.4	43
8	The activity of sulfono- \hat{l}^3 -AApeptide helical foldamers that mimic GLP-1. Science Advances, 2020, 6, eaaz4988.	10.3	36
9	One-Bead–Two-Compound Thioether Bridged Macrocyclic γ-AApeptide Screening Library against EphA2. Journal of Medicinal Chemistry, 2017, 60, 9290-9298.	6.4	32
10	Aggregationâ€Induced Emissive and Circularly Polarized Homogeneous Sulfonoâ€Î³â€AApeptide Foldamers. Advanced Optical Materials, 2020, 8, 1902122.	7.3	24
11	Antimicrobial AApeptides. Current Topics in Medicinal Chemistry, 2017, 17, 1266-1279.	2.1	19
12	Rational Design and Synthesis of Right-Handed <scp>d</scp> -Sulfono-γ-AApeptide Helical Foldamers as Potent Inhibitors of Protein–Protein Interactions. Journal of Organic Chemistry, 2020, 85, 10552-10560.	3.2	16
13	Rational Design of Right-Handed Heterogeneous Peptidomimetics as Inhibitors of Protein–Protein Interactions. Journal of Medicinal Chemistry, 2020, 63, 13187-13196.	6.4	15
14	\hat{I}^3 -AApeptides as a New Strategy for Therapeutic Development. Current Medicinal Chemistry, 2019, 26, 2313-2329.	2.4	14
15	Polymyxin derivatives as broad-spectrum antibiotic agents. Chemical Communications, 2019, 55, 13104-13107.	4.1	10
16	Activation of E6AP/UBE3A-Mediated Protein Ubiquitination and Degradation Pathways by a Cyclic Î ³ -AA Peptide. Journal of Medicinal Chemistry, 2022, 65, 2497-2506.	6.4	10
17	$\hat{l}\pm/S$ ulfono- \hat{l}^3 -AApeptide Hybrid Analogues of Glucagon with Enhanced Stability and Prolonged In Vivo Activity. Journal of Medicinal Chemistry, 2021, 64, 13893-13901.	6.4	9
18	Dimeric lipo- \hat{l} ±/sulfono- \hat{l} 3-AA hybrid peptides as broad-spectrum antibiotic agents. Biomaterials Science, 2021, 9, 3410-3424.	5.4	8

#	Article	IF	CITATIONS
19	Discovery of a macrocyclic \hat{l}^3 -AApeptide binding to lncRNA GAS5 and its therapeutic implication in Type 2 diabetes. Future Medicinal Chemistry, 2019, 11, 2233-2235.	2.3	6
20	Dimeric \hat{I}^3 -AApeptides With Potent and Selective Antibacterial Activity. Frontiers in Chemistry, 2020, 8, 441.	3.6	6
21	 	4.1	5
22	The Activity of Small Ureaâ€Î³â€AApeptides Toward Gramâ€Positive Bacteria. ChemMedChem, 2019, 14, 1963-1	967.2	1
23	Discovery of α-helix-mimicking sulfono-γ-AApeptides as p53â^³MDM2 inhibitors. Future Medicinal Chemistry, 2021, 13, 1021-1023.	2.3	1