Tian-Miao Ou

List of Publications by Year in descending order

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201385 315357 2,161 38 27 38 h-index citations g-index papers 38 38 38 1865 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Rational design of small-molecules to recognize G-quadruplexes of c-MYC promoter and telomere and the evaluation of their <i>in vivo</i> antitumor activity against breast cancer. Nucleic Acids Research, 2022, 50, 1829-1848.	6.5	25
2	Design, Synthesis, and Evaluation of New Quinazolinone Derivatives that Inhibit Bloom Syndrome Protein (BLM) Helicase, Trigger DNA Damage at the Telomere Region, and Synergize with PARP Inhibitors. Journal of Medicinal Chemistry, 2020, 63, 9752-9772.	2.9	26
3	Developing Novel G-Quadruplex Ligands: from Interaction with Nucleic Acids to Interfering with Nucleic Acid–Protein Interaction. Molecules, 2019, 24, 396.	1.7	85
4	Probes and drugs that interfere with protein translation via targeting to the RNAs or RNA-protein interactions. Methods, 2019 , 167 , $124-133$.	1.9	5
5	MYC modulators in cancer: a patent review. Expert Opinion on Therapeutic Patents, 2019, 29, 353-367.	2.4	17
6	Discovery of Isaindigotone Derivatives as Novel Bloom's Syndrome Protein (BLM) Helicase Inhibitors That Disrupt the BLM/DNA Interactions and Regulate the Homologous Recombination Repair. Journal of Medicinal Chemistry, 2019, 62, 3147-3162.	2.9	32
7	Discovery of a New Four-Leaf Clover-Like Ligand as a Potent <i>c-MYC</i> Transcription Inhibitor Specifically Targeting the Promoter G-Quadruplex. Journal of Medicinal Chemistry, 2018, 61, 2447-2459.	2.9	86
8	Tracking the Dynamic Folding and Unfolding of RNA Gâ€Quadruplexes in Live Cells. Angewandte Chemie, 2018, 130, 4792-4796.	1.6	27
9	Discovery of Novel Schizocommunin Derivatives as Telomeric G-Quadruplex Ligands That Trigger Telomere Dysfunction and the Deoxyribonucleic Acid (DNA) Damage Response. Journal of Medicinal Chemistry, 2018, 61, 3436-3453.	2.9	33
10	Design, Synthesis, and Evaluation of Novel $\langle i \rangle p \langle i \rangle$ -(Methylthio)styryl Substituted Quindoline Derivatives as Neuroblastoma RAS (NRAS) Repressors via Specific Stabilizing the RNA G-Quadruplex. Journal of Medicinal Chemistry, 2018, 61, 6629-6646.	2.9	26
11	Design, Synthesis, and Evaluation of Isaindigotone Derivatives To Downregulate <i>c-myc</i> Transcription via Disrupting the Interaction of NM23-H2 with G-Quadruplex. Journal of Medicinal Chemistry, 2017, 60, 1292-1308.	2.9	40
12	Discovery of Novel 11-Triazole Substituted Benzofuro[3,2- <i>b</i>]quinolone Derivatives as <i>c-myc</i> G-Quadruplex Specific Stabilizers via Click Chemistry. Journal of Medicinal Chemistry, 2017, 60, 5407-5423.	2.9	68
13	Discovery of Small Molecules for Repressing Cap-Independent Translation of Human Vascular Endothelial Growth Factor ($h < VEGF < li>$) as Novel Antitumor Agents. Journal of Medicinal Chemistry, 2017, 60, 5306-5319.	2.9	16
14	New Disubstituted Quindoline Derivatives Inhibiting Burkitt's Lymphoma Cell Proliferation by Impeding <i>c-MYC</i> Transcription. Journal of Medicinal Chemistry, 2017, 60, 5438-5454.	2.9	46
15	Design, Synthesis, and Evaluation of New Selective NM23-H2 Binders as <i>c-MYC</i> Transcription Inhibitors via Disruption of the NM23-H2/G-Quadruplex Interaction. Journal of Medicinal Chemistry, 2017, 60, 6924-6941.	2.9	32
16	Specific targeting of telomeric multimeric G-quadruplexes by a new triaryl-substituted imidazole. Nucleic Acids Research, 2017, 45, 1606-1618.	6.5	86
17	Accurate high-throughput identification of parallel G-quadruplex topology by a new tetraaryl-substituted imidazole. Biosensors and Bioelectronics, 2016, 83, 77-84.	5.3	24
18	Synthesis and Mechanism Studies of 1,3-Benzoazolyl Substituted Pyrrolo[2,3- <i>b</i>)pyrazine Derivatives as Nonintercalative Topoisomerase II Catalytic Inhibitors. Journal of Medicinal Chemistry, 2016, 59, 238-252.	2.9	45

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19	A new application of click chemistry in situ: development of fluorescent probe for specific G-quadruplex topology. Scientific Reports, 2015, 5, 17202.	1.6	28
20	Chemical intervention of the NM23-H2 transcriptional programme on <i>c-MYC</i> via a novel small molecule. Nucleic Acids Research, 2015, 43, 6677-6691.	6.5	33
21	Stabilization of VEGF G-quadruplex and inhibition of angiogenesis by quindoline derivatives. Biochimica Et Biophysica Acta - General Subjects, 2014, 1840, 2970-2977.	1.1	34
22	Synthesis and evaluation of new BODIPY-benzofuroquinoline conjugates for sensitive and selective DNA detection. Dyes and Pigments, 2014, 107, 97-105.	2.0	9
23	Development of a new colorimetric and red-emitting fluorescent dual probe for G-quadruplex nucleic acids. Chemical Communications, 2014, 50, 6927-6930.	2.2	57
24	Discovery of a new fluorescent light-up probe specific to parallel G-quadruplexes. Chemical Communications, 2014, 50, 12173-12176.	2.2	48
25	New quinazoline derivatives for telomeric G-quadruplex DNA: Effects of an added phenyl group on quadruplex binding ability. European Journal of Medicinal Chemistry, 2013, 63, 1-13.	2.6	24
26	Benzofuroquinoline Derivatives Had Remarkable Improvement of their Selectivity for Telomeric G-Quadruplex DNA over Duplex DNA upon Introduction of Peptidyl Group. Bioconjugate Chemistry, 2012, 23, 1821-1831.	1.8	20
27	Design, synthesis and evaluation of isaindigotone derivatives as dual inhibitors for acetylcholinesterase and amyloid beta aggregation. Bioorganic and Medicinal Chemistry, 2012, 20, 2527-2534.	1.4	47
28	Disubstituted quinazoline derivatives as a new type of highly selective ligands for telomeric G-quadruplex DNA. European Journal of Medicinal Chemistry, 2012, 47, 299-311.	2.6	42
29	Inhibition of Cell Proliferation by Quindoline Derivative (SYUIQ-05) through its Preferential Interaction with <i>c</i> - <i>myc</i> Promoter G-Quadruplex. Journal of Medicinal Chemistry, 2011, 54, 5671-5679.	2.9	102
30	Impact of planarity of unfused aromatic molecules on G-quadruplex binding: Learning from isaindigotone derivatives. Organic and Biomolecular Chemistry, 2011, 9, 6422.	1.5	34
31	Pharmacophore-based discovery of triaryl-substituted imidazole as new telomeric G-quadruplex ligand. Bioorganic and Medicinal Chemistry Letters, 2011, 21, 1004-1009.	1.0	41
32	Quinolino-benzo-[5, 6]-dihydroisoquindolium compounds derived from berberine: A new class of highly selective ligands for G-quadruplex DNA in c-myc oncogene. European Journal of Medicinal Chemistry, 2011, 46, 1906-1913.	2.6	46
33	Turning off Transcription of the <i>bcl-2</i> Gene by Stabilizing the <i>bcl-2</i> Promoter Quadruplex with Quindoline Derivatives. Journal of Medicinal Chemistry, 2010, 53, 4390-4398.	2.9	117
34	Isaindigotone Derivatives: A New Class of Highly Selective Ligands for Telomeric G-Quadruplex DNA. Journal of Medicinal Chemistry, 2009, 52, 2825-2835.	2.9	87
35	9-N-Substituted berberine derivatives: Stabilization of G-quadruplex DNA and down-regulation of oncogene c-myc. Bioorganic and Medicinal Chemistry, 2008, 16, 7582-7591.	1.4	112
36	5- <i>N</i> -Methylated Quindoline Derivatives as Telomeric G-Quadruplex Stabilizing Ligands: Effects of 5- <i>N</i> Positive Charge on Quadruplex Binding Affinity and Cell Proliferation. Journal of Medicinal Chemistry, 2008, 51, 6381-6392.	2.9	123

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#	Article	IF	CITATION
37	Stabilization of G-Quadruplex DNA and Down-Regulation of Oncogenec-mycby Quindoline Derivatives. Journal of Medicinal Chemistry, 2007, 50, 1465-1474.	2.9	273
38	Synthesis and Evaluation of Quindoline Derivatives as G-Quadruplex Inducing and Stabilizing Ligands and Potential Inhibitors of Telomerase. Journal of Medicinal Chemistry, 2005, 48, 7315-7321.	2.9	165