

Yves Pommier

List of Publications by Year in descending order

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Version: 2024-02-01

257
papers

29,327
citations

6250

80
h-index

5677

162
g-index

283
all docs

283
docs citations

283
times ranked

28281
citing authors

#	ARTICLE	IF	CITATIONS
1	Prognostic impact of Schlafen 11 in bladder cancer patients treated with platinum-based chemotherapy. <i>Cancer Science</i> , 2022, 113, 784-795.	1.7	10
2	The ubiquitin-dependent ATPase p97 removes cytotoxic trapped PARP1 from chromatin. <i>Nature Cell Biology</i> , 2022, 24, 62-73.	4.6	66
3	Topoisomerase I (TOP1) dynamics: conformational transition from open to closed states. <i>Nature Communications</i> , 2022, 13, 59.	5.8	11
4	Human topoisomerases and their roles in genome stability and organization. <i>Nature Reviews Molecular Cell Biology</i> , 2022, 23, 407-427.	16.1	125
5	Synthesis of 11-aminoalkoxy substituted benzophenanthridine derivatives as tyrosyl-DNA phosphodiesterase 1 inhibitors and their anticancer activity. <i>Bioorganic Chemistry</i> , 2022, 123, 105789.	2.0	4
6	SUMO: A Swiss Army Knife for Eukaryotic Topoisomerases. <i>Frontiers in Molecular Biosciences</i> , 2022, 9, 871161.	1.6	7
7	Cancer/Testis Antigen 55 is required for cancer cell proliferation and mitochondrial DNA maintenance. <i>Mitochondrion</i> , 2022, 64, 19-26.	1.6	2
8	2-Arylquinolines as novel anticancer agents with dual EGFR/FAK kinase inhibitory activity: synthesis, biological evaluation, and molecular modelling insights. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2022, 37, 355-378.	2.5	15
9	TOP1-DNA Trapping by Exatecan and Combination Therapy with ATR Inhibitor. <i>Molecular Cancer Therapeutics</i> , 2022, 21, 1090-1102.	1.9	13
10	From Antarctica to cancer research: a novel human DNA topoisomerase 1B inhibitor from Antarctic sponge <i>Dendrilla antarctica</i> . <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2022, 37, 1404-1410.	2.5	5
11	Replication Stress Defines Distinct Molecular Subtypes Across Cancers. <i>Cancer Research Communications</i> , 2022, 2, 503-517.	0.7	12
12	Structural, molecular, and functional insights into Schlafen proteins. <i>Experimental and Molecular Medicine</i> , 2022, 54, 730-738.	3.2	17
13	CDK7 Inhibition Synergizes with Topoisomerase I Inhibition in Small Cell Lung Cancer Cells by Inducing Ubiquitin-Mediated Proteolysis of RNA Polymerase II. <i>Molecular Cancer Therapeutics</i> , 2022, 21, 1430-1438.	1.9	3
14	Resolution of R-loops by topoisomerase III- β (TOP3B) in coordination with the DEAD-box helicase DDX5. <i>Cell Reports</i> , 2022, 40, 111067.	2.9	19
15	Immunohistochemical analysis of SLFN11 expression uncovers potential non-responders to DNA-damaging agents overlooked by tissue RNA-seq. <i>Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin</i> , 2021, 478, 569-579.	1.4	25
16	An interplay of NOX1-derived ROS and oxygen determines the spermatogonial stem cell self-renewal efficiency under hypoxia. <i>Genes and Development</i> , 2021, 35, 250-260.	2.7	19
17	Small molecule microarray identifies inhibitors of tyrosyl-DNA phosphodiesterase 1 that simultaneously access the catalytic pocket and two substrate binding sites. <i>Chemical Science</i> , 2021, 12, 3876-3884.	3.7	18
18	Whole-exome sequencing reveals germline-mutated small cell lung cancer subtype with favorable response to DNA repair-targeted therapies. <i>Science Translational Medicine</i> , 2021, 13, .	5.8	35

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19	SLFN11 promotes CDT1 degradation by CUL4 in response to replicative DNA damage, while its absence leads to synthetic lethality with ATR/CHK1 inhibitors. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2021, 118, .	3.3	30
20	Exonuclease VII repairs quinolone-induced damage by resolving DNA gyrase cleavage complexes. <i>Science Advances</i> , 2021, 7, .	4.7	6
21	SLFN11 Inactivation Induces Proteotoxic Stress and Sensitizes Cancer Cells to Ubiquitin Activating Enzyme Inhibitor TAK-243. <i>Cancer Research</i> , 2021, 81, 3067-3078.	0.4	23
22	Discovery of 4-alkoxy-2-aryl-6,7-dimethoxyquinolines as a new class of topoisomerase I inhibitors endowed with potent in vitro anticancer activity. <i>European Journal of Medicinal Chemistry</i> , 2021, 215, 113261.	2.6	24
23	Autophagy-Dependent Sensitization of Triple-Negative Breast Cancer Models to Topoisomerase II Poisons by Inhibition of the Nucleosome Remodeling Factor. <i>Molecular Cancer Research</i> , 2021, 19, 1338-1349.	1.5	9
24	Therapeutic targeting of ATR yields durable regressions in small cell lung cancers with high replication stress. <i>Cancer Cell</i> , 2021, 39, 566-579.e7.	7.7	107
25	Synthesis of Methoxy-, Methyleneoxy-, Hydroxy-, and Halo-Substituted Benzophenanthridinone Derivatives as DNA Topoisomerase IB (TOP1) and Tyrosyl-DNA Phosphodiesterase 1 (TDP1) Inhibitors and Their Biological Activity for Drug-Resistant Cancer. <i>Journal of Medicinal Chemistry</i> , 2021, 64, 7617-7629.	2.9	14
26	Novel and Highly Potent ATR Inhibitor M4344 Kills Cancer Cells With Replication Stress, and Enhances the Chemotherapeutic Activity of Widely Used DNA Damaging Agents. <i>Molecular Cancer Therapeutics</i> , 2021, 20, 1431-1441.	1.9	58
27	Design and synthesis of C-aryl angular luotonins via a one-pot aza-Nazarov-Friedlander sequence and their Topo-I inhibition studies along with C-aryl vasicinones and luotonins. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2021, 41, 127998.	1.0	5
28	PARylation prevents the proteasomal degradation of topoisomerase I DNA-protein crosslinks and induces their deubiquitylation. <i>Nature Communications</i> , 2021, 12, 5010.	5.8	26
29	Precision Oncology with Drugs Targeting the Replication Stress, ATR, and Schlafen 11. <i>Cancers</i> , 2021, 13, 4601.	1.7	19
30	Genomic and evolutionary classification of lung cancer in never smokers. <i>Nature Genetics</i> , 2021, 53, 1348-1359.	9.4	81
31	Schlafen 11 Expression in Human Acute Leukemia Cells with Gain-of-Function Mutations in the Interferon-JAK Signaling Pathway. <i>iScience</i> , 2021, 24, 103173.	1.9	6
32	Replication-dependent cytotoxicity and Spartan-mediated repair of trapped PARP1-DNA complexes. <i>Nucleic Acids Research</i> , 2021, 49, 10493-10506.	6.5	16
33	Epigenetic suppression of SLFN11 in germinal center B-cells during B-cell development. <i>PLoS ONE</i> , 2021, 16, e0237554.	1.1	20
34	A polymer index-matched to water enables diverse applications in fluorescence microscopy. <i>Lab on A Chip</i> , 2021, 21, 1549-1562.	3.1	18
35	Functions of the CSB Protein at Topoisomerase 2 Inhibitors-Induced DNA Lesions. <i>Frontiers in Cell and Developmental Biology</i> , 2021, 9, 727836.	1.8	0
36	Multiview confocal super-resolution microscopy. <i>Nature</i> , 2021, 600, 279-284.	13.7	55

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37	Novel deazaflavin tyrosyl-DNA phosphodiesterase 2 (TDP2) inhibitors. <i>DNA Repair</i> , 2020, 85, 102747.	1.3	15
38	SCLC-CellMiner: A Resource for Small Cell Lung Cancer Cell Line Genomics and Pharmacology Based on Genomic Signatures. <i>Cell Reports</i> , 2020, 33, 108296.	2.9	86
39	Debulking of topoisomerase DNA-protein crosslinks (TOP-DPC) by the proteasome, non-proteasomal and non-proteolytic pathways. <i>DNA Repair</i> , 2020, 94, 102926.	1.3	48
40	Resistance to the CHK1 inhibitor prexasertib involves functionally distinct CHK1 activities in BRCA wild-type ovarian cancer. <i>Oncogene</i> , 2020, 39, 5520-5535.	2.6	28
41	The first evidence for SLFN11 expression as an independent prognostic factor for patients with esophageal cancer after chemoradiotherapy. <i>BMC Cancer</i> , 2020, 20, 1123.	1.1	21
42	Recifin A, Initial Example of the Tyr-Lock Peptide Structural Family, Is a Selective Allosteric Inhibitor of Tyrosyl-DNA Phosphodiesterase I. <i>Journal of the American Chemical Society</i> , 2020, 142, 21178-21188.	6.6	7
43	A conserved SUMO pathway repairs topoisomerase DNA-protein cross-links by engaging ubiquitin-mediated proteasomal degradation. <i>Science Advances</i> , 2020, 6, .	4.7	76
44	MGMT Status as a Clinical Biomarker in Glioblastoma. <i>Trends in Cancer</i> , 2020, 6, 380-391.	3.8	131
45	Response to Letter to the Editor by Yang etÂal.. <i>Journal of Thoracic Oncology</i> , 2020, 15, e91.	0.5	0
46	The Indenoisoquinoline LMP517: A Novel Antitumor Agent Targeting both TOP1 and TOP2. <i>Molecular Cancer Therapeutics</i> , 2020, 19, 1589-1597.	1.9	10
47	Topoisomerase I-driven repair of UV-induced damage in NER-deficient cells. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2020, 117, 14412-14420.	3.3	16
48	Excision repair of topoisomerase DNA-protein crosslinks (TOP-DPC). <i>DNA Repair</i> , 2020, 89, 102837.	1.3	62
49	Chromatin Remodeling and Immediate Early Gene Activation by SLFN11 in Response to Replication Stress. <i>Cell Reports</i> , 2020, 30, 4137-4151.e6.	2.9	48
50	BRCAness, SLFN11, and RB1 loss predict response to topoisomerase I inhibitors in triple-negative breast cancers. <i>Science Translational Medicine</i> , 2020, 12, .	5.8	86
51	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. <i>Journal of Thoracic Oncology</i> , 2020, 15, 843-859.	0.5	51
52	BAMscale: quantification of next-generation sequencing peaks and generation of scaled coverage tracks. <i>Epigenetics and Chromatin</i> , 2020, 13, 21.	1.8	40
53	DNA and RNA Cleavage Complexes and Repair Pathway for TOP3B RNA- and DNA-Protein Crosslinks. <i>Cell Reports</i> , 2020, 33, 108569.	2.9	27
54	Acquired SETD2 mutation and impaired CREB1 activation confer cisplatin resistance in metastatic non-small cell lung cancer. <i>Oncogene</i> , 2019, 38, 180-193.	2.6	35

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55	The Indenoisoquinoline TOP1 Inhibitors Selectively Target Homologous Recombination-Deficient and Schlafen 11-Positive Cancer Cells and Synergize with Olaparib. <i>Clinical Cancer Research</i> , 2019, 25, 6206-6216.	3.2	34
56	Beyond the unwinding: role of TOP1MT in mitochondrial translation. <i>Cell Cycle</i> , 2019, 18, 2377-2384.	1.3	11
57	Discovery of Novel Integrase Inhibitors Acting outside the Active Site Through High-Throughput Screening. <i>Molecules</i> , 2019, 24, 3675.	1.7	5
58	Identification of Schlafen-11 as a Target of CD47 Signaling That Regulates Sensitivity to Ionizing Radiation and Topoisomerase Inhibitors. <i>Frontiers in Oncology</i> , 2019, 9, 994.	1.3	22
59	Dual Processing of R-Loops and Topoisomerase I Induces Transcription-Dependent DNA Double-Strand Breaks. <i>Cell Reports</i> , 2019, 28, 3167-3181.e6.	2.9	108
60	Identification of a ligand binding hot spot and structural motifs replicating aspects of tyrosyl-DNA phosphodiesterase I (TDP1) phosphoryl recognition by crystallographic fragment cocktail screening. <i>Nucleic Acids Research</i> , 2019, 47, 10134-10150.	6.5	27
61	Targeting Topoisomerase I in the Era of Precision Medicine. <i>Clinical Cancer Research</i> , 2019, 25, 6581-6589.	3.2	184
62	Mammalian Tyrosyl-DNA Phosphodiesterases in the Context of Mitochondrial DNA Repair. <i>International Journal of Molecular Sciences</i> , 2019, 20, 3015.	1.8	6
63	Synthesis and biological evaluation of 5-aminoethyl benzophenanthridone derivatives as DNA topoisomerase IB inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2019, 178, 81-92.	2.6	11
64	Topoisomerase II-Induced Chromosome Breakage and Translocation Is Determined by Chromosome Architecture and Transcriptional Activity. <i>Molecular Cell</i> , 2019, 75, 252-266.e8.	4.5	145
65	Schlafen 11 (SLFN11), a restriction factor for replicative stress induced by DNA-targeting anti-cancer therapies. , 2019, 201, 94-102.		106
66	Phosphatase 1 Nuclear Targeting Subunit, a Novel DNA Repair Partner of PARP1. <i>Cancer Research</i> , 2019, 79, 2460-2461.	0.4	3
67	Novel Deazaflavin Analogues Potently Inhibited Tyrosyl DNA Phosphodiesterase 2 (TDP2) and Strongly Sensitized Cancer Cells toward Treatment with Topoisomerase II (TOP2) Poison Etoposide. <i>Journal of Medicinal Chemistry</i> , 2019, 62, 4669-4682.	2.9	13
68	The antitumor activity of CYB-L10, a human topoisomerase IB catalytic inhibitor. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2019, 34, 818-822.	2.5	5
69	Mitochondrial tyrosyl-DNA phosphodiesterase 2 and its TDP2 short isoform. <i>EMBO Reports</i> , 2018, 19, .	2.0	19
70	SLFN11 Blocks Stressed Replication Forks Independently of ATR. <i>Molecular Cell</i> , 2018, 69, 371-384.e6.	4.5	177
71	Overcoming Resistance to DNA-Targeted Agents by Epigenetic Activation of Schlafen 11 (SLFN11) Expression with Class I Histone Deacetylase Inhibitors. <i>Clinical Cancer Research</i> , 2018, 24, 1944-1953.	3.2	65
72	PRMT5-mediated arginine methylation of TDP1 for the repair of topoisomerase I covalent complexes. <i>Nucleic Acids Research</i> , 2018, 46, 5601-5617.	6.5	40

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73	New fluorescence-based high-throughput screening assay for small molecule inhibitors of tyrosyl-DNA phosphodiesterase 2 (TDP2). <i>European Journal of Pharmaceutical Sciences</i> , 2018, 118, 67-79.	1.9	14
74	Synthesis, anti-cancer screening and tyrosyl-DNA phosphodiesterase 1 (Tdp1) inhibition activity of novel piperidinyl sulfamides. <i>European Journal of Pharmaceutical Sciences</i> , 2018, 111, 337-348.	1.9	13
75	Application of Sequential Palladium Catalysis for the Discovery of Janus Kinase Inhibitors in the Benzo[<i>c</i>]pyrrolo[2,3- <i>h</i>][1,6]naphthyridin-5-one (BPN) Series. <i>Journal of Medicinal Chemistry</i> , 2018, 61, 10440-10462.	2.9	14
76	CellMinerCDB for Integrative Cross-Database Genomics and Pharmacogenomics Analyses of Cancer Cell Lines. <i>IScience</i> , 2018, 10, 247-264.	1.9	117
77	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. <i>Journal of Medicinal Chemistry</i> , 2018, 61, 9908-9930.	2.9	37
78	Probing the evolutionary conserved residues Y204, F259, S400 and W590 that shape the catalytic groove of human TDP1 for 3'- and 5'-phosphodiester-DNA bond cleavage. <i>DNA Repair</i> , 2018, 66-67, 64-71.	1.3	4
79	NCI Comparative Oncology Program Testing of Non-Camptothecin Indenoisoquinoline Topoisomerase I Inhibitors in Naturally Occurring Canine Lymphoma. <i>Clinical Cancer Research</i> , 2018, 24, 5830-5840.	3.2	36
80	HIV-1 Integrase-Targeted Short Peptides Derived from a Viral Protein R Sequence. <i>Molecules</i> , 2018, 23, 1858.	1.7	3
81	Novel Fluoroindenoisoquinoline Non-Camptothecin Topoisomerase I Inhibitors. <i>Molecular Cancer Therapeutics</i> , 2018, 17, 1694-1704.	1.9	30
82	Endogenous single-strand DNA breaks at RNA polymerase II promoters in <i>Saccharomyces cerevisiae</i> . <i>Nucleic Acids Research</i> , 2018, 46, 10649-10668.	6.5	12
83	Characterization and structure-activity relationships of indenoisoquinoline-derived topoisomerase I inhibitors in unsilencing the dormant <i>Ube3a</i> gene associated with Angelman syndrome. <i>Molecular Autism</i> , 2018, 9, 45.	2.6	28
84	TDP1 suppresses mis-joining of radiomimetic DNA double-strand breaks and cooperates with Artemis to promote optimal nonhomologous end joining. <i>Nucleic Acids Research</i> , 2018, 46, 8926-8939.	6.5	15
85	The evolving landscape of predictive biomarkers of response to PARP inhibitors. <i>Journal of Clinical Investigation</i> , 2018, 128, 1727-1730.	3.9	47
86	Novel screen for anti-cancer drugs that elevate chromosome instability (CIN) using human artificial chromosome (HAC). <i>Oncotarget</i> , 2018, 9, 36833-36835.	0.8	2
87	Novel clinical indenoisoquinoline topoisomerase I inhibitors: a twist around the camptothecins. <i>Oncotarget</i> , 2018, 9, 37286-37288.	0.8	19
88	DNA-Targeted Precision Medicine; Have We Been Caught Sleeping?. <i>Trends in Cancer</i> , 2017, 3, 2-6.	3.8	18
89	A subset of platinum-containing chemotherapeutic agents kills cells by inducing ribosome biogenesis stress. <i>Nature Medicine</i> , 2017, 23, 461-471.	15.2	379
90	Temozolomide in the Era of Precision Medicine. <i>Cancer Research</i> , 2017, 77, 823-826.	0.4	91

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91	Synthesis and Biological Evaluation of the First Triple Inhibitors of Human Topoisomerase 1, Tyrosyl-DNA Phosphodiesterase 1 (Tdp1), and Tyrosyl-DNA Phosphodiesterase 2 (Tdp2). <i>Journal of Medicinal Chemistry</i> , 2017, 60, 3275-3288.	2.9	43
92	The NCI-60 Methylome and Its Integration into CellMiner. <i>Cancer Research</i> , 2017, 77, 601-612.	0.4	48
93	Topoisomerase mediated cleavage at unrepaired ribonucleotides generates DNA double-strand breaks. <i>EMBO Journal</i> , 2017, 36, 361-373.	3.5	59
94	Effects of camptothecin or TOP1 overexpression on genetic stability in <i>Saccharomyces cerevisiae</i> . <i>DNA Repair</i> , 2017, 59, 69-75.	1.3	9
95	HTLV-1 bZIP factor suppresses TDP1 expression through inhibition of NRF-1 in adult T-cell leukemia. <i>Scientific Reports</i> , 2017, 7, 12849.	1.6	13
96	Transcription profiling suggests that mitochondrial topoisomerase IB acts as a topological barrier and regulator of mitochondrial DNA transcription. <i>Journal of Biological Chemistry</i> , 2017, 292, 20162-20172.	1.6	17
97	Distribution bias and biochemical characterization of TOP1MT single nucleotide variants. <i>Scientific Reports</i> , 2017, 7, 8614.	1.6	12
98	Genome Organization Drives Chromosome Fragility. <i>Cell</i> , 2017, 170, 507-521.e18.	13.5	311
99	Structure-Guided Optimization of HIV Integrase Strand Transfer Inhibitors. <i>Journal of Medicinal Chemistry</i> , 2017, 60, 7315-7332.	2.9	44
100	TDP1 is Critical for the Repair of DNA Breaks Induced by Sapacitabine, a Nucleoside also Targeting ATM- and BRCA-Deficient Tumors. <i>Molecular Cancer Therapeutics</i> , 2017, 16, 2543-2551.	1.9	25
101	Identification of Natural Products That Inhibit the Catalytic Function of Human Tyrosyl-DNA Phosphodiesterase (TDP1). <i>SLAS Discovery</i> , 2017, 22, 1093-1105.	1.4	12
102	Design and Synthesis of Chlorinated and Fluorinated 7-Azaindenoisoquinolines as Potent Cytotoxic Anticancer Agents That Inhibit Topoisomerase I. <i>Journal of Medicinal Chemistry</i> , 2017, 60, 5364-5376.	2.9	29
103	Cytidine Deaminase Deficiency Reveals New Therapeutic Opportunities against Cancer. <i>Clinical Cancer Research</i> , 2017, 23, 2116-2126.	3.2	28
104	Phosphorylated fraction of H2AX as a measurement for DNA damage in cancer cells and potential applications of a novel assay. <i>PLoS ONE</i> , 2017, 12, e0171582.	1.1	72
105	ALC1/CHD1L, a chromatin-remodeling enzyme, is required for efficient base excision repair. <i>PLoS ONE</i> , 2017, 12, e0188320.	1.1	34
106	The dominant role of proofreading exonuclease activity of replicative polymerase δ in cellular tolerance to cytarabine (Ara-C). <i>Oncotarget</i> , 2017, 8, 33457-33474.	0.8	24
107	Parallel analysis of ribonucleotide-dependent deletions produced by yeast Top1 <i>in vitro</i> and <i>in vivo</i> . <i>Nucleic Acids Research</i> , 2016, 44, 7714-7721.	6.5	15
108	Epigenetic inactivation of the putative DNA/RNA helicase SLFN11 in human cancer confers resistance to platinum drugs. <i>Oncotarget</i> , 2016, 7, 3084-3097.	0.8	120

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109	Characterization of DNA topoisomerase I in three SN-38 resistant human colon cancer cell lines reveals a new pair of resistance-associated mutations. <i>Journal of Experimental and Clinical Cancer Research</i> , 2016, 35, 56.	3.5	23
110	Clinical and pharmacologic evaluation of two dosing schedules of indotecan (LMP400), a novel indenoisoquinoline, in patients with advanced solid tumors. <i>Cancer Chemotherapy and Pharmacology</i> , 2016, 78, 73-81.	1.1	32
111	RNA Polymerase II Regulates Topoisomerase 1 Activity to Favor Efficient Transcription. <i>Cell</i> , 2016, 165, 357-371.	13.5	211
112	Investigation of the Structure-Activity Relationships of Aza-A-Ring Indenoisoquinoline Topoisomerase I Poisons. <i>Journal of Medicinal Chemistry</i> , 2016, 59, 3840-3853.	2.9	35
113	Design, Synthesis, and Biological Evaluation of Potential Prodrugs Related to the Experimental Anticancer Agent Indotecan (LMP400). <i>Journal of Medicinal Chemistry</i> , 2016, 59, 4890-4899.	2.9	42
114	Deazaflavin Inhibitors of Tyrosyl-DNA Phosphodiesterase 2 (TDP2) Specific for the Human Enzyme and Active against Cellular TDP2. <i>ACS Chemical Biology</i> , 2016, 11, 1925-1933.	1.6	32
115	Selectivity for strand-transfer over 3'-processing and susceptibility to clinical resistance of HIV-1 integrase inhibitors are driven by key enzyme-DNA interactions in the active site. <i>Nucleic Acids Research</i> , 2016, 44, 6896-6906.	6.5	16
116	Novel TDP2-ubiquitin interactions and their importance for the repair of topoisomerase II-mediated DNA damage. <i>Nucleic Acids Research</i> , 2016, 44, gkw719.	6.5	17
117	RNA topoisomerase is prevalent in all domains of life and associates with polyribosomes in animals. <i>Nucleic Acids Research</i> , 2016, 44, 6335-6349.	6.5	63
118	Phenanthriplatin Acts As a Covalent Poison of Topoisomerase II Cleavage Complexes. <i>ACS Chemical Biology</i> , 2016, 11, 2996-3001.	1.6	19
119	Roles of eukaryotic topoisomerases in transcription, replication and genomic stability. <i>Nature Reviews Molecular Cell Biology</i> , 2016, 17, 703-721.	16.1	695
120	Small cell lung cancer: Time to revisit DNA-damaging chemotherapy. <i>Science Translational Medicine</i> , 2016, 8, 346fs12.	5.8	40
121	Laying a trap to kill cancer cells: PARP inhibitors and their mechanisms of action. <i>Science Translational Medicine</i> , 2016, 8, 362ps17.	5.8	518
122	<i>topIb</i> , a phylogenetic hallmark gene of Thaumarchaeota encodes a functional eukaryote-like topoisomerase IB. <i>Nucleic Acids Research</i> , 2016, 44, 2795-2805.	6.5	5
123	F10 cytotoxicity via topoisomerase I cleavage complex repair consistent with a unique mechanism for thymineless death. <i>Future Oncology</i> , 2016, 12, 2183-2188.	1.1	10
124	Chromatin Regulators as a Guide for Cancer Treatment Choice. <i>Molecular Cancer Therapeutics</i> , 2016, 15, 1768-1777.	1.9	18
125	rCellMiner: exploring molecular profiles and drug response of the NCI-60 cell lines in R. <i>Bioinformatics</i> , 2016, 32, 1272-1274.	1.8	39
126	HIV-1 Integrase Strand Transfer Inhibitors with Reduced Susceptibility to Drug Resistant Mutant Integrases. <i>ACS Chemical Biology</i> , 2016, 11, 1074-1081.	1.6	35

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127	Isoquinoline-1,3-diones as Selective Inhibitors of Tyrosyl DNA Phosphodiesterase II (TDP2). <i>Journal of Medicinal Chemistry</i> , 2016, 59, 2734-2746.	2.9	52
128	Synthesis and biological evaluation of new fluorinated and chlorinated indenoisoquinoline topoisomerase I poisons. <i>Bioorganic and Medicinal Chemistry</i> , 2016, 24, 1469-1479.	1.4	22
129	Resistance to PARP inhibitors by SLFN11 inactivation can be overcome by ATR inhibition. <i>Oncotarget</i> , 2016, 7, 76534-76550.	0.8	219
130	Analogues of the novel phytohormone, strigolactone, trigger apoptosis and synergize with PARP inhibitors by inducing DNA damage and inhibiting DNA repair. <i>Oncotarget</i> , 2016, 7, 13984-14001.	0.8	30
131	Camptothecin targets WRN protein: mechanism and relevance in clinical breast cancer. <i>Oncotarget</i> , 2016, 7, 13269-13284.	0.8	38
132	Relative contribution of four nucleases, CtIP, Dna2, Exo1 and Mre11, to the initial step of DNA double-strand break repair by homologous recombination in both the chicken DT40 and human TK6 cell lines. <i>Genes To Cells</i> , 2015, 20, 1059-1076.	0.5	46
133	Activation of RAF1 (c-RAF) by the Marine Alkaloid Lasonolide A Induces Rapid Premature Chromosome Condensation. <i>Marine Drugs</i> , 2015, 13, 3625-3639.	2.2	15
134	Topoisomerase I Alone Is Sufficient to Produce Short DNA Deletions and Can Also Reverse Nicks at Ribonucleotide Sites. <i>Journal of Biological Chemistry</i> , 2015, 290, 14068-14076.	1.6	52
135	Anti-HIV-1 activity of a tripodal receptor that recognizes mannose oligomers. <i>European Journal of Medicinal Chemistry</i> , 2015, 106, 132-143.	2.6	10
136	Single-Molecule Supercoil Relaxation Assay as a Screening Tool to Determine the Mechanism and Efficacy of Human Topoisomerase IB Inhibitors. <i>Molecular Cancer Therapeutics</i> , 2015, 14, 2552-2559.	1.9	13
137	Alterations of DNA repair genes in the NCI-60 cell lines and their predictive value for anticancer drug activity. <i>DNA Repair</i> , 2015, 28, 107-115.	1.3	55
138	Using CellMiner 1.6 for Systems Pharmacology and Genomic Analysis of the NCI-60. <i>Clinical Cancer Research</i> , 2015, 21, 3841-3852.	3.2	80
139	Interfacial inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2015, 25, 3961-3965.	1.0	34
140	Synthesis and biological evaluation of 6-substituted indolizinoquinolinediones as catalytic DNA topoisomerase I inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2015, 101, 525-533.	2.6	15
141	Discovery of Potent Indenoisoquinoline Topoisomerase I Poisons Lacking the 3-Nitro Toxicophore. <i>Journal of Medicinal Chemistry</i> , 2015, 58, 3997-4015.	2.9	40
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