Yves Pommier

List of Publications by Year in descending order

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257 papers 29,327 citations

80 h-index 161 g-index

283 all docs 283
docs citations

times ranked

283

28281 citing authors

#	Article	IF	CITATIONS
1	Topoisomerase I inhibitors: camptothecins and beyond. Nature Reviews Cancer, 2006, 6, 789-802.	12.8	1,824
2	Trapping of PARP1 and PARP2 by Clinical PARP Inhibitors. Cancer Research, 2012, 72, 5588-5599.	0.4	1,657
3	DNA Topoisomerases and Their Poisoning by Anticancer and Antibacterial Drugs. Chemistry and Biology, 2010, 17, 421-433.	6.2	1,507
4	Î ³ H2AX and cancer. Nature Reviews Cancer, 2008, 8, 957-967.	12.8	1,423
5	A gene expression database for the molecular pharmacology of cancer. Nature Genetics, 2000, 24, 236-244.	9.4	1,357
6	Drugging Topoisomerases: Lessons and Challenges. ACS Chemical Biology, 2013, 8, 82-95.	1.6	698
7	Roles of eukaryotic topoisomerases in transcription, replication and genomic stability. Nature Reviews Molecular Cell Biology, 2016, 17, 703-721.	16.1	695
8	Stereospecific PARP Trapping by BMN 673 and Comparison with Olaparib and Rucaparib. Molecular Cancer Therapeutics, 2014, 13, 433-443.	1.9	627
9	Integrase inhibitors to treat HIV/Aids. Nature Reviews Drug Discovery, 2005, 4, 236-248.	21.5	612
10	DNA Topoisomerase I Inhibitors: Chemistry, Biology, and Interfacial Inhibition. Chemical Reviews, 2009, 109, 2894-2902.	23.0	609
11	Apoptosis defects and chemotherapy resistance: molecular interaction maps and networks. Oncogene, 2004, 23, 2934-2949.	2.6	524
12	Laying a trap to kill cancer cells: PARP inhibitors and their mechanisms of action. Science Translational Medicine, 2016, 8, 362ps17.	5.8	518
13	Structures of Three Classes of Anticancer Agents Bound to the Human Topoisomerase Iâ^DNA Covalent Complex. Journal of Medicinal Chemistry, 2005, 48, 2336-2345.	2.9	447
14	Topoisomerase I suppresses genomic instability by preventing interference between replication and transcription. Nature Cell Biology, 2009, 11, 1315-1324.	4.6	445
15	A subset of platinum-containing chemotherapeutic agents kills cells by inducing ribosome biogenesis stress. Nature Medicine, 2017, 23, 461-471.	15.2	379
16	Antiproliferative activity of ecteinascidin 743 is dependent upon transcription-coupled nucleotide-excision repair. Nature Medicine, 2001, 7, 961-966.	15.2	339
17	Conversion of Topoisomerase I Cleavage Complexes on the Leading Strand of Ribosomal DNA into 5′-Phosphorylated DNA Double-Strand Breaks by Replication Runoff. Molecular and Cellular Biology, 2000, 20, 3977-3987.	1.1	314
18	Genome Organization Drives Chromosome Fragility. Cell, 2017, 170, 507-521.e18.	13.5	311

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19	DNA Sequence- and Structure-Selective Alkylation of Guanine N2 in the DNA Minor Groove by Ecteinascidin 743, a Potent Antitumor Compound from the Caribbean TunicateEcteinascidia turbinata. Biochemistry, 1996, 35, 13303-13309.	1.2	288
20	CellMiner: a relational database and query tool for the NCI-60 cancer cell lines. BMC Genomics, 2009, 10, 277.	1.2	273
21	Tyrosyl-DNA-phosphodiesterases (TDP1 and TDP2). DNA Repair, 2014, 19, 114-129.	1.3	253
22	Putative DNA/RNA helicase Schlafen-11 (SLFN11) sensitizes cancer cells to DNA-damaging agents. Proceedings of the National Academy of Sciences of the United States of America, 2012, 109, 15030-15035.	3.3	252
23	Mutagenic Processing of Ribonucleotides in DNA by Yeast Topoisomerase I. Science, 2011, 332, 1561-1564.	6.0	251
24	Repair of Topoisomerase Iâ€Mediated DNA Damage. Progress in Molecular Biology and Translational Science, 2006, 81, 179-229.	1.9	247
25	The Exomes of the NCI-60 Panel: A Genomic Resource for Cancer Biology and Systems Pharmacology. Cancer Research, 2013, 73, 4372-4382.	0.4	239
26	Rationale for Poly(ADP-ribose) Polymerase (PARP) Inhibitors in Combination Therapy with Camptothecins or Temozolomide Based on PARP Trapping versus Catalytic Inhibition. Journal of Pharmacology and Experimental Therapeutics, 2014, 349, 408-416.	1.3	237
27	Curcumin Analogs with Altered Potencies against HIV-1 Integrase as Probes for Biochemical Mechanisms of Drug Action. Journal of Medicinal Chemistry, 1997, 40, 3057-3063.	2.9	228
28	Resistance to PARP inhibitors by SLFN11 inactivation can be overcome by ATR inhibition. Oncotarget, 2016, 7, 76534-76550.	0.8	219
29	RNA Polymerase II Regulates Topoisomerase 1 Activity to Favor Efficient Transcription. Cell, 2016, 165, 357-371.	13.5	211
30	Ataxia telangiectasia mutated activation by transcription―and topoisomerase l―induced DNA doubleâ€strand breaks. EMBO Reports, 2009, 10, 887-893.	2.0	208
31	Cellular Determinants of Sensitivity and Resistance to DNA Topoisomerase Inhibitors. Cancer Investigation, 1994, 12, 530-542.	0.6	204
32	Interfacial inhibitors: targeting macromolecular complexes. Nature Reviews Drug Discovery, 2012, 11, 25-36.	21.5	204
33	Targeting Topoisomerase I in the Era of Precision Medicine. Clinical Cancer Research, 2019, 25, 6581-6589.	3.2	184
34	Topoisomerase I-mediated DNA damage. Advances in Cancer Research, 2001, 80, 189-216.	1.9	182
35	Local sequence requirements for DNA cleavage by mammalian topoisomerase II in the presence of doxorubicin. Nucleic Acids Research, 1990, 18, 6611-6619.	6.5	179
36	SLFN11 Blocks Stressed Replication Forks Independently of ATR. Molecular Cell, 2018, 69, 371-384.e6.	4.5	177

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37	Quantitative Structure-Antitumor Activity Relationships of Camptothecin Analogues:  Cluster Analysis and Genetic Algorithm-Based Studies. Journal of Medicinal Chemistry, 2001, 44, 3254-3263.	2.9	171
38	Induction of Reversible Complexes between Eukaryotic DNA Topoisomerase I and DNA-containing Oxidative Base Damages. Journal of Biological Chemistry, 1999, 274, 8516-8523.	1.6	168
39	Effects of Uracil Incorporation, DNA Mismatches, and Abasic Sites on Cleavage and Religation Activities of Mammalian Topoisomerase I. Journal of Biological Chemistry, 1997, 272, 7792-7796.	1.6	164
40	PARP1–TDP1 coupling for the repair of topoisomerase l–induced DNA damage. Nucleic Acids Research, 2014, 42, 4435-4449.	6.5	163
41	Interfacial inhibition of macromolecular interactions: nature's paradigm for drug discovery. Trends in Pharmacological Sciences, 2005, 26, 138-145.	4.0	161
42	Topoisomerase I inhibitors: selectivity and cellular resistance. Drug Resistance Updates, 1999, 2, 307-318.	6.5	158
43	Tyrosyl-DNA Phosphodiesterase 1 (TDP1) Repairs DNA Damage Induced by Topoisomerases I and II and Base Alkylation in Vertebrate Cells. Journal of Biological Chemistry, 2012, 287, 12848-12857.	1.6	155
44	Trapping of Mammalian Topoisomerase I and Recombinations Induced by Damaged DNA Containing Nicks or Gaps. Journal of Biological Chemistry, 1997, 272, 26441-26447.	1.6	153
45	The Novel Silatecan 7-tert-Butyldimethylsilyl-10-hydroxycamptothecin Displays High Lipophilicity, Improved Human Blood Stability, and Potent Anticancer Activity. Journal of Medicinal Chemistry, 2000, 43, 3970-3980.	2.9	147
46	Topoisomerase II-Induced Chromosome Breakage and Translocation Is Determined by Chromosome Architecture and Transcriptional Activity. Molecular Cell, 2019, 75, 252-266.e8.	4.5	145
47	The indenoisoquinoline noncamptothecin topoisomerase I inhibitors: update and perspectives. Molecular Cancer Therapeutics, 2009, 8, 1008-1014.	1.9	144
48	Ion Selective Folding of Loop Domains in a Potent Anti-HIV Oligonucleotideâ€. Biochemistry, 1997, 36, 12498-12505.	1.2	139
49	Mus81-mediated DNA cleavage resolves replication forks stalled by topoisomerase l–DNA complexes. Journal of Cell Biology, 2011, 195, 739-749.	2.3	138
50	Local base sequence preferences for DNA cleavage by mammalian topoisomerase II in the presence of amsacrine or teniposide. Nucleic Acids Research, 1991, 19, 5973-5980.	6.5	136
51	Production of Extrachromosomal MicroDNAs Is Linked to Mismatch Repair Pathways and Transcriptional Activity. Cell Reports, 2015, 11, 1749-1759.	2.9	135
52	Tyrosyl-DNA Phosphodiesterase as a Target for Anticancer Therapy. Anti-Cancer Agents in Medicinal Chemistry, 2008, 8, 381-389.	0.9	133
53	Poly(ADP-ribose) polymerase and XPF–ERCC1 participate in distinct pathways for the repair of topoisomerase I-induced DNA damage in mammalian cells. Nucleic Acids Research, 2011, 39, 3607-3620.	6.5	132
54	ATR Inhibitors VE-821 and VX-970 Sensitize Cancer Cells to Topoisomerase I Inhibitors by Disabling DNA Replication Initiation and Fork Elongation Responses. Cancer Research, 2014, 74, 6968-6979.	0.4	131

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55	MGMT Status as a Clinical Biomarker in Glioblastoma. Trends in Cancer, 2020, 6, 380-391.	3.8	131
56	Mechanisms of Camptothecin Resistance by Human Topoisomerase I Mutations. Journal of Molecular Biology, 2004, 339, 773-784.	2.0	129
57	Optimal function of the DNA repair enzyme TDP1 requires its phosphorylation by ATM and/or DNA-PK. EMBO Journal, 2009, 28, 3667-3680.	3.5	125
58	Human topoisomerases and their roles in genome stability and organization. Nature Reviews Molecular Cell Biology, 2022, 23, 407-427.	16.1	125
59	Role of tyrosyl-DNA phosphodiesterase (TDP1) in mitochondria. Proceedings of the National Academy of Sciences of the United States of America, 2010, 107, 19790-19795.	3.3	124
60	Synthesis of Cytotoxic Indenoisoquinoline Topoisomerase I Poisons. Journal of Medicinal Chemistry, 1999, 42, 446-457.	2.9	122
61	Epigenetic inactivation of the putative DNA/RNA helicase SLFN11 in human cancer confers resistance to platinum drugs. Oncotarget, 2016, 7, 3084-3097.	0.8	120
62	Processing of nucleopeptides mimicking the topoisomerase I-DNA covalent complex by tyrosyl-DNA phosphodiesterase. Nucleic Acids Research, 2002, 30, 1198-1204.	6.5	119
63	CellMinerCDB for Integrative Cross-Database Genomics and Pharmacogenomics Analyses of Cancer Cell Lines. IScience, 2018, 10, 247-264.	1.9	117
64	Implication of Checkpoint Kinase-dependent Up-regulation of Ribonucleotide Reductase R2 in DNA Damage Response. Journal of Biological Chemistry, 2009, 284, 18085-18095.	1.6	116
65	Differential and Common DNA Repair Pathways for Topoisomerase I- and II-Targeted Drugs in a Genetic DT40 Repair Cell Screen Panel. Molecular Cancer Therapeutics, 2014, 13, 214-220.	1.9	116
66	Apoptotic response to camptothecin and 7-hydroxystaurosporine (UCN-01) in the 8 human breast cancer cell lines of the NCI anticancer drug screen: Multifactorial relationships with topoisomerase i, protein kinase C, Bcl-2, p53, MDM-2 and caspase pathways., 1999, 82, 396-404.		111
67	Dual Processing of R-Loops and Topoisomerase I Induces Transcription-Dependent DNA Double-Strand Breaks. Cell Reports, 2019, 28, 3167-3181.e6.	2.9	108
68	Therapeutic targeting of ATR yields durable regressions in small cell lung cancers with high replication stress. Cancer Cell, 2021, 39, 566-579.e7.	7.7	107
69	Schlafen 11 (SLFN11), a restriction factor for replicative stress induced by DNA-targeting anti-cancer therapies., 2019, 201, 94-102.		106
70	Proteolytic Degradation of Topoisomerase II (Top2) Enables the Processing of Top2·DNA and Top2·RNA Covalent Complexes by Tyrosyl-DNA-Phosphodiesterase 2 (TDP2). Journal of Biological Chemistry, 2014, 289, 17960-17969.	1.6	103
71	Synthesis and Mechanism of Action Studies of a Series of Norindenoisoquinoline Topoisomerase I Poisons Reveal an Inhibitor with a Flipped Orientation in the Ternary DNAâ^'Enzymeâ^'Inhibitor Complex As Determined by X-ray Crystallographic Analysis. Journal of Medicinal Chemistry, 2005, 48, 4803-4814.	2.9	102
72	Mitochondrial Topoisomerase I (Top 1 mt) Is a Novel Limiting Factor of Doxorubicin Cardiotoxicity. Clinical Cancer Research, 2014, 20, 4873-4881.	3.2	102

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73	Nonclassic Functions of Human Topoisomerase I: Genome-Wide and Pharmacologic Analyses. Cancer Research, 2007, 67, 8752-8761.	0.4	93
74	Mitochondrial Topoisomerase I is Critical for Mitochondrial Integrity and Cellular Energy Metabolism. PLoS ONE, 2012, 7, e41094.	1.1	93
75	DNA cleavage assay for the identification of topoisomerase I inhibitors. Nature Protocols, 2008, 3, 1736-1750.	5.5	92
76	Temozolomide in the Era of Precision Medicine. Cancer Research, 2017, 77, 823-826.	0.4	91
77	<i>SLFN11</i> Is a Transcriptional Target of EWS-FLI1 and a Determinant of Drug Response in Ewing Sarcoma. Clinical Cancer Research, 2015, 21, 4184-4193.	3.2	89
78	Camptothecins and Topoisomerase I; A Foot in the Door. Targeting the Genome Beyond Topoisomerase I with Camptothecins and Novel Anticancer Drugs; Importance of DNA Replication, Repair and Cell Cycle Checkpoints. Anti-Cancer Agents in Medicinal Chemistry, 2004, 4, 429-434.	7.0	87
79	TDP2 promotes repair of topoisomerase I-mediated DNA damage in the absence of TDP1. Nucleic Acids Research, 2012, 40, 8371-8380.	6.5	86
80	TDP1 repairs nuclear and mitochondrial DNA damage induced by chain-terminating anticancer and antiviral nucleoside analogs. Nucleic Acids Research, 2013, 41, 7793-7803.	6.5	86
81	SCLC-CellMiner: A Resource for Small Cell Lung Cancer Cell Line Genomics and Pharmacology Based on Genomic Signatures. Cell Reports, 2020, 33, 108296.	2.9	86
82	BRCAness, SLFN11, and RB1 loss predict response to topoisomerase I inhibitors in triple-negative breast cancers. Science Translational Medicine, 2020, 12, .	5.8	86
83	Synthesis and Biological Evaluation of the First Dual Tyrosyl-DNA Phosphodiesterase I (Top1) Inhibitors. Journal of Medicinal Chemistry, 2012, 55, 4457-4478.	2.9	85
84	Gemcitabine (2',2'-difluoro-2'-deoxycytidine), an antimetabolite that poisons topoisomerase I. Clinical Cancer Research, 2002, 8, 2499-504.	3.2	82
85	Genomic and evolutionary classification of lung cancer in never smokers. Nature Genetics, 2021, 53, 1348-1359.	9.4	81
86	Activation of the Fas pathway independently of Fas ligand during apoptosis induced by camptothecin in p53 mutant human colon carcinoma cells. Oncogene, 2001, 20, 1852-1859.	2.6	80
87	Using CellMiner 1.6 for Systems Pharmacology and Genomic Analysis of the NCI-60. Clinical Cancer Research, 2015, 21, 3841-3852.	3.2	80
88	Apoptosis Induced by DNA Topoisomerase I and II Inhibitors in Human Leukemic HL-60 Cells. Leukemia and Lymphoma, 1994, 15, 21-32.	0.6	78
89	Chk2 Molecular Interaction Map and Rationale for Chk2 Inhibitors: Fig. 1 Clinical Cancer Research, 2006, 12, 2657-2661.	3.2	78
90	Tyrosyl-DNA Phosphodiesterase 1 (Tdp1) inhibitors. Expert Opinion on Therapeutic Patents, 2011, 21, 1285-1292.	2.4	78

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91	A conserved SUMO pathway repairs topoisomerase DNA-protein cross-links by engaging ubiquitin-mediated proteasomal degradation. Science Advances, 2020, 6, .	4.7	76
92	Gene Expression Correlations in Human Cancer Cell Lines Define Molecular Interaction Networks for Epithelial Phenotype. PLoS ONE, 2014, 9, e99269.	1.1	76
93	Phosphorylated fraction of H2AX as a measurement for DNA damage in cancer cells and potential applications of a novel assay. PLoS ONE, 2017, 12, e0171582.	1.1	72
94	8-Oxoguanine rearranges the active site of human topoisomerase I. Proceedings of the National Academy of Sciences of the United States of America, 2002, 99, 12102-12107.	3.3	68
95	Increased negative supercoiling of mtDNA in TOP1mt knockout mice and presence of topoisomerases IIÂ and IIÂ in vertebrate mitochondria. Nucleic Acids Research, 2014, 42, 7259-7267.	6.5	67
96	The ubiquitin-dependent ATPase p97 removes cytotoxic trapped PARP1 from chromatin. Nature Cell Biology, 2022, 24, 62-73.	4.6	66
97	Interfacial Inhibitors of Protein-Nucleic Acid Interactions. Anti-Cancer Agents in Medicinal Chemistry, 2005, 5, 421-429.	7.0	65
98	Overcoming Resistance to DNA-Targeted Agents by Epigenetic Activation of Schlafen 11 (<i>SLFN11)</i> Expression with Class I Histone Deacetylase Inhibitors. Clinical Cancer Research, 2018, 24, 1944-1953.	3.2	65
99	Molecular and Biological Determinants of the Cytotoxic Actions of Camptothecins: Perspective for the Development of New Topoisomerase I Inhibitors. Annals of the New York Academy of Sciences, 2000, 922, 11-26.	1.8	64
100	RNA topoisomerase is prevalent in all domains of life and associates with polyribosomes in animals. Nucleic Acids Research, 2016, 44, 6335-6349.	6.5	63
101	Antisense inhibition of Chk2/hCds1 expression attenuates DNA damage-induced S and G2 checkpoints and enhances apoptotic activity in HEK-293 cells. FEBS Letters, 2001, 505, 7-12.	1.3	62
102	Excision repair of topoisomerase DNA-protein crosslinks (TOP-DPC). DNA Repair, 2020, 89, 102837.	1.3	62
103	Topoisomerase lâ€mediated cleavage at unrepaired ribonucleotides generates DNA doubleâ€strand breaks. EMBO Journal, 2017, 36, 361-373.	3.5	59
104	Novel and Highly Potent ATR Inhibitor M4344 Kills Cancer Cells With Replication Stress, and Enhances the Chemotherapeutic Activity of Widely Used DNA Damaging Agents. Molecular Cancer Therapeutics, 2021, 20, 1431-1441.	1.9	58
105	Differential cytotoxicity of clinically important camptothecin derivatives in P-glycoprotein-overexpressing cell lines. Cancer Chemotherapy and Pharmacology, 1997, 40, 433-438.	1.1	57
106	Nucleosome positioning as a critical determinant for the DNA cleavage sites of mammalian DNA topoisomerase in reconstituted Simian virus 40 chromatin. Nucleic Acids Research, 1990, 18, 4553-4559.	6. 5	56
107	Poisoning of Mitochondrial Topoisomerase I by Lamellarin D. Molecular Pharmacology, 2014, 86, 193-199.	1.0	56
108	Hyperphosphorylation of RNA Polymerase II in Response to Topoisomerase I Cleavage Complexes and Its Association with Transcription- and BRCA1-dependent Degradation of Topoisomerase I. Journal of Molecular Biology, 2008, 381, 540-549.	2.0	55

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109	Alterations of DNA repair genes in the NCI-60 cell lines and their predictive value for anticancer drug activity. DNA Repair, 2015, 28, 107-115.	1.3	55
110	Multiview confocal super-resolution microscopy. Nature, 2021, 600, 279-284.	13.7	55
111	Biochemical Characterization of Human Tyrosyl-DNA Phosphodiesterase 2 (TDP2/TTRAP). Journal of Biological Chemistry, 2012, 287, 30842-30852.	1.6	54
112	Structural basis for recognition of 5′-phosphotyrosine adducts by Tdp2. Nature Structural and Molecular Biology, 2012, 19, 1372-1377.	3.6	53
113	Topoisomerase I Alone Is Sufficient to Produce Short DNA Deletions and Can Also Reverse Nicks at Ribonucleotide Sites. Journal of Biological Chemistry, 2015, 290, 14068-14076.	1.6	52
114	Isoquinoline-1,3-diones as Selective Inhibitors of Tyrosyl DNA Phosphodiesterase II (TDP2). Journal of Medicinal Chemistry, 2016, 59, 2734-2746.	2.9	52
115	Sensitivity of Mesothelioma Cells to PARP Inhibitors Is Not Dependent on BAP1 but Is Enhanced by Temozolomide in Cells With High-Schlafen 11 and Low-O6-methylguanine-DNA Methyltransferase Expression. Journal of Thoracic Oncology, 2020, 15, 843-859.	0.5	51
116	Synthesis and Biological Evaluation of Nitrated 7-, 8-, 9-, and 10-Hydroxyindenoisoquinolines as Potential Dual Topoisomerase I (Top1)–Tyrosyl-DNA Phosphodiesterase I (TDP1) Inhibitors. Journal of Medicinal Chemistry, 2015, 58, 3188-3208.	2.9	50
117	Lack of mitochondrial topoisomerase I (<i>TOP1mt</i>) impairs liver regeneration. Proceedings of the National Academy of Sciences of the United States of America, 2015, 112, 11282-11287.	3.3	50
118	The NCI-60 Methylome and Its Integration into CellMiner. Cancer Research, 2017, 77, 601-612.	0.4	48
119	Debulking of topoisomerase DNA-protein crosslinks (TOP-DPC) by the proteasome, non-proteasomal and non-proteolytic pathways. DNA Repair, 2020, 94, 102926.	1.3	48
120	Chromatin Remodeling and Immediate Early Gene Activation by SLFN11 in Response to Replication Stress. Cell Reports, 2020, 30, 4137-4151.e6.	2.9	48
121	DNA recombinase activity of eukaryotic DNA topoisomerase I; effects of camptothecin and other inhibitors. Mutation Research DNA Repair, 1995, 337, 135-145.	3.8	47
122	DNA double-strand breaks and ATM activation by transcription-blocking DNA lesions. Cell Cycle, 2010, 9, 274-278.	1.3	47
123	A kinetic clutch governs religation by type IB topoisomerases and determines camptothecin sensitivity. Proceedings of the National Academy of Sciences of the United States of America, 2012, 109, 16125-16130.	3.3	47
124	The evolving landscape of predictive biomarkers of response to PARP inhibitors. Journal of Clinical Investigation, 2018, 128, 1727-1730.	3.9	47
125	Relative contribution of four nucleases, CtIP, Dna2, Exo1 and Mre11, to the initial step of DNA doubleâ€strand break repair by homologous recombination in both the chicken DT40 and human TK6 cell lines. Genes To Cells, 2015, 20, 1059-1076.	0.5	46
126	Structure-Guided Optimization of HIV Integrase Strand Transfer Inhibitors. Journal of Medicinal Chemistry, 2017, 60, 7315-7332.	2.9	44

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127	Synthesis and Biological Evaluation of the First Triple Inhibitors of Human Topoisomerase 1, Tyrosyl–DNA Phosphodiesterase 1 (Tdp1), and Tyrosyl–DNA Phosphodiesterase 2 (Tdp2). Journal of Medicinal Chemistry, 2017, 60, 3275-3288.	2.9	43
128	Design, Synthesis, and Biological Evaluation of Potential Prodrugs Related to the Experimental Anticancer Agent Indotecan (LMP400). Journal of Medicinal Chemistry, 2016, 59, 4890-4899.	2.9	42
129	Structural Impact of the Leukemia Drug 1-ॆ-d-Arabinofuranosylcytosine (Ara-C) on the Covalent Human Topoisomerase I-DNA Complex. Journal of Biological Chemistry, 2003, 278, 12461-12466.	1.6	41
130	Discovery of Potent Indenoisoquinoline Topoisomerase I Poisons Lacking the 3-Nitro Toxicophore. Journal of Medicinal Chemistry, 2015, 58, 3997-4015.	2.9	40
131	Small cell lung cancer: Time to revisit DNA-damaging chemotherapy. Science Translational Medicine, 2016, 8, 346fs12.	5.8	40
132	PRMT5-mediated arginine methylation of TDP1 for the repair of topoisomerase I covalent complexes. Nucleic Acids Research, 2018, 46, 5601-5617.	6.5	40
133	BAMscale: quantification of next-generation sequencing peaks and generation of scaled coverage tracks. Epigenetics and Chromatin, 2020, 13, 21.	1.8	40
134	rcellminer: exploring molecular profiles and drug response of the NCI-60 cell lines in R. Bioinformatics, 2016, 32, 1272-1274.	1.8	39
135	<i>N</i> -Substituted Quinolinonyl Diketo Acid Derivatives as HIV Integrase Strand Transfer Inhibitors and Their Activity against RNase H Function of Reverse Transcriptase. Journal of Medicinal Chemistry, 2015, 58, 4610-4623.	2.9	38
136	Camptothecin targets WRN protein: mechanism and relevance in clinical breast cancer. Oncotarget, 2016, 7, 13269-13284.	0.8	38
137	Negative regulation of mitochondrial transcription by mitochondrial topoisomerase I. Nucleic Acids Research, 2013, 41, 9848-9857.	6.5	37
138	Discovery, Synthesis, and Evaluation of Oxynitidine Derivatives as Dual Inhibitors of DNA Topoisomerase IB (TOP1) and Tyrosyl-DNA Phosphodiesterase 1 (TDP1), and Potential Antitumor Agents. Journal of Medicinal Chemistry, 2018, 61, 9908-9930.	2.9	37
139	Different Effects on Human Topoisomerase I by Minor Groove and Intercalated Deoxyguanosine Adducts Derived from Two Polycyclic Aromatic Hydrocarbon Diol Epoxides at or Near a Normal Cleavage Site. Journal of Biological Chemistry, 2002, 277, 13666-13672.	1.6	36
140	NCI Comparative Oncology Program Testing of Non-Camptothecin Indenoisoquinoline Topoisomerase I Inhibitors in Naturally Occurring Canine Lymphoma. Clinical Cancer Research, 2018, 24, 5830-5840.	3.2	36
141	High Resolution Copy Number Variation Data in the NCI-60 Cancer Cell Lines from Whole Genome Microarrays Accessible through CellMiner. PLoS ONE, 2014, 9, e92047.	1.1	36
142	Substitutions of Asn-726 in the Active Site of Yeast DNA Topoisomerase I Define Novel Mechanisms of Stabilizing the Covalent Enzyme-DNA Intermediate. Journal of Biological Chemistry, 2000, 275, 15246-15253.	1.6	35
143	Investigation of the Structure–Activity Relationships of Aza-A-Ring Indenoisoquinoline Topoisomerase I Poisons. Journal of Medicinal Chemistry, 2016, 59, 3840-3853.	2.9	35
144	HIV-1 Integrase Strand Transfer Inhibitors with Reduced Susceptibility to Drug Resistant Mutant Integrases. ACS Chemical Biology, 2016, 11, 1074-1081.	1.6	35

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145	Acquired SETD2 mutation and impaired CREB1 activation confer cisplatin resistance in metastatic non-small cell lung cancer. Oncogene, 2019, 38, 180-193.	2.6	35
146	Whole-exome sequencing reveals germline-mutated small cell lung cancer subtype with favorable response to DNA repair–targeted therapies. Science Translational Medicine, 2021, 13, .	5.8	35
147	Human DNA topoisomerase I-mediated cleavage and recombination of duck hepatitis B virus DNA in vitro. Nucleic Acids Research, 1999, 27, 1919-1925.	6.5	34
148	Thirteen-exon-motif signature for vertebrate nuclear and mitochondrial type IB topoisomerases. Nucleic Acids Research, 2004, 32, 2087-2092.	6.5	34
149	Interfacial inhibitors. Bioorganic and Medicinal Chemistry Letters, 2015, 25, 3961-3965.	1.0	34
150	The Indenoisoquinoline TOP1 Inhibitors Selectively Target Homologous Recombination-Deficient and Schlafen 11-Positive Cancer Cells and Synergize with Olaparib. Clinical Cancer Research, 2019, 25, 6206-6216.	3.2	34
151	ALC1/CHD1L, a chromatin-remodeling enzyme, is required for efficient base excision repair. PLoS ONE, 2017, 12, e0188320.	1.1	34
152	Clinical and pharmacologic evaluation of two dosing schedules of indotecan (LMP400), a novel indenoisoquinoline, in patients with advanced solid tumors. Cancer Chemotherapy and Pharmacology, 2016, 78, 73-81.	1.1	32
153	Deazaflavin Inhibitors of Tyrosyl-DNA Phosphodiesterase 2 (TDP2) Specific for the Human Enzyme and Active against Cellular TDP2. ACS Chemical Biology, 2016, 11, 1925-1933.	1.6	32
154	Novel Fluoroindenoisoquinoline Non-Camptothecin Topoisomerase I Inhibitors. Molecular Cancer Therapeutics, 2018, 17, 1694-1704.	1.9	30
155	SLFN11 promotes CDT1 degradation by CUL4 in response to replicative DNA damage, while its absence leads to synthetic lethality with ATR/CHK1 inhibitors. Proceedings of the National Academy of Sciences of the United States of America, 2021, 118 , .	3.3	30
156	Analogs of the novel phytohormone, strigolactone, trigger apoptosis and synergize with PARP inhibitors by inducing DNA damage and inhibiting DNA repair. Oncotarget, 2016, 7, 13984-14001.	0.8	30
157	Design and Synthesis of Chlorinated and Fluorinated 7-Azaindenoisoquinolines as Potent Cytotoxic Anticancer Agents That Inhibit Topoisomerase I. Journal of Medicinal Chemistry, 2017, 60, 5364-5376.	2.9	29
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