

# Stanley T Crooke

## List of Publications by Year in descending order

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211  
papers

14,636  
citations

17405

63  
h-index

21474

114  
g-index

214  
all docs

214  
docs citations

214  
times ranked

10495  
citing authors

#	ARTICLE	IF	CITATIONS
1	Mipomersen, an apolipoprotein B synthesis inhibitor, for lowering of LDL cholesterol concentrations in patients with homozygous familial hypercholesterolaemia: a randomised, double-blind, placebo-controlled trial. <i>Lancet, The</i> , 2010, 375, 998-1006.	6.3	813
2	Antisense oligonucleotides targeting apolipoprotein(a) in people with raised lipoprotein(a): two randomised, double-blind, placebo-controlled, dose-ranging trials. <i>Lancet, The</i> , 2016, 388, 2239-2253.	6.3	584
3	RNA-Targeted Therapeutics. <i>Cell Metabolism</i> , 2018, 27, 714-739.	7.2	556
4	Mitomycin C: a review. <i>Cancer Treatment Reviews</i> , 1976, 3, 121-139.	3.4	430
5	Cellular uptake and trafficking of antisense oligonucleotides. <i>Nature Biotechnology</i> , 2017, 35, 230-237.	9.4	416
6	Efficient Reduction of Target RNAs by Small Interfering RNA and RNase H-dependent Antisense Agents. <i>Journal of Biological Chemistry</i> , 2003, 278, 7108-7118.	1.6	403
7	Antisense Oligonucleotide Inhibition of Apolipoprotein C-III Reduces Plasma Triglycerides in Rodents, Nonhuman Primates, and Humans. <i>Circulation Research</i> , 2013, 112, 1479-1490.	2.0	326
8	Antisense technology: an overview and prospectus. <i>Nature Reviews Drug Discovery</i> , 2021, 20, 427-453.	21.5	299
9	Molecular cloning and complete amino-acid sequence of form-I phosphoinositide-specific phospholipase C. <i>Nature</i> , 1988, 334, 268-270.	13.7	266
10	Antisense inhibition of proprotein convertase subtilisin/kexin type 9 reduces serum LDL in hyperlipidemic mice. <i>Journal of Lipid Research</i> , 2007, 48, 763-767.	2.0	266
11	Molecular Mechanisms of Antisense Oligonucleotides. <i>Nucleic Acid Therapeutics</i> , 2017, 27, 70-77.	2.0	255
12	Determination of the Role of the Human RNase H1 in the Pharmacology of DNA-like Antisense Drugs. <i>Journal of Biological Chemistry</i> , 2004, 279, 17181-17189.	1.6	250
13	Single-Stranded RNAs Use RNAi to Potently and Allele-Selectively Inhibit Mutant Huntingtin Expression. <i>Cell</i> , 2012, 150, 895-908.	13.5	250
14	Single-Stranded siRNAs Activate RNAi in Animals. <i>Cell</i> , 2012, 150, 883-894.	13.5	239
15	Correlation of the in vitro cytotoxic and in vivo antitumor activities of gold(I) coordination complexes. <i>Journal of Medicinal Chemistry</i> , 1986, 29, 218-223.	2.9	222
16	Chemical modification of PS-ASO therapeutics reduces cellular protein-binding and improves the therapeutic index. <i>Nature Biotechnology</i> , 2019, 37, 640-650.	9.4	205
17	RNase H1-Dependent Antisense Oligonucleotides Are Robustly Active in Directing RNA Cleavage in Both the Cytoplasm and the Nucleus. <i>Molecular Therapy</i> , 2017, 25, 2075-2092.	3.7	168
18	Rapid identification and strain-typing of respiratory pathogens for epidemic surveillance. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2005, 102, 8012-8017.	3.3	165

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19	Human RNase III Is a 160-kDa Protein Involved in Preribosomal RNA Processing. <i>Journal of Biological Chemistry</i> , 2000, 275, 36957-36965.	1.6	164
20	Phosphorothioate modified oligonucleotide-protein interactions. <i>Nucleic Acids Research</i> , 2020, 48, 5235-5253.	6.5	163
21	Antisense Strategies. <i>Current Molecular Medicine</i> , 2004, 4, 465-487.	0.6	157
22	Etoposide (VP-16-213). <i>Cancer Treatment Reviews</i> , 1979, 6, 107-124.	3.4	155
23	Antitumor activity of bis(diphenylphosphino)alkanes, their gold(I) coordination complexes, and related compounds. <i>Journal of Medicinal Chemistry</i> , 1987, 30, 2181-2190.	2.9	155
24	Identification and characterization of intracellular proteins that bind oligonucleotides with phosphorothioate linkages. <i>Nucleic Acids Research</i> , 2015, 43, 2927-2945.	6.5	151
25	Antisense technology: A review. <i>Journal of Biological Chemistry</i> , 2021, 296, 100416.	1.6	149
26	Properties of Cloned and Expressed Human RNase H1. <i>Journal of Biological Chemistry</i> , 1999, 274, 28270-28278.	1.6	148
27	Maytansine. <i>Cancer Treatment Reviews</i> , 1978, 5, 199-207.	3.4	145
28	Clinical pharmacological properties of mipomersen (Kynamro), a second generation antisense inhibitor of apolipoprotein B. <i>British Journal of Clinical Pharmacology</i> , 2013, 76, 269-276.	1.1	145
29	Hepatotoxicity of high affinity gapmer antisense oligonucleotides is mediated by RNase H1 dependent promiscuous reduction of very long pre-mRNA transcripts. <i>Nucleic Acids Research</i> , 2016, 44, 2093-2109.	6.5	142
30	Progress in antisense technology: The end of the beginning. <i>Methods in Enzymology</i> , 2000, 313, 3-45.	0.4	141
31	Clinical and Preclinical Pharmacokinetics and Pharmacodynamics of Mipomersen (Kynamro®): A Second-Generation Antisense Oligonucleotide Inhibitor of Apolipoprotein B. <i>Clinical Pharmacokinetics</i> , 2015, 54, 133-146.	1.6	138
32	Translation efficiency of mRNAs is increased by antisense oligonucleotides targeting upstream open reading frames. <i>Nature Biotechnology</i> , 2016, 34, 875-880.	9.4	137
33	Binding Affinity and Specificity of Escherichia coli RNase H1: Impact on the Kinetics of Catalysis of Antisense Oligonucleotide-RNA Hybrids. <i>Biochemistry</i> , 1997, 36, 390-398.	1.2	127
34	Progress toward oligonucleotide therapeutics: pharmacodynamic properties. <i>FASEB Journal</i> , 1993, 7, 533-539.	0.2	118
35	An Overview of Progress in Antisense Therapeutics. <i>Oligonucleotides</i> , 1998, 8, 115-122.	4.4	106
36	Binding and Cleavage Specificities of Human Argonaute2. <i>Journal of Biological Chemistry</i> , 2009, 284, 26017-26028.	1.6	104

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37	Peptidoleukotrienes: Distinct receptors for leukotriene C4 and D4 in the Guinea-pig lung. <i>Biochemical and Biophysical Research Communications</i> , 1983, 116, 1136-1143.	1.0	103
38	The Effects of 2'-O-Methoxyethyl Containing Antisense Oligonucleotides on Platelets in Human Clinical Trials. <i>Nucleic Acid Therapeutics</i> , 2017, 27, 121-129.	2.0	101
39	2'-Fluoro-modified phosphorothioate oligonucleotide can cause rapid degradation of P54nrb and PSF. <i>Nucleic Acids Research</i> , 2015, 43, 4569-4578.	6.5	97
40	Cellular association, intracellular distribution, and efflux of auranofin via sequential ligand exchange reactions. <i>Biochemical Pharmacology</i> , 1986, 35, 923-932.	2.0	95
41	Rapid Identification of Emerging Pathogens: Coronavirus. <i>Emerging Infectious Diseases</i> , 2005, 11, 373-379.	2.0	94
42	Molecular Mechanisms of Antisense Drugs: RNase H. Oligonucleotides, 1998, 8, 133-134.	4.4	92
43	The signal transduction system of the leukotriene D4 receptor. <i>Trends in Pharmacological Sciences</i> , 1989, 10, 103-107.	4.0	91
44	Integrated Safety Assessment of 2'-O-Methoxyethyl Chimeric Antisense Oligonucleotides in NonHuman Primates and Healthy Human Volunteers. <i>Molecular Therapy</i> , 2016, 24, 1771-1782.	3.7	91
45	Phosphorothioate oligonucleotides can displace NEAT1 RNA and form nuclear paraspeckle-like structures. <i>Nucleic Acids Research</i> , 2014, 42, 8648-8662.	6.5	87
46	Integrated Assessment of the Clinical Performance of GalNAc <sub>3</sub> -Conjugated 2'-O-Methoxyethyl Chimeric Antisense Oligonucleotides: I. Human Volunteer Experience. <i>Nucleic Acid Therapeutics</i> , 2019, 29, 16-32.	2.0	85
47	Vitravene, "Another Piece in the Mosaic. Oligonucleotides, 1998, 8, vii-viii.	4.4	84
48	Viable RNaseH1 knockout mice show RNaseH1 is essential for R loop processing, mitochondrial and liver function. <i>Nucleic Acids Research</i> , 2016, 44, 5299-5312.	6.5	84
49	Reduced levels of Ago2 expression result in increased siRNA competition in mammalian cells. <i>Nucleic Acids Research</i> , 2007, 35, 6598-6610.	6.5	83
50	Antisense oligonucleotides targeting translation inhibitory elements in 5' UTRs can selectively increase protein levels. <i>Nucleic Acids Research</i> , 2017, 45, 9528-9546.	6.5	83
51	Human RNase H1 Discriminates between Subtle Variations in the Structure of the Heteroduplex Substrate. <i>Molecular Pharmacology</i> , 2007, 71, 83-91.	1.0	82
52	Cloning and expression of a yeast copper metallothionein gene. <i>Gene</i> , 1984, 27, 23-33.	1.0	81
53	TCP1 complex proteins interact with phosphorothioate oligonucleotides and can co-localize in oligonucleotide-induced nuclear bodies in mammalian cells. <i>Nucleic Acids Research</i> , 2014, 42, 7819-7832.	6.5	80
54	Multiplexed Screening of Neutral Mass-Tagged RNA Targets against Ligand Libraries with Electrospray Ionization FTICR MS: A Paradigm for High-Throughput Affinity Screening. <i>Analytical Chemistry</i> , 1999, 71, 3436-3440.	3.2	78

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55	Site-specific replacement of phosphorothioate with alkyl phosphonate linkages enhances the therapeutic profile of gapmer ASOs by modulating interactions with cellular proteins. <i>Nucleic Acids Research</i> , 2019, 47, 5465-5479.	6.5	77
56	The Interaction of Phosphorothioate-Containing RNA Targeted Drugs with Proteins Is a Critical Determinant of the Therapeutic Effects of These Agents. <i>Journal of the American Chemical Society</i> , 2020, 142, 14754-14771.	6.6	77
57	Role of intravesical mitomycin c in management of superficial bladder tumors. <i>Urology</i> , 1980, 16, 11-15.	0.5	76
58	A pharmacokinetic evaluation of <sup>14</sup> C-labeled afovirsen sodium in patients with genital warts. <i>Clinical Pharmacology and Therapeutics</i> , 1994, 56, 641-646.	2.3	75
59	Potential roles of antisense technology in cancer chemotherapy. <i>Oncogene</i> , 2000, 19, 6651-6659.	2.6	74
60	Investigating the Structure of Human RNase H1 by Site-directed Mutagenesis. <i>Journal of Biological Chemistry</i> , 2001, 276, 23547-23553.	1.6	74
61	Structural Requirements at the Catalytic Site of the Heteroduplex Substrate for Human RNase H1 Catalysis. <i>Journal of Biological Chemistry</i> , 2004, 279, 36317-36326.	1.6	72
62	Acute hepatotoxicity of 2'-fluoro-modified 5'-10' gapmer phosphorothioate oligonucleotides in mice correlates with intracellular protein binding and the loss of DBHS proteins. <i>Nucleic Acids Research</i> , 2018, 46, 2204-2217.	6.5	71
63	Identification of metabolically stable 5'-phosphate analogs that support single-stranded siRNA activity. <i>Nucleic Acids Research</i> , 2015, 43, 2993-3011.	6.5	67
64	Hsp90 protein interacts with phosphorothioate oligonucleotides containing hydrophobic 2'-modifications and enhances antisense activity. <i>Nucleic Acids Research</i> , 2016, 44, 3892-3907.	6.5	65
65	Bleomycin serum pharmacokinetics as determined by a radioimmunoassay and a microbiologic assay in a patient with compromised renal function. <i>Cancer</i> , 1977, 39, 1430-1434.	2.0	63
66	Understanding the effect of controlling phosphorothioate chirality in the DNA gap on the potency and safety of gapmer antisense oligonucleotides. <i>Nucleic Acids Research</i> , 2020, 48, 1691-1700.	6.5	63
67	Interactions of gold coordination complexes with DNA. <i>Biochemical Pharmacology</i> , 1986, 35, 1427-1433.	2.0	62
68	Vascular vasopressin receptors mediate phosphatidylinositol turnover and calcium efflux in an established smooth muscle cell line. <i>Life Sciences</i> , 1986, 39, 37-45.	2.0	62
69	Efficient and specific knockdown of small non-coding RNAs in mammalian cells and in mice. <i>Nucleic Acids Research</i> , 2011, 39, e13-e13.	6.5	62
70	Effect of leukotrienes, bradykinin and calcium ionophore (A 23187) on bovine endothelial cells: Release of prostacyclin. <i>Prostaglandins</i> , 1986, 31, 157-166.	1.2	61
71	Towards next generation antisense oligonucleotides: mesylphosphoramidate modification improves therapeutic index and duration of effect of gapmer antisense oligonucleotides. <i>Nucleic Acids Research</i> , 2021, 49, 9026-9041.	6.5	61
72	Intra-endosomal trafficking mediated by lysobisphosphatidic acid contributes to intracellular release of phosphorothioate-modified antisense oligonucleotides. <i>Nucleic Acids Research</i> , 2017, 45, 5309-5322.	6.5	60

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73	The rates of the major steps in the molecular mechanism of RNase H1-dependent antisense oligonucleotide induced degradation of RNA. <i>Nucleic Acids Research</i> , 2015, 43, 8955-8963.	6.5	59
74	Annexin A2 facilitates endocytic trafficking of antisense oligonucleotides. <i>Nucleic Acids Research</i> , 2016, 44, gkw595.	6.5	58
75	Cellular uptake mediated by epidermal growth factor receptor facilitates the intracellular activity of phosphorothioate-modified antisense oligonucleotides. <i>Nucleic Acids Research</i> , 2018, 46, 3579-3594.	6.5	58
76	Molecular mechanisms of action of auranofin and other gold complexes as related to their biologic activities. <i>American Journal of Medicine</i> , 1983, 75, 109-113.	0.6	57
77	Cleavage of Single Strand RNA Adjacent to RNA-DNA Duplex Regions by <i>Escherichia coli</i> RNase H1. <i>Journal of Biological Chemistry</i> , 1997, 272, 27513-27516.	1.6	57
78	Development of a Quantitative BRET Affinity Assay for Nucleic Acid-Protein Interactions. <i>PLoS ONE</i> , 2016, 11, e0161930.	1.1	57
79	RNA cleavage products generated by antisense oligonucleotides and siRNAs are processed by the RNA surveillance machinery. <i>Nucleic Acids Research</i> , 2016, 44, 3351-3363.	6.5	57
80	Antisense Oligonucleotides Capable of Promoting Specific Target mRNA Reduction via Competing RNase H1-Dependent and Independent Mechanisms. <i>PLoS ONE</i> , 2014, 9, e108625.	1.1	56
81	Origins of the Increased Affinity of Phosphorothioate-Modified Therapeutic Nucleic Acids for Proteins. <i>Journal of the American Chemical Society</i> , 2020, 142, 7456-7468.	6.6	56
82	Identification and characterization of leukotriene D4 receptors in adult and fetal human lung. <i>Biochemical Pharmacology</i> , 1985, 34, 4311-4317.	2.0	55
83	The Effects of 2'-Methoxyethyl Oligonucleotides on Renal Function in Humans. <i>Nucleic Acid Therapeutics</i> , 2018, 28, 10-22.	2.0	55
84	Antisense drug discovery and development technology considered in a pharmacological context. <i>Biochemical Pharmacology</i> , 2021, 189, 114196.	2.0	55
85	Molecular Cloning and Expression of cDNA for Human RNase H. <i>Oligonucleotides</i> , 1998, 8, 53-61.	4.4	54
86	Differential effects of manoalide on secreted and intracellular phospholipases. <i>Biochemical Pharmacology</i> , 1987, 36, 733-740.	2.0	53
87	Thiol competition for Et3PAuS-albumin: a nonenzymatic mechanism for Et3PO formation. <i>Journal of Inorganic Biochemistry</i> , 1987, 30, 177-187.	1.5	52
88	The cellular pharmacology of auranofin. <i>Seminars in Arthritis and Rheumatism</i> , 1987, 17, 71-80.	1.6	51
89	Advances in Understanding the Pharmacological Properties of Antisense Oligonucleotides. <i>Advances in Pharmacology</i> , 1997, 40, 1-49.	1.2	51
90	Identification and Partial Purification of Human Double Strand RNase Activity. <i>Journal of Biological Chemistry</i> , 1998, 273, 2532-2542.	1.6	50

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91	Depletion of key protein components of the RISC pathway impairs pre-ribosomal RNA processing. <i>Nucleic Acids Research</i> , 2011, 39, 4875-4889.	6.5	50
92	Nucleic acid binding proteins affect the subcellular distribution of phosphorothioate antisense oligonucleotides. <i>Nucleic Acids Research</i> , 2017, 45, 10649-10671.	6.5	50
93	Antisense Therapeutics. <i>Biotechnology and Genetic Engineering Reviews</i> , 1998, 15, 121-158.	2.4	48
94	Lipid Conjugates Enhance Endosomal Release of Antisense Oligonucleotides Into Cells. <i>Nucleic Acid Therapeutics</i> , 2019, 29, 245-255.	2.0	48
95	Low-Molecular-Weight Nuclear RNAs. <i>Perspectives in Biology and Medicine</i> , 1971, 15, 117-139.	0.3	47
96	The Influence of Antisense Oligonucleotide-induced RNA Structure on Escherichia coli RNase H1 Activity. <i>Journal of Biological Chemistry</i> , 1997, 272, 18191-18199.	1.6	47
97	Hepatic distribution of a phosphorothioate oligodeoxynucleotide within rodents following intravenous administration <sup>212</sup> Abbreviations: HSV, herpes simplex virus; CGE, capillary gel electrophoresis; and HPLC/ES-MS, high performance liquid chromatography/electrospray-mass spectrometry. Johnston J, ISIS Pharmaceuticals, personal communication. Cited with permission. <i>Biochemical Pharmacology</i> , 2001, 62, 297-306.	2.0	47
98	RNA Therapeutics in Oncology: Advances, Challenges, and Future Directions. <i>Journal of Clinical Pharmacology</i> , 2017, 57, S43-S59.	1.0	46
99	Progress in antisense therapeutics. , 1996, 16, 319-344.		45
100	Defining the Factors That Contribute to On-Target Specificity of Antisense Oligonucleotides. <i>PLoS ONE</i> , 2014, 9, e101752.	1.1	45
101	Dynamic nucleoplasmic and nucleolar localization of mammalian RNase H1 in response to RNAP I transcriptional R-loops. <i>Nucleic Acids Research</i> , 2017, 45, 10672-10692.	6.5	44
102	Human RNase H1 Is Associated with Protein P32 and Is Involved in Mitochondrial Pre-rRNA Processing. <i>PLoS ONE</i> , 2013, 8, e71006.	1.1	43
103	Human Dicer Binds Short Single-strand and Double-strand RNA with High Affinity and Interacts with Different Regions of the Nucleic Acids. <i>Journal of Biological Chemistry</i> , 2009, 284, 2535-2548.	1.6	42
104	Lipid Nanoparticles Improve Activity of Single-Stranded siRNA and Gapmer Antisense Oligonucleotides in Animals. <i>ACS Chemical Biology</i> , 2013, 8, 1402-1406.	1.6	41
105	Application of a tissue culture microtiter test for the detection of cytotoxic agents from natural products.. <i>Journal of Antibiotics</i> , 1985, 38, 758-766.	1.0	40
106	Identification of specific binding sites for leukotriene C4 in human fetal lung. <i>Prostaglandins</i> , 1984, 27, 961-974.	1.2	39
107	Human RNase H1 Uses One Tryptophan and Two Lysines to Position the Enzyme at the 3' end-DNA/5' end-RNA Terminus of the Heteroduplex Substrate. <i>Journal of Biological Chemistry</i> , 2003, 278, 49860-49867.	1.6	39
108	Transfection of siRNAs can alter miRNA levels and trigger non-specific protein degradation in mammalian cells. <i>Biochimica Et Biophysica Acta - Gene Regulatory Mechanisms</i> , 2013, 1829, 455-468.	0.9	36



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109	Binding of a novel dopaminergic agonist radioligand [3H]-fenoldopam1 (SKF 82526) to D-1 receptors in rat striatum. <i>Life Sciences</i> , 1985, 36, 1427-1436.	2.0	35
110	The Positional Influence of the Helical Geometry of the Heteroduplex Substrate on Human RNase H1 Catalysis. <i>Molecular Pharmacology</i> , 2007, 71, 73-82.	1.0	35
111	mRNA levels can be reduced by antisense oligonucleotides via no-go decay pathway. <i>Nucleic Acids Research</i> , 2019, 47, 6900-6916.	6.5	32
112	Golgi-endosome transport mediated by M6PR facilitates release of antisense oligonucleotides from endosomes. <i>Nucleic Acids Research</i> , 2020, 48, 1372-1391.	6.5	32
113	Mitotane (o, p-DDD). <i>Cancer Treatment Reviews</i> , 1980, 7, 49-55.	3.4	31
114	Off-target and a portion of target-specific siRNA mediated mRNA degradation is Ago2 Slicer™ independent and can be mediated by Ago1. <i>Nucleic Acids Research</i> , 2009, 37, 6927-6941.	6.5	31
115	COPII vesicles can affect the activity of antisense oligonucleotides by facilitating the release of oligonucleotides from endocytic pathways. <i>Nucleic Acids Research</i> , 2018, 46, 10225-10245.	6.5	31
116	Delivery of Oligonucleotides and Polynucleotides. <i>Journal of Drug Targeting</i> , 1995, 3, 185-190.	2.1	30
117	Pharmacokinetics in mice of a [3H]-labeled phosphorothioate oligonucleotide formulated in the presence and absence of a cationic lipid. <i>Journal of Controlled Release</i> , 1996, 41, 121-130.	4.8	30
118	Human RNases H. <i>Methods in Enzymology</i> , 2001, 341, 430-440.	0.4	30
119	Proof of Mechanism of Antisense Drugs. <i>Oligonucleotides</i> , 1996, 6, 145-147.	4.4	28
120	Differential effects of aspirin and dexamethasone on phospholipase A2 and C activities and arachidonic acid release from endothelial cells in response to bradykinin and leukotriene D4. <i>Prostaglandins</i> , 1986, 32, 703-708.	1.2	27
121	Kinetic and subcellular analysis of PS-ASO/protein interactions with P54nrb and RNase H1. <i>Nucleic Acids Research</i> , 2019, 47, 10865-10880.	6.5	27
122	Site-specific incorporation of 5-methyl DNA enhances the therapeutic profile of gapmer ASOs. <i>Nucleic Acids Research</i> , 2021, 49, 1828-1839.	6.5	26
123	Depletion of NEAT1 lncRNA attenuates nucleolar stress by releasing sequestered P54nrb and PSF to facilitate c-Myc translation. <i>PLoS ONE</i> , 2017, 12, e0173494.	1.1	26
124	Cellular interactions of auranofin and a related gold complex with raw 264.7 macrophages. <i>Biochemical Pharmacology</i> , 1987, 36, 647-654.	2.0	23
125	Altered expression and transcription of the topoisomerase II gene in nitrogen mustard-resistant human cells. <i>Biochemical Pharmacology</i> , 1988, 37, 4413-4416.	2.0	23
126	Leukotriene C4 ([3H] - LTC4) binding to membranes isolated from a hamster smooth muscle cell line (DDTIMF2). <i>Life Sciences</i> , 1984, 35, 441-448.	2.0	22



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127	Identification of leukotriene D4 specific binding sites in the membrane preparation isolated from guinea pig lung. Prostaglandins, 1984, 28, 805-822.	1.2	22
128	Oligonucleotide therapy. Current Opinion in Biotechnology, 1992, 3, 656-661.	3.3	22
129	Antisense Oligonucleotides Targeting Human Protein Kinase C- $\beta$ Inhibit Phorbol Ester-Induced Reduction of Bradykinin-Evoked Calcium Mobilization in A549 Cells. Molecular Pharmacology, 1997, 51, 209-216.	1.0	22
130	A call to arms against ultra-rare diseases. Nature Biotechnology, 2021, 39, 671-677.	9.4	22
131	Cyclophosphamide, doxorubicin, vincristine, and low-dose continuous infusion bleomycin in non-Hodgkin's lymphoma: Cancer and leukemia group B study #7804. Cancer, 1982, 49, 1346-1352.	2.0	21
132	Effects of coordinated gold compounds on in vitro and in situ DNA replication. Biochemical Pharmacology, 1985, 34, 3243-3250.	2.0	21
133	Inter-strand cross-links and single-strand breaks produced by gold(I) and gold(III) coordination complexes. Biochemical Pharmacology, 1986, 35, 1435-1443.	2.0	21
134	Human RNase H1 Activity Is Regulated by a Unique Redox Switch Formed between Adjacent Cysteines. Journal of Biological Chemistry, 2003, 278, 14906-14912.	1.6	21
135	Cisplatin pharmacokinetics in a patient with renal dysfunction. Medical and Pediatric Oncology, 1978, 5, 183-188.	1.0	20
136	Effect of Antisense Oligonucleotides on Cytokine Release from Human Keratinocytes in an in Vitro Model of Skin. Toxicology and Applied Pharmacology, 1996, 140, 85-93.	1.3	20
137	The mechanism of acute cytotoxicity of triethylphosphine gold(I) complexes. Toxicology and Applied Pharmacology, 1987, 90, 377-390.	1.3	19
138	U1 adaptors result in reduction of multiple pre-mRNA species principally by sequestering U1snRNP. Nucleic Acids Research, 2011, 39, e71-e71.	6.5	19
139	Regulation of agonist and antagonist binding to striatal D-1 dopamine receptors: Studies using the selective D-1 antagonist [3H]SK&F R-83566. Life Sciences, 1986, 38, 2087-2096.	2.0	18
140	Desensitization of vasopressin sensitive adenylate cyclase by vasopressin and phorbol esters. Cellular Signalling, 1990, 2, 153-160.	1.7	18
141	Comparison of bleomycin A2 and talisomycin a specific fragmentation of linear duplex DNA. Biochemical and Biophysical Research Communications, 1979, 91, 871-877.	1.0	17
142	Targeting of Repeated Sequences Unique to a Gene Results in Significant Increases in Antisense Oligonucleotide Potency. PLoS ONE, 2014, 9, e110615.	1.1	17
143	Induction of functional $\beta_2$ -adrenergic receptors in rat aortic smooth muscle cells by sodium butyrate. Biochemical Pharmacology, 1986, 35, 3813-3820.	2.0	16
144	Phorbol 12-myristate 13-acetate inhibition of leukotriene D4-induced signal transduction was rapidly reversed by staurosporine. Biochemical and Biophysical Research Communications, 1988, 157, 521-529.	1.0	16

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145	The effects of prior exposure to bleomycin on the incidence of pulmonary toxicities in a group of patients with disseminated testicular carcinomas. <i>Medical and Pediatric Oncology</i> , 1978, 5, 93-98.	1.0	15
146	Zinostatin (neocarzinostatin). <i>Cancer Treatment Reviews</i> , 1979, 6, 239-249.	3.4	15
147	Evaluating the Mechanism of Action of Antiproliferative Antisense Drugs. <i>Oligonucleotides</i> , 2000, 10, 123-126.	4.4	15
148	Translation can affect the antisense activity of RNase H1-dependent oligonucleotides targeting mRNAs. <i>Nucleic Acids Research</i> , 2018, 46, 293-313.	6.5	15
149	Membrane Destabilization Induced by Lipid Species Increases Activity of Phosphorothioate-Antisense Oligonucleotides. <i>Molecular Therapy - Nucleic Acids</i> , 2018, 13, 686-698.	2.3	15
150	Some ASOs that bind in the coding region of mRNAs and induce RNase H1 cleavage can cause increases in the pre-mRNAs that may blunt total activity. <i>Nucleic Acids Research</i> , 2020, 48, 9840-9858.	6.5	14
151	The transition metal binding properties of a 3RD generation bleomycin analogue, tallysomyin. <i>Biochemical and Biophysical Research Communications</i> , 1978, 85, 1407-1414.	1.0	13
152	Variations in mRNA Content Have No Effect on the Potency of Antisense Oligonucleotides. <i>Oligonucleotides</i> , 2000, 10, 453-461.	4.4	13
153	Site-specific Incorporation of 2'-5'-Linked Nucleic Acids Enhances Therapeutic Profile of Antisense Oligonucleotides. <i>ACS Medicinal Chemistry Letters</i> , 2021, 12, 922-927.	1.3	13
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