

Hiroyuki Seimiya

List of Publications by Year in descending order

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Version: 2024-02-01

99
papers

4,635
citations

126907

33
h-index

102487

66
g-index

101
all docs

101
docs citations

101
times ranked

6790
citing authors

#	ARTICLE	IF	CITATIONS
1	Role of EMT in the DNA damage response, double-strand break repair pathway choice and its implications in cancer treatment. <i>Cancer Science</i> , 2022, , .	3.9	4
2	Design and synthesis of 14 and 15-membered macrocyclic scaffolds exhibiting inhibitory activities of hypoxia-inducible factor 1 α . <i>Bioorganic and Medicinal Chemistry</i> , 2021, 30, 115949.	3.0	3
3	G-quadruplex-forming nucleic acids interact with splicing factor 3B subunit 2 and suppress innate immune gene expression. <i>Genes To Cells</i> , 2021, 26, 65-82.	1.2	10
4	Serum IL-8 level as a candidate prognostic marker of response to anti-angiogenic therapy for metastatic colorectal cancer. <i>International Journal of Colorectal Disease</i> , 2021, 36, 131-139.	2.2	3
5	Novel tankyrase inhibitors suppress TDP-43 aggregate formation. <i>Biochemical and Biophysical Research Communications</i> , 2021, 537, 85-92.	2.1	4
6	Lamellarin 14, a derivative of marine alkaloids, inhibits the T790M/C797S mutant epidermal growth factor receptor. <i>Cancer Science</i> , 2021, 112, 1963-1974.	3.9	13
7	Neutralization of the induced VEGF-A potentiates the therapeutic effect of an anti-VEGFR2 antibody on gastric cancer in vivo. <i>Scientific Reports</i> , 2021, 11, 15125.	3.3	8
8	Chemical targeting of G-quadruplexes in telomeres and beyond for molecular cancer therapeutics. <i>Journal of Antibiotics</i> , 2021, 74, 617-628.	2.0	10
9	Pericentromeric noncoding RNA changes DNA binding of CTCF and inflammatory gene expression in senescence and cancer. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2021, 118, .	7.1	38
10	A phase I study to determine the maximum tolerated dose of trifluridine/tipiracil and oxaliplatin in patients with refractory metastatic colorectal cancer: LUPIN study. <i>Investigational New Drugs</i> , 2020, 38, 111-119.	2.6	6
11	Tankyrase promotes primary precursor miRNA processing to precursor miRNA. <i>Biochemical and Biophysical Research Communications</i> , 2020, 522, 945-951.	2.1	3
12	Target identification of a macrocyclic hexaoxazole G-quadruplex ligand using post-target-binding visualization. <i>Chemical Communications</i> , 2020, 56, 12905-12908.	4.1	17
13	Crossroads of telomere biology and anticancer drug discovery. <i>Cancer Science</i> , 2020, 111, 3089-3099.	3.9	28
14	Design and Discovery of an Orally Efficacious Spiroindolinone-Based Tankyrase Inhibitor for the Treatment of Colon Cancer. <i>Journal of Medicinal Chemistry</i> , 2020, 63, 4183-4204.	6.4	25
15	A fully synthetic 6-aza-artemisinin bearing an amphiphilic chain generates aggregates and exhibits anti-cancer activities. <i>Organic and Biomolecular Chemistry</i> , 2020, 18, 5339-5343.	2.8	10
16	Epidermal growth factor receptor mRNA expression: A potential molecular escape mechanism from regorafenib. <i>Cancer Science</i> , 2020, 111, 441-450.	3.9	8
17	ALDH1A3-mTOR axis as a therapeutic target for anticancer drug-tolerant persister cells in gastric cancer. <i>Cancer Science</i> , 2020, 111, 962-973.	3.9	36
18	Tankyrase Inhibitors Target Colorectal Cancer Stem Cells via AXIN-Dependent Downregulation of c-KIT Tyrosine Kinase. <i>Molecular Cancer Therapeutics</i> , 2020, 19, 765-776.	4.1	14

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19	G-quadruplex in cancer biology and drug discovery. <i>Biochemical and Biophysical Research Communications</i> , 2020, 531, 45-50.	2.1	48
20	c-KIT regulates stability of cancer stemness in CD44-positive colorectal cancer cells. <i>Biochemical and Biophysical Research Communications</i> , 2020, 527, 1014-1020.	2.1	7
21	In silico chemical screening identifies epidermal growth factor receptor as a therapeutic target of drug-tolerant CD44v9-positive gastric cancer cells. <i>British Journal of Cancer</i> , 2019, 121, 846-856.	6.4	13
22	Revisiting Telomere Shortening in Cancer. <i>Cells</i> , 2019, 8, 107.	4.1	98
23	Discovery of Novel Spiroindoline Derivatives as Selective Tankyrase Inhibitors. <i>Journal of Medicinal Chemistry</i> , 2019, 62, 3407-3427.	6.4	43
24	From the wings to the center stage of chromosomes. <i>Journal of Biological Chemistry</i> , 2019, 294, 17723-17724.	3.4	3
25	Calpain-10 regulates actin dynamics by proteolysis of microtubule-associated protein 1B. <i>Scientific Reports</i> , 2018, 8, 16756.	3.3	10
26	Cell-based chemical fingerprinting identifies telomeres and lamin A as modifiers of DNA damage response in cancer cells. <i>Scientific Reports</i> , 2018, 8, 14827.	3.3	17
27	<sc>RK</sc>â€287107, a potent and specific tankyrase inhibitor, blocks colorectal cancer cell growth in a preclinical model. <i>Cancer Science</i> , 2018, 109, 4003-4014.	3.9	60
28	Development of G-quadruplex ligands for selective induction of a parallel-type topology. <i>Organic and Biomolecular Chemistry</i> , 2018, 16, 7375-7382.	2.8	18
29	MERIT40-dependent recruitment of tankyrase to damaged DNA and its implication for cell sensitivity to DNA-damaging anticancer drugs. <i>Oncotarget</i> , 2018, 9, 35844-35855.	1.8	15
30	Tankyrase-Binding Protein TNKS1BP1 Regulates Actin Cytoskeleton Rearrangement and Cancer Cell Invasion. <i>Cancer Research</i> , 2017, 77, 2328-2338.	0.9	33
31	<i>APC</i> Mutations as a Potential Biomarker for Sensitivity to Tankyrase Inhibitors in Colorectal Cancer. <i>Molecular Cancer Therapeutics</i> , 2017, 16, 752-762.	4.1	67
32	Targeting glioma stem cells in vivo by a G-quadruplex-stabilizing synthetic macrocyclic hexaoxazole. <i>Scientific Reports</i> , 2017, 7, 3605.	3.3	40
33	mTOR signaling mediates resistance to tankyrase inhibitors in Wnt-driven colorectal cancer. <i>Oncotarget</i> , 2017, 8, 47902-47915.	1.8	20
34	Tankyrases. , 2017, , 4446-4449.		0
35	Serum VEGF-A and CCL5 levels as candidate biomarkers for efficacy and toxicity of regorafenib in patients with metastatic colorectal cancer. <i>Oncotarget</i> , 2016, 7, 34811-34823.	1.8	43
36	Recent advances in telomere biology for new cancer medicine. <i>Annals of Oncology</i> , 2016, 27, vii70.	1.2	0

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37	Design, Synthesis and Evaluation of an L-Dopa-Derived Macrocyclic Hexaoxazole (6otd) as a G-Quadruplex-Selective Ligand. <i>Heterocycles</i> , 2016, 92, 305.	0.7	3
38	Report on the use of non-clinical studies in the regulatory evaluation of oncology drugs. <i>Cancer Science</i> , 2016, 107, 189-202.	3.9	6
39	Senescence from glioma stem cell differentiation promotes tumor growth. <i>Biochemical and Biophysical Research Communications</i> , 2016, 470, 275-281.	2.1	24
40	G-quadruplex ligand-induced DNA damage response coupled with telomere dysfunction and replication stress in glioma stem cells. <i>Biochemical and Biophysical Research Communications</i> , 2016, 471, 75-81.	2.1	30
41	Design and Synthesis of a Berberine Dimer: A Fluorescent Ligand with High Affinity towards G-Quadruplexes. <i>Chemistry - A European Journal</i> , 2015, 21, 14519-14528.	3.3	28
42	Comprehensive transcriptomic analysis of molecularly targeted drugs in cancer for target pathway evaluation. <i>Cancer Science</i> , 2015, 106, 909-920.	3.9	18
43	Telomeric repeat-containing RNA/G-quadruplex-forming sequences cause genome-wide alteration of gene expression in human cancer cells in vivo. <i>Nucleic Acids Research</i> , 2015, 43, 2022-2032.	14.5	62
44	Predicting Risk at the End of the End: Telomere G-tail as a Biomarker. <i>EBioMedicine</i> , 2015, 2, 804-805.	6.1	4
45	Reprogramming Suppresses Premature Senescence Phenotypes of Werner Syndrome Cells and Maintains Chromosomal Stability over Long-Term Culture. <i>PLoS ONE</i> , 2014, 9, e112900.	2.5	52
46	TRF1 Ensures the Centromeric Function of Aurora-B and Proper Chromosome Segregation. <i>Molecular and Cellular Biology</i> , 2014, 34, 2464-2478.	2.3	29
47	Inhibition of ATP citrate lyase induces triglyceride accumulation with altered fatty acid composition in cancer cells. <i>International Journal of Cancer</i> , 2014, 135, 37-47.	5.1	52
48	TRIB1 Supports Prostate Tumorigenesis and Tumor-Propagating Cell Survival by Regulation of Endoplasmic Reticulum Chaperone Expression. <i>Cancer Research</i> , 2014, 74, 4888-4897.	0.9	53
49	Fission Yeast Pot1 and RecQ Helicase Are Required for Efficient Chromosome Segregation. <i>Molecular and Cellular Biology</i> , 2014, 34, 2551-2552.	2.3	0
50	Tankyrases. , 2014, , 1-4.		0
51	Interaction of long telomeric DNAs with macrocyclic hexaoxazole as a G-quadruplex ligand. <i>MedChemComm</i> , 2013, 4, 260-264.	3.4	7
52	Inhibition of ATP Citrate Lyase Induces an Anticancer Effect via Reactive Oxygen Species. <i>American Journal of Pathology</i> , 2013, 182, 1800-1810.	3.8	44
53	Telomere Length Influences Cancer Cell Differentiation <i>In Vivo</i> . <i>Molecular and Cellular Biology</i> , 2013, 33, 2988-2995.	2.3	45
54	Development of a gene expression database and related analysis programs for evaluation of anticancer compounds. <i>Cancer Science</i> , 2013, 104, 360-368.	3.9	9

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73	Therapeutic Targets and Drugs III: Tankyrase 1, Telomere-Binding Proteins, and Inhibitors. , 2009, , 281-291.		0
74	Telomere elongation by a mutant tankyrase 1 without TRF1 poly(ADP-ribosyl)ation. Experimental Cell Research, 2008, 314, 1115-1124.	2.6	9
75	ATP Citrate Lyase: Activation and Therapeutic Implications in Non-“Small Cell Lung Cancer. Cancer Research, 2008, 68, 8547-8554.	0.9	326
76	HnRNP A3 binds to and protects mammalian telomeric repeats in vitro. Biochemical and Biophysical Research Communications, 2007, 358, 608-614.	2.1	27
77	Cross-species difference in telomeric function of tankyrase 1. Cancer Science, 2007, 98, 850-857.	3.9	28
78	Evaluation of Tankyrase Inhibition in Whole Cells. Methods in Molecular Biology, 2007, 405, 133-146.	0.9	7
79	The telomeric PARP, tankyrases, as targets for cancer therapy. British Journal of Cancer, 2006, 94, 341-345.	6.4	88
80	Cancer therapy targeting the telomere maintenance system. Drug Delivery System, 2006, 21, 24-31.	0.0	2
81	Tankyrase 1 as a target for telomere-directed molecular cancer therapeutics. Cancer Cell, 2005, 7, 25-37.	16.8	160
82	Protein tyrosine phosphatase receptor-type O (PTPRO) exhibits characteristics of a candidate tumor suppressor in human lung cancer. Proceedings of the National Academy of Sciences of the United States of America, 2004, 101, 13844-13849.	7.1	102
83	Functional Subdomain in the Ankyrin Domain of Tankyrase 1 Required for Poly(ADP-Ribosyl)ation of TRF1 and Telomere Elongation. Molecular and Cellular Biology, 2004, 24, 1944-1955.	2.3	83
84	The Telomeric Poly(ADP-ribose) Polymerase, Tankyrase 1, Contains Multiple Binding Sites for Telomeric Repeat Binding Factor 1 (TRF1) and a Novel Acceptor, 182-kDa Tankyrase-binding Protein (TAB182). Journal of Biological Chemistry, 2002, 277, 14116-14126.	3.4	129
85	Telomere shortening and growth inhibition of human cancer cells by novel synthetic telomerase inhibitors MST-312, MST-295, and MST-1991. Molecular Cancer Therapeutics, 2002, 1, 657-65.	4.1	99
86	Pim-1 Negatively Regulates the Activity of PTP-U2S Phosphatase and Influences Terminal Differentiation and Apoptosis of Monoblastoid Leukemia Cells. Archives of Biochemistry and Biophysics, 2001, 390, 9-18.	3.0	41
87	ASK1 mediates apoptotic cell death induced by genotoxic stress. Oncogene, 1999, 18, 173-180.	5.9	169
88	Activation of c-Abl tyrosine kinase requires caspase activation and is not involved in JNK/SAPK activation during apoptosis of human monocytic leukemia U937 cells. Oncogene, 1999, 18, 1277-1283.	5.9	30
89	Hypoxia Up-Regulates Telomerase Activity via Mitogen-Activated Protein Kinase Signaling in Human Solid Tumor Cells. Biochemical and Biophysical Research Communications, 1999, 260, 365-370.	2.1	90
90	Apoptosis resistance in tumor cells. Cytotechnology, 1998, 27, 293-308.	1.6	5

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91	2-deoxyglucose inhibits chemotherapeutic drug-induced apoptosis in human monocytic leukemia U937 cells with inhibition of c-Jun N-terminal kinase 1/stress-activated protein kinase activation. , 1998, 76, 86-90.		16
92	Telomerase Inhibition, Telomere Shortening, and Senescence of Cancer Cells by Tea Catechins. Biochemical and Biophysical Research Communications, 1998, 249, 391-396.	2.1	226
93	Functional Involvement of PTP-U2L in Apoptosis Subsequent to Terminal Differentiation of Monoblastoid Leukemia Cells. Journal of Biological Chemistry, 1998, 273, 21187-21193.	3.4	20
94	Apoptosis resistance in tumor cells. , 1998, , 293-308.		0
95	c-Jun NH2-terminal Kinase-mediated Activation of Interleukin-1 β Converting Enzyme/CED-3-like Protease during Anticancer Drug-induced Apoptosis. Journal of Biological Chemistry, 1997, 272, 4631-4636.	3.4	182
96	Inhibition of the Association with Nuclear Matrix of pRB, p70 and p40 Proteins Along with the Specific Suppression of c-MYC Expression by Geldanamycin, an Inhibitor of Src Tyrosine Kinase.. Journal of Antibiotics, 1995, 48, 1021-1026.	2.0	20
97	M-CSF gene transduction in multidrug-resistant human cancer cells to enhance anti-P-glycoprotein antibody-dependent macrophage-mediated cytotoxicity. International Journal of Cancer, 1993, 54, 851-857.	5.1	20
98	T Cell Receptor-Extracellular Constant Regions as Hetero-Cross-Linkers for Immunoglobulin Variable Regions. Journal of Biochemistry, 1993, 113, 687-691.	1.7	4
99	Mouse-Human Chimeric Antibody MH171 against the Multidrug Transporter P-glycoprotein. Japanese Journal of Cancer Research, 1992, 83, 515-521.	1.7	11