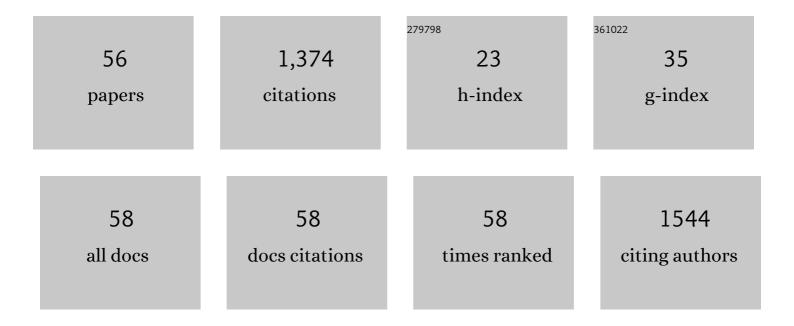
Robert L Eoff

List of Publications by Year in descending order

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POREDT L FOEE

#	Article	IF	CITATIONS
1	323 Generation of a functional precision medicine pipeline which combines comparative transcriptomics and tumor organoid modeling to identify bespoke treatment strategies for glioblastoma. Journal of Clinical and Translational Science, 2022, 6, 58-58.	0.6	0
2	Monensin and its analogues show antiâ€glioblastoma activity in an organoid model of cancer. FASEB Journal, 2022, 36, .	0.5	0
3	Human Rev1 relies on insert-2 to promote selective binding and accurate replication of stabilized G-quadruplex motifs. Nucleic Acids Research, 2021, 49, 2065-2084.	14.5	13
4	Inhibition of tryptophan 2,3-dioxygenase impairs DNA damage tolerance and repair in glioma cells. NAR Cancer, 2021, 3, zcab014.	3.1	10
5	DNA Polymerase Kappa Acts as a Barrier to Unrestrained Replication in Glioblastoma. FASEB Journal, 2021, 35, .	0.5	0
6	Deletion of putative xenobiotic response elements (XREs) in hpol κ alters the replication stress response and overall genomic instability in glioblastoma cells. FASEB Journal, 2021, 35, .	0.5	0
7	Selective Binding Of Human Rev1 With Gâ€Quadruplex DNA Is Determined By A Region Unique to Higher Eukaryotes. FASEB Journal, 2021, 35, .	0.5	0
8	Single and double modified salinomycin analogs target stem-like cells in 2D and 3D breast cancer models. Biomedicine and Pharmacotherapy, 2021, 141, 111815.	5.6	7
9	Inositol serves as a natural inhibitor of mitochondrial fission by directly targeting AMPK. Molecular Cell, 2021, 81, 3803-3819.e7.	9.7	39
10	Biobanked Glioblastoma Patient-Derived Organoids as a Precision Medicine Model to Study Inhibition of Invasion. International Journal of Molecular Sciences, 2021, 22, 10720.	4.1	11
11	Site-Specific Synthesis of Oligonucleotides Containing 6-Oxo-M ₁ dG, the Genomic Metabolite of M ₁ dG, and Liquid Chromatography–Tandem Mass Spectrometry Analysis of Its In Vitro Bypass by Human Polymerase I¹. Chemical Research in Toxicology, 2021, 34, 2567-2578.	3.3	2
12	A Functional Precision Medicine Pipeline Combines Comparative Transcriptomics and Tumor Organoid Modeling to Identify Bespoke Treatment Strategies for Glioblastoma. Cells, 2021, 10, 3400.	4.1	15
13	A Facile Semisynthesis and Evaluation of Garcinoic Acid and Its Analogs for the Inhibition of Human DNA Polymerase β. Molecules, 2020, 25, 5847.	3.8	2
14	Novel Salinomycin Analogs Show Improved Selectivity Towards Breast Cancer Stem Cells. FASEB Journal, 2020, 34, 1-1.	0.5	0
15	Inhibition of Human DNA Polymerases Eta and Kappa by Indole-Derived Molecules Occurs through Distinct Mechanisms. ACS Chemical Biology, 2019, 14, 1337-1351.	3.4	18
16	LC8/DYNLL1 is a 53BP1 effector and regulates checkpoint activation. Nucleic Acids Research, 2019, 47, 6236-6249.	14.5	34
17	A Small-Molecule Inhibitor of Human DNA Polymerase η Potentiates the Effects of Cisplatin in Tumor Cells. Biochemistry, 2018, 57, 1262-1273.	2.5	27
18	Synthesis and Evaluation of 2-Naphthaleno trans-Stilbenes and Cyanostilbenes as Anticancer Agents. Anti-Cancer Agents in Medicinal Chemistry, 2018, 18, 556-564.	1.7	7

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19	Residues in the RecQ C-terminal Domain of the Human Werner Syndrome Helicase Are Involved in Unwinding G-quadruplex DNA. Journal of Biological Chemistry, 2017, 292, 3154-3163.	3.4	19
20	Translesion DNA Synthesis in Cancer: Molecular Mechanisms and Therapeutic Opportunities. Chemical Research in Toxicology, 2017, 30, 1942-1955.	3.3	37
21	A catch and release program for single-stranded DNA. Journal of Biological Chemistry, 2017, 292, 13085-13086.	3.4	3
22	Evidence That G-quadruplex DNA Accumulates in the Cytoplasm and Participates in Stress Granule Assembly in Response to Oxidative Stress. Journal of Biological Chemistry, 2016, 291, 18041-18057.	3.4	71
23	Dioxol and dihydrodioxin analogs of 2- and 3-phenylacetonitriles as potent anti-cancer agents with nanomolar activity against a variety of human cancer cells. Bioorganic and Medicinal Chemistry Letters, 2016, 26, 2164-2169.	2.2	9
24	Aberrant Kynurenine Signaling Modulates DNA Replication Stress Factors and Promotes Genomic Instability in Gliomas. Chemical Research in Toxicology, 2016, 29, 1369-1380.	3.3	11
25	Human Translesion Polymerase κ Exhibits Enhanced Activity and Reduced Fidelity Two Nucleotides from G-Quadruplex DNA. Biochemistry, 2016, 55, 5218-5229.	2.5	36
26	Antinociceptive effects of the 6- O -sulfate ester of morphine in normal and diabetic rats: Comparative role of mu- and delta-opioid receptors. Pharmacological Research, 2016, 113, 335-347.	7.1	21
27	Effects of Twelve Germline Missense Variations on DNA Lesion and G-Quadruplex Bypass Activities of Human DNA Polymerase REV1. Chemical Research in Toxicology, 2016, 29, 367-379.	3.3	12
28	Kynurenine Signaling Increases DNA Polymerase Kappa Expression and Promotes Genomic Instability in Glioblastoma Cells. Chemical Research in Toxicology, 2016, 29, 101-108.	3.3	27
29	Evidence for the Kinetic Partitioning of Polymerase Activity on G-Quadruplex DNA. Biochemistry, 2015, 54, 3218-3230.	2.5	37
30	Synthesis, anticancer activity and molecular docking studies on a series of heterocyclic trans-cyanocombretastatin analogues as antitubulin agents. European Journal of Medicinal Chemistry, 2015, 92, 212-220.	5.5	18
31	Synthesis and evaluation of a series of resveratrol analogues as potent anti-cancer agents that target tubulin. MedChemComm, 2015, 6, 788-794.	3.4	31
32	Synthesis and anti-cancer screening of novel heterocyclic-(2H)-1,2,3-triazoles as potential anti-cancer agents. MedChemComm, 2015, 6, 1535-1543.	3.4	49
33	1-Benzyl-2-methyl-3-indolylmethylene barbituric acid derivatives: Anti-cancer agents that target nucleophosmin 1 (NPM1). Bioorganic and Medicinal Chemistry, 2015, 23, 7226-7233.	3.0	35
34	Synthesis and biological evaluation of novel 4,5-disubstituted 2H-1,2,3-triazoles as cis-constrained analogues of combretastatin A-4. European Journal of Medicinal Chemistry, 2015, 103, 123-132.	5.5	56
35	Human Rev1 polymerase disrupts G-quadruplex DNA. Nucleic Acids Research, 2014, 42, 3272-3285.	14.5	62
36	The Werner syndrome protein limits the error-prone 8-oxo-dG lesion bypass activity of human DNA polymerase kappa. Nucleic Acids Research, 2014, 42, 12027-12040.	14.5	11

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37	<i>N</i> -Aroyl Indole Thiobarbituric Acids as Inhibitors of DNA Repair and Replication Stress Response Polymerases. ACS Chemical Biology, 2013, 8, 1722-1729.	3.4	25
38	Leukotriene Biosynthesis Inhibitor MK886 Impedes DNA Polymerase Activity. Chemical Research in Toxicology, 2013, 26, 221-232.	3.3	17
39	Identification and characterization of novel small molecule inhibitors of the human Yâ€family DNA polymerases. FASEB Journal, 2013, 27, 543.1.	0.5	0
40	Enhancement of Human DNA Polymerase η Activity and Fidelity Is Dependent Upon a Bipartite Interaction with the Werner Syndrome Protein. Journal of Biological Chemistry, 2012, 287, 42312-42323.	3.4	14
41	A Comprehensive Strategy to Discover Inhibitors of the Translesion Synthesis DNA Polymerase κ. PLoS ONE, 2012, 7, e45032.	2.5	32
42	Roles of the Four DNA Polymerases of the Crenarchaeon Sulfolobus solfataricus and Accessory Proteins in DNA Replication. Journal of Biological Chemistry, 2011, 286, 31180-31193.	3.4	51
43	Selective Modulation of DNA Polymerase Activity by Fixedâ€Conformation Nucleoside Analogues. Angewandte Chemie, 2010, 122, 7643-7647.	2.0	1
44	Selective Modulation of DNA Polymerase Activity by Fixedâ€Conformation Nucleoside Analogues. Angewandte Chemie - International Edition, 2010, 49, 7481-7485.	13.8	15
45	Mechanistic Studies with DNA Polymerases Reveal Complex Outcomes following Bypass of DNA Damage. Journal of Nucleic Acids, 2010, 2010, 1-12.	1.2	9
46	<i>In Vitro</i> Bypass of the Major Malondialdehyde- and Base Propenal-Derived DNA Adduct by Human Y-family DNA Polymerases l̂², l̂¹, and Rev1. Biochemistry, 2010, 49, 8415-8424.	2.5	24
47	Kinetic Mechanism for DNA Unwinding by Multiple Molecules of Dda Helicase Aligned on DNA. Biochemistry, 2010, 49, 4543-4553.	2.5	21
48	Conformational Changes during Nucleotide Selection by Sulfolobus solfataricus DNA Polymerase Dpo4. Journal of Biological Chemistry, 2009, 284, 21090-21099.	3.4	23
49	Structural and Functional Elucidation of the Mechanism Promoting Error-prone Synthesis by Human DNA Polymerase lº Opposite the 7,8-Dihydro-8-oxo-2′-deoxyguanosine Adduct. Journal of Biological Chemistry, 2009, 284, 22467-22480.	3.4	78
50	Translesion DNA Synthesis by Human DNA Polymerase Î∙ on Templates Containing a Pyrimidopurinone Deoxyguanosine Adduct, 3-(2′-Deoxy-β-d-erythro-pentofuranosyl)pyrimido-[1,2-a]purin-10(3H)-one. Biochemistry, 2009, 48, 471-480.	2.5	15
51	Structural and Functional Analysis of <i>Sulfolobus solfataricus</i> Y-Family DNA Polymerase Dpo4-Catalyzed Bypass of the Malondialdehydeâ^Deoxyguanosine Adduct [,] . Biochemistry, 2009, 48, 7079-7088.	2.5	30
52	Molecular Basis of Selectivity of Nucleoside Triphosphate Incorporation Opposite O6-Benzylguanine by Sulfolobus solfataricus DNA Polymerase Dpo4. Journal of Biological Chemistry, 2007, 282, 13573-13584.	3.4	58
53	Sulfolobus solfataricus DNA Polymerase Dpo4 Is Partially Inhibited by "Wobble―Pairing between O6-Methylguanine and Cytosine, but Accurate Bypass Is Preferred. Journal of Biological Chemistry, 2007, 282, 1456-1467.	3.4	79
54	Hydrogen Bonding of 7,8-Dihydro-8-oxodeoxyguanosine with a Charged Residue in the Little Finger Domain Determines Miscoding Events in Sulfolobus solfataricus DNA Polymerase Dpo4. Journal of Biological Chemistry, 2007, 282, 19831-19843.	3.4	71

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55	Intermediates revealed in the kinetic mechanism for DNA unwinding by a monomeric helicase. Nature Structural and Molecular Biology, 2006, 13, 242-249.	8.2	46
56	Chemically Modified DNA Substrates Implicate the Importance of Electrostatic Interactions for DNA Unwinding by Dda Helicase. Biochemistry, 2005, 44, 666-674.	2.5	27