

Marco Folini

List of Publications by Year in descending order

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Version: 2024-02-01

83
papers

8,528
citations

94433

37
h-index

76900

74
g-index

84
all docs

84
docs citations

84
times ranked

18974
citing authors

#	ARTICLE	IF	CITATIONS
1	Telomere as a Therapeutic Target in Dedifferentiated Liposarcoma. <i>Cancers</i> , 2022, 14, 2624.	3.7	1
2	On the Road to Fight Cancer: The Potential of G-Quadruplex Ligands as Novel Therapeutic Agents. <i>International Journal of Molecular Sciences</i> , 2021, 22, 5947.	4.1	45
3	The Role of Alternative Lengthening of Telomeres Mechanism in Cancer: Translational and Therapeutic Implications. <i>Cancers</i> , 2020, 12, 949.	3.7	29
4	Comparative Assessment of Antitumor Effects and Autophagy Induction as a Resistance Mechanism by Cytotoxics and EZH2 Inhibition in INI1-Negative Epithelioid Sarcoma Patient-Derived Xenograft. <i>Cancers</i> , 2019, 11, 1015.	3.7	21
5	The Oncogenic Signaling Pathways in BRAF-Mutant Melanoma Cells are Modulated by Naphthalene Diimide-Like G-Quadruplex Ligands. <i>Cells</i> , 2019, 8, 1274.	4.1	12
6	Distinct biological responses of metastatic castration resistant prostate cancer cells upon exposure to G-quadruplex interacting naphthalenediimide derivatives. <i>European Journal of Medicinal Chemistry</i> , 2019, 177, 401-413.	5.5	16
7	Luminescent dinuclear rhenium(II) PNA conjugates for microRNA-21 targeting: Synthesis, chemico-physical and biological characterization. <i>Journal of Organometallic Chemistry</i> , 2019, 887, 32-39.	1.8	7
8	miR-205 enhances radiation sensitivity of prostate cancer cells by impairing DNA damage repair through PKC μ and ZEB1 inhibition. <i>Journal of Experimental and Clinical Cancer Research</i> , 2019, 38, 51.	8.6	64
9	Down-Regulation of the Androgen Receptor by G-Quadruplex Ligands Sensitizes Castration-Resistant Prostate Cancer Cells to Enzalutamide. <i>Journal of Medicinal Chemistry</i> , 2018, 61, 8625-8638.	6.4	28
10	Naphthalene diimide derivatives G-quadruplex ligands induce cell proliferation inhibition, mild telomeric dysfunction and cell cycle perturbation in U251MG glioma cells. <i>FEBS Journal</i> , 2018, 285, 3769-3785.	4.7	21
11	CPAM type 2-derived mesenchymal stem cells: Malignancy risk study in a 14-month-old boy. <i>Pediatric Pulmonology</i> , 2017, 52, 990-999.	2.0	8
12	Synthesis and Superpotent Anticancer Activity of Tubulysins Carrying Non-hydrolysable N-Substituents on Tubuvaline. <i>Chemistry - A European Journal</i> , 2017, 23, 5842-5850.	3.3	9
13	miR-380-5p-mediated repression of TEP1 and TSPYL5 interferes with telomerase activity and favours the emergence of an "ALT-like" phenotype in diffuse malignant peritoneal mesothelioma cells. <i>Journal of Hematology and Oncology</i> , 2017, 10, 140.	17.0	23
14	Emerging Role of G-quadruplex DNA as Target in Anticancer Therapy. <i>Current Pharmaceutical Design</i> , 2017, 22, 6612-6624.	1.9	67
15	Guidelines for the use and interpretation of assays for monitoring autophagy (3rd edition). <i>Autophagy</i> , 2016, 12, 1-222.	9.1	4,701
16	Targeting of <i>RET</i> oncogene by naphthalene diimide-mediated gene promoter G-quadruplex stabilization exerts anti-tumor activity in oncogene-addicted human medullary thyroid cancer. <i>Oncotarget</i> , 2016, 7, 49649-49663.	1.8	22
17	miR-342 overexpression results in a synthetic lethal phenotype in <i>BRCA1</i> -mutant HCC1937 breast cancer cells. <i>Oncotarget</i> , 2016, 7, 18594-18604.	1.8	20
18	MicroRNAs and the Response of Prostate Cancer to Anti-Cancer Drugs. <i>Current Drug Targets</i> , 2016, 17, 257-265.	2.1	5

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19	Assessment of gene promoter G-quadruplex binding and modulation by a naphthalene diimide derivative in tumor cells. <i>International Journal of Oncology</i> , 2015, 46, 369-380.	3.3	28
20	Unravelling off-target effects of redox-active polymers and polymer multilayered capsules in prostate cancer cells. <i>Nanoscale</i> , 2015, 7, 6261-6270.	5.6	9
21	Water-soluble isoindolo[2,1-a]quinoxalin-6-imines: In vitro antiproliferative activity and molecular mechanism(s) of action. <i>European Journal of Medicinal Chemistry</i> , 2015, 94, 149-162.	5.5	51
22	Redox-Sensitive PEG-Polypeptide Nanoporous Particles for Survivin Silencing in Prostate Cancer Cells. <i>Biomacromolecules</i> , 2015, 16, 2168-2178.	5.4	38
23	Double stranded promoter region of BRAF undergoes to structural rearrangement in nearly physiological conditions. <i>FEBS Letters</i> , 2015, 589, 2117-2123.	2.8	11
24	Naphthalene diimides as red fluorescent pH sensors for functional cell imaging. <i>Organic and Biomolecular Chemistry</i> , 2015, 13, 570-576.	2.8	54
25	Editorial (Thematic Issue: Targeting Telomere Maintenance Mechanisms in Cancer Therapy). <i>Current Pharmaceutical Design</i> , 2014, 20, 6359-6360.	1.9	1
26	Synergistic Cooperation Between Sunitinib and Cisplatin Promotes Apoptotic Cell Death in Human Medullary Thyroid Cancer. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2014, 99, 498-509.	3.6	23
27	Senescent stroma promotes prostate cancer progression: The role of miR-210. <i>Molecular Oncology</i> , 2014, 8, 1729-1746.	4.6	102
28	miR-205 impairs the autophagic flux and enhances cisplatin cytotoxicity in castration-resistant prostate cancer cells. <i>Biochemical Pharmacology</i> , 2014, 87, 579-597.	4.4	83
29	MicroRNA-dependent Regulation of Telomere Maintenance Mechanisms: A Field as Much Unexplored as Potentially Promising. <i>Current Pharmaceutical Design</i> , 2014, 20, 6404-6421.	1.9	14
30	Nestling telomere length does not predict longevity, but covaries with adult body size in wild barn swallows. <i>Biology Letters</i> , 2013, 9, 20130340.	2.3	30
31	Targeting Loop Adenines in G-Quadruplex by a Selective Oxirane. <i>Chemistry - A European Journal</i> , 2013, 19, 78-81.	3.3	77
32	G-Quadruplex Structures in the Human Genome as Novel Therapeutic Targets. <i>Molecules</i> , 2013, 18, 12368-12395.	3.8	125
33	Autophagy acts as a safeguard mechanism against G-quadruplex ligand-mediated DNA damage. <i>Autophagy</i> , 2012, 8, 1185-1196.	9.1	51
34	RNA Interference-Mediated Validation of Survivin and Apollon/BRUCE as New Therapeutic Targets for Cancer Therapy. <i>Current Topics in Medicinal Chemistry</i> , 2012, 12, 69-78.	2.1	12
35	Hybrid ligand-alkylating agents targeting telomeric G-quadruplex structures. <i>Organic and Biomolecular Chemistry</i> , 2012, 10, 2798.	2.8	94
36	Telomere maintenance mechanisms in malignant peripheral nerve sheath tumors: expression and prognostic relevance. <i>Neuro-Oncology</i> , 2012, 14, 736-744.	1.2	21

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37	miR-205 regulates basement membrane deposition in human prostate: implications for cancer development. <i>Cell Death and Differentiation</i> , 2012, 19, 1750-1760.	11.2	77
38	Redox-Active Polymer Microcapsules for the Delivery of a Survivin-Specific siRNA in Prostate Cancer Cells. <i>ACS Nano</i> , 2011, 5, 1335-1344.	14.6	99
39	Telomeres as targets for anticancer therapies. <i>Expert Opinion on Therapeutic Targets</i> , 2011, 15, 579-593.	3.4	45
40	MicroRNAs as new therapeutic targets and tools in cancer. <i>Expert Opinion on Therapeutic Targets</i> , 2011, 15, 265-279.	3.4	81
41	MicroRNAs in Prostate Cancer: A Possible Role as Novel Biomarkers and Therapeutic Targets?. , 2011, , 145-162.		0
42	Targeting Survivin in Cancer Therapy: Pre-clinical Studies. , 2010, , 147-168.		1
43	Remarkable interference with telomeric function by a G-quadruplex selective bisantrene regioisomer. <i>Biochemical Pharmacology</i> , 2010, 79, 1781-1790.	4.4	17
44	miR-21: an oncomir on strike in prostate cancer. <i>Molecular Cancer</i> , 2010, 9, 12.	19.2	189
45	Apollon gene silencing induces apoptosis in breast cancer cells through p53 stabilisation and caspase-3 activation. <i>British Journal of Cancer</i> , 2009, 100, 739-746.	6.4	47
46	miR-205 Exerts Tumor-Suppressive Functions in Human Prostate through Down-regulation of Protein Kinase C μ . <i>Cancer Research</i> , 2009, 69, 2287-2295.	0.9	334
47	Targeting the telosome: Therapeutic implications. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2009, 1792, 309-316.	3.8	37
48	Towards the definition of prostate cancer-related microRNAs: where are we now?. <i>Trends in Molecular Medicine</i> , 2009, 15, 381-390.	6.7	54
49	RNA Interference-Mediated Validation of Genes Involved in Telomere Maintenance and Evasion of Apoptosis as Cancer Therapeutic Targets. <i>Methods in Molecular Biology</i> , 2009, 487, 1-28.	0.9	12
50	Targeting survivin in cancer therapy. <i>Expert Opinion on Therapeutic Targets</i> , 2008, 12, 463-476.	3.4	154
51	Validation of Telomerase and Survivin as Anticancer Therapeutic Targets Using Ribozymes and Small-Interfering RNAs. , 2007, 361, 239-264.		17
52	Photochemical Internalization: A New Tool for Drug Delivery. <i>Current Pharmaceutical Biotechnology</i> , 2007, 8, 362-372.	1.6	116
53	Targeting survivin in cancer therapy: fulfilled promises and open questions. <i>Carcinogenesis</i> , 2007, 28, 1133-1139.	2.8	217
54	Dimerizable Redox-Sensitive Triazine-Based Cationic Lipids for inâ€...vitro Gene Delivery. <i>ChemMedChem</i> , 2007, 2, 292-296.	3.2	38

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55	Photochemically enhanced delivery of a cell-penetrating peptide nucleic acid conjugate targeting human telomerase reverse transcriptase: effects on telomere status and proliferative potential of human prostate cancer cells. <i>Cell Proliferation</i> , 2007, 40, 905-920.	5.3	24
56	Down-regulation of human telomerase reverse transcriptase through specific activation of RNAi pathway quickly results in cancer cell growth impairment. <i>Biochemical Pharmacology</i> , 2007, 73, 1703-1714.	4.4	45
57	Therapeutic Uses of Peptide Nucleic Acids (PNA) in Oncology. , 2006, , 171-180.		2
58	Silencing of survivin gene by small interfering RNAs produces supra-additive growth suppression in combination with 17-allylamino-17-demethoxygeldanamycin in human prostate cancer cells. <i>Molecular Cancer Therapeutics</i> , 2006, 5, 179-186.	4.1	73
59	Therapeutic uses of peptide nucleic acids (PNA) in oncology. <i>International Journal of Peptide Research and Therapeutics</i> , 2005, 10, 287-296.	1.9	0
60	Targeting Telomerase by Antisense-Based Approaches: Perspectives for New Anti-Cancer Therapies. <i>Current Pharmaceutical Design</i> , 2005, 11, 1105-1117.	1.9	30
61	Antisense oligonucleotide-mediated inhibition of hTERT, but not hTERC, induces rapid cell growth decline and apoptosis in the absence of telomere shortening in human prostate cancer cells. <i>European Journal of Cancer</i> , 2005, 41, 624-634.	2.8	80
62	Ribozyme-mediated down-regulation of survivin expression sensitizes human melanoma cells to topotecan in vitro and in vivo. <i>Carcinogenesis</i> , 2004, 25, 1129-1136.	2.8	57
63	Ribozyme-mediated inhibition of survivin expression increases spontaneous and drug-induced apoptosis and decreases the tumorigenic potential of human prostate cancer cells. <i>Oncogene</i> , 2004, 23, 386-394.	5.9	92
64	Oligomer-mediated modulation of hTERT alternative splicing induces telomerase inhibition and cell growth decline in human prostate cancer cells. <i>Cellular and Molecular Life Sciences</i> , 2004, 61, 1764-74.	5.4	29
65	Use of ribozymes in validation of targets involved in tumor progression. <i>Drug Discovery Today: Technologies</i> , 2004, 1, 119-124.	4.0	0
66	Therapeutic uses of peptide nucleic acids (PNA) in oncology. <i>International Journal of Peptide Research and Therapeutics</i> , 2003, 10, 287-296.	0.1	0
67	Radiosensitization of Human Melanoma Cells by Ribozyme-Mediated Inhibition of Survivin Expression. <i>Journal of Investigative Dermatology</i> , 2003, 120, 648-654.	0.7	90
68	Ribozyme-mediated inhibition of PKC β sensitizes androgen-independent human prostate cancer cells to cisplatin-induced apoptosis. <i>Prostate</i> , 2003, 54, 133-143.	2.3	24
69	Therapeutic uses of peptide nucleic acids (PNA) in oncology. <i>International Journal of Peptide Research and Therapeutics</i> , 2003, 10, 287-296.	1.9	1
70	Inhibition of telomerase activity by geldanamycin and 17-allylamino, 17-demethoxygeldanamycin in human melanoma cells. <i>Carcinogenesis</i> , 2003, 24, 851-859.	2.8	43
71	Approaches for the Inhibition of Human Telomerase Based on the Use of Peptide Nucleic Acids and Hammerhead Ribozymes. <i>Mini-Reviews in Medicinal Chemistry</i> , 2003, 3, 51-60.	2.4	6
72	Photochemical internalization of a peptide nucleic acid targeting the catalytic subunit of human telomerase. <i>Cancer Research</i> , 2003, 63, 3490-4.	0.9	55

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73	Targeting Human Telomerase by Antisense Oligonucleotides and Ribozymes. <i>Anti-Cancer Agents in Medicinal Chemistry</i> , 2002, 2, 605-612.	7.0	21
74	Ribozyme-mediated attenuation of survivin expression sensitizes human melanoma cells to cisplatin-induced apoptosis. <i>Journal of Clinical Investigation</i> , 2002, 109, 285-286.	8.2	73
75	Ribozyme-mediated attenuation of survivin expression sensitizes human melanoma cells to cisplatin-induced apoptosis. <i>Journal of Clinical Investigation</i> , 2002, 109, 285-286.	8.2	51
76	Possible Regulation of Telomerase Activity by Transcription and Alternative Splicing of Telomerase Reverse Transcriptase in Human Melanoma. <i>Journal of Investigative Dermatology</i> , 2001, 116, 867-873.	0.7	37
77	Inhibition of Telomerase Activity by a Hammerhead Ribozyme Targeting the RNA Component of Telomerase in Human Melanoma Cells. <i>Journal of Investigative Dermatology</i> , 2000, 114, 259-267.	0.7	68
78	Telomerase activity and telomere length in human ovarian cancer and melanoma cell lines: correlation with sensitivity to DNA damaging agents. <i>International Journal of Oncology</i> , 2000, 16, 995-1002.	3.3	13
79	Attenuation of telomerase activity does not increase sensitivity of human melanoma cells to anticancer agents. <i>European Journal of Cancer</i> , 2000, 36, 2137-2145.	2.8	28
80	Inhibition of telomerase activity by a cell-penetrating peptide nucleic acid construct in human melanoma cells. <i>FEBS Letters</i> , 2000, 473, 241-248.	2.8	82
81	Macrophage populations of different origins have distinct susceptibilities to lipid peroxidation induced by Fe^{2+} -haematin (malaria pigment). <i>FEBS Letters</i> , 1998, 433, 215-218.	2.8	21
82	Telomerase Activity in Benign and Malignant Breast Lesions: a Pilot Prospective Study on Fine-Needle Aspirates. <i>Journal of the National Cancer Institute</i> , 1998, 90, 537-539.	6.3	25
83	Schedule-dependent modulation of idarubicin cytotoxicity by lonidamine in human lymphoma cell lines. <i>International Journal of Oncology</i> , 1997, 11, 675-9.	3.3	0