

Philip A Cole

List of Publications by Year in descending order

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112
papers

11,923
citations

47006

47
h-index

29157

104
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140
all docs

140
docs citations

140
times ranked

15224
citing authors

#	ARTICLE	IF	CITATIONS
1	Histone Demethylation Mediated by the Nuclear Amine Oxidase Homolog LSD1. <i>Cell</i> , 2004, 119, 941-953.	28.9	3,626
2	Virtual Ligand Screening of the p300/CBP Histone Acetyltransferase: Identification of a Selective Small Molecule Inhibitor. <i>Chemistry and Biology</i> , 2010, 17, 471-482.	6.0	538
3	Discovery of a selective catalytic p300/CBP inhibitor that targets lineage-specific tumours. <i>Nature</i> , 2017, 550, 128-132.	27.8	498
4	Protein Lysine Acetylation by p300/CBP. <i>Chemical Reviews</i> , 2015, 115, 2419-2452.	47.7	398
5	The structural basis of protein acetylation by the p300/CBP transcriptional coactivator. <i>Nature</i> , 2008, 451, 846-850.	27.8	381
6	HATs off. <i>Molecular Cell</i> , 2000, 5, 589-595.	9.7	376
7	Regulation of the p300 HAT domain via a novel activation loop. <i>Nature Structural and Molecular Biology</i> , 2004, 11, 308-315.	8.2	374
8	Time-Resolved Analysis Reveals Rapid Dynamics and Broad Scope of the CBP/p300 Acetylome. <i>Cell</i> , 2018, 174, 231-244.e12.	28.9	313
9	Chemical probes for histone-modifying enzymes. <i>Nature Chemical Biology</i> , 2008, 4, 590-597.	8.0	231
10	Inhibition of the Acetyltransferases p300 and CBP Reveals a Targetable Function for p300 in the Survival and Invasion Pathways of Prostate Cancer Cell Lines. <i>Molecular Cancer Therapeutics</i> , 2011, 10, 1644-1655.	4.1	188
11	Mechanism-based design of a protein kinase inhibitor. <i>Nature Structural Biology</i> , 2001, 8, 37-41.	9.7	185
12	Targeting the CoREST complex with dual histone deacetylase and demethylase inhibitors. <i>Nature Communications</i> , 2018, 9, 53.	12.8	175
13	Inhibition of p300 impairs Foxp3+ T regulatory cell function and promotes antitumor immunity. <i>Nature Medicine</i> , 2013, 19, 1173-1177.	30.7	168
14	How IGF-1 activates its receptor. <i>ELife</i> , 2014, 3, .	6.0	154
15	Structure and chemistry of the p300/CBP and Rtt109 histone acetyltransferases: implications for histone acetyltransferase evolution and function. <i>Current Opinion in Structural Biology</i> , 2008, 18, 741-747.	5.7	152
16	Comparative Analysis of Small Molecules and Histone Substrate Analogues as LSD1 Lysine Demethylase Inhibitors. <i>Journal of the American Chemical Society</i> , 2010, 132, 3164-3176.	13.7	149
17	Regulation of CK2 by phosphorylation and O-GlcNAcylation revealed by semisynthesis. <i>Nature Chemical Biology</i> , 2012, 8, 262-269.	8.0	148
18	Genetically encoded biosensors for visualizing live-cell biochemical activity at super-resolution. <i>Nature Methods</i> , 2017, 14, 427-434.	19.0	138

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19	LSD1 and the chemistry of histone demethylation. <i>Current Opinion in Chemical Biology</i> , 2007, 11, 561-568.	6.1	128
20	Re-programing Chromatin with a Bifunctional LSD1/HDAC Inhibitor Induces Therapeutic Differentiation in DIPG. <i>Cancer Cell</i> , 2019, 36, 528-544.e10.	16.8	128
21	Down-regulation of p300/CBP histone acetyltransferase activates a senescence checkpoint in human melanocytes. <i>Cancer Research</i> , 2002, 62, 6231-9.	0.9	120
22	Chemical Rescue of a Mutant Enzyme in Living Cells. <i>Science</i> , 2006, 311, 1293-1297.	12.6	111
23	Site-Specific Introduction of an Acetyl-Lysine Mimic into Peptides and Proteins by Cysteine Alkylation. <i>Journal of the American Chemical Society</i> , 2010, 132, 9986-9987.	13.7	107
24	Catalytic Mechanisms and Regulation of Protein Kinases. <i>Methods in Enzymology</i> , 2014, 548, 1-21.	1.0	107
25	BCL6 repression of EP300 in human diffuse large B cell lymphoma cells provides a basis for rational combinatorial therapy. <i>Journal of Clinical Investigation</i> , 2010, 120, 4569-4582.	8.2	101
26	p300/CBP-associated Factor Histone Acetyltransferase Processing of a Peptide Substrate. <i>Journal of Biological Chemistry</i> , 2000, 275, 21953-21959.	3.4	100
27	Akt Kinase Activation Mechanisms Revealed Using Protein Semisynthesis. <i>Cell</i> , 2018, 174, 897-907.e14.	28.9	96
28	Synthetic approaches to protein phosphorylation. <i>Current Opinion in Chemical Biology</i> , 2015, 28, 115-122.	6.1	93
29	Phosphorylation-mediated PTEN conformational closure and deactivation revealed with protein semisynthesis. <i>ELife</i> , 2013, 2, e00691.	6.0	89
30	A Selective Phenelzine Analogue Inhibitor of Histone Demethylase LSD1. <i>ACS Chemical Biology</i> , 2014, 9, 1284-1293.	3.4	88
31	Mechanistic Analysis of a Suicide Inactivator of Histone Demethylase LSD1. <i>Biochemistry</i> , 2007, 46, 6892-6902.	2.5	87
32	Selective Inhibition of p300 HAT Blocks Cell Cycle Progression, Induces Cellular Senescence, and Inhibits the DNA Damage Response in Melanoma Cells. <i>Journal of Investigative Dermatology</i> , 2013, 133, 2444-2452.	0.7	87
33	The regulatory enzymes and protein substrates for the lysine ϵ^2 -hydroxybutyrylation pathway. <i>Science Advances</i> , 2021, 7, .	10.3	87
34	Protein tyrosine kinases Src and Csk: a tail's tale. <i>Current Opinion in Chemical Biology</i> , 2003, 7, 580-585.	6.1	84
35	A Tunable Brake for HECT Ubiquitin Ligases. <i>Molecular Cell</i> , 2017, 66, 345-357.e6.	9.7	83
36	Protein semisynthesis and expressed protein ligation: chasing a protein's tail. <i>Current Opinion in Chemical Biology</i> , 2005, 9, 561-569.	6.1	79

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37	The Role of the Phospho-CDK2/Cyclin A Recruitment Site in Substrate Recognition. <i>Journal of Biological Chemistry</i> , 2006, 281, 23167-23179.	3.4	79
38	The Chemical Biology of Reversible Lysine Post-translational Modifications. <i>Cell Chemical Biology</i> , 2020, 27, 953-969.	5.2	76
39	Two Histone/Protein Acetyltransferases, CBP and p300, Are Indispensable for Foxp3 ⁺ T-Regulatory Cell Development and Function. <i>Molecular and Cellular Biology</i> , 2014, 34, 3993-4007.	2.3	75
40	Mechanism of Crosstalk between the LSD1 Demethylase and HDAC1 Deacetylase in the CoREST Complex. <i>Cell Reports</i> , 2020, 30, 2699-2711.e8.	6.4	74
41	Histone deacetylase inhibitors decrease NHEJ both by acetylation of repair factors and trapping of PARP1 at DNA double-strand breaks in chromatin. <i>Leukemia Research</i> , 2016, 45, 14-23.	0.8	65
42	Structural basis of nSH2 regulation and lipid binding in PI3K β . <i>Oncotarget</i> , 2014, 5, 5198-5208.	1.8	62
43	CBP Regulates Recruitment and Release of Promoter-Proximal RNA Polymerase II. <i>Molecular Cell</i> , 2017, 68, 491-503.e5.	9.7	59
44	Endotoxemia-mediated activation of acetyltransferase P300 impairs insulin signaling in obesity. <i>Nature Communications</i> , 2017, 8, 131.	12.8	59
45	p300/CBP-associated Factor Drives DEK into Interchromatin Granule Clusters. <i>Journal of Biological Chemistry</i> , 2005, 280, 31760-31767.	3.4	53
46	Discovery of Spiro Oxazolidinediones as Selective, Orally Bioavailable Inhibitors of p300/CBP Histone Acetyltransferases. <i>ACS Medicinal Chemistry Letters</i> , 2018, 9, 28-33.	2.8	53
47	YcgC represents a new protein deacetylase family in prokaryotes. <i>ELife</i> , 2015, 4, .	6.0	52
48	Combined Targeting of the BRD4 \rightarrow NUT \rightarrow p300 Axis in NUT Midline Carcinoma by Dual Selective Bromodomain Inhibitor, NEO2734. <i>Molecular Cancer Therapeutics</i> , 2020, 19, 1406-1414.	4.1	51
49	Azalsine Analogues as Probes for Protein Lysine Deacetylation and Demethylation. <i>Journal of the American Chemical Society</i> , 2012, 134, 5138-5148.	13.7	49
50	Enzyme-catalyzed expressed protein ligation. <i>Nature Methods</i> , 2016, 13, 925-927.	19.0	49
51	In Vitro Enzymatic Characterization of Near Full Length EGFR in Activated and Inhibited States. <i>Biochemistry</i> , 2009, 48, 6624-6632.	2.5	47
52	Live-Cell Studies of p300/CBP Histone Acetyltransferase Activity and Inhibition. <i>ChemBioChem</i> , 2012, 13, 2113-2121.	2.6	47
53	MITF Expression Predicts Therapeutic Vulnerability to p300 Inhibition in Human Melanoma. <i>Cancer Research</i> , 2019, 79, 2649-2661.	0.9	47
54	Lysine-14 acetylation of histone H3 in chromatin confers resistance to the deacetylase and demethylase activities of an epigenetic silencing complex. <i>ELife</i> , 2018, 7, .	6.0	43

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55	Molecular Features of Phosphatase and Tensin Homolog (PTEN) Regulation by C-terminal Phosphorylation. <i>Journal of Biological Chemistry</i> , 2016, 291, 14160-14169.	3.4	41
56	Modulation of p300/CBP Acetylation of Nucleosomes by Bromodomain Ligand I-CBP112. <i>Biochemistry</i> , 2016, 55, 3727-3734.	2.5	41
57	Inhibiting the coregulator CoREST impairs Foxp3+ Treg function and promotes antitumor immunity. <i>Journal of Clinical Investigation</i> , 2020, 130, 1830-1842.	8.2	41
58	The structural determinants of PH domain-mediated regulation of Akt revealed by segmental labeling. <i>ELife</i> , 2020, 9, .	6.0	41
59	Diverse nucleosome Site-Selectivity among histone deacetylase complexes. <i>ELife</i> , 2020, 9, .	6.0	37
60	Site-Specific Protein Labeling with <i>N</i> -Hydroxysuccinimide-Esters and the Analysis of Ubiquitin Ligase Mechanisms. <i>Journal of the American Chemical Society</i> , 2018, 140, 9374-9378.	13.7	36
61	Enzymatic Analysis of PTEN Ubiquitylation by WWP2 and NEDD4-1 E3 Ligases. <i>Biochemistry</i> , 2016, 55, 3658-3666.	2.5	34
62	AKTivation mechanisms. <i>Current Opinion in Structural Biology</i> , 2019, 59, 47-53.	5.7	34
63	Hydrazide Mimics for Protein Lysine Acylation To Assess Nucleosome Dynamics and Deubiquitinase Action. <i>Journal of the American Chemical Society</i> , 2018, 140, 9478-9485.	13.7	33
64	Investigation of N-Terminal Phospho-Regulation of Uracil DNA Glycosylase Using Protein Semisynthesis. <i>Biophysical Journal</i> , 2017, 113, 393-401.	0.5	30
65	Combination Targeting of the Bromodomain and Acetyltransferase Active Site of p300/CBP. <i>Biochemistry</i> , 2019, 58, 2133-2143.	2.5	30
66	CREB Promotes Beta Cell Gene Expression by Targeting Its Coactivators to Tissue-Specific Enhancers. <i>Molecular and Cellular Biology</i> , 2019, 39, .	2.3	29
67	Lysine-Specific Demethylase 1 Mediates AKT Activity and Promotes Epithelial-to-Mesenchymal Transition in <i>PIK3CA</i> -Mutant Colorectal Cancer. <i>Molecular Cancer Research</i> , 2020, 18, 264-277.	3.4	29
68	Chemical Approaches to Reversible Protein Phosphorylation. <i>Accounts of Chemical Research</i> , 2003, 36, 444-452.	15.6	26
69	Regulation of S-Adenosylhomocysteine Hydrolase by Lysine Acetylation. <i>Journal of Biological Chemistry</i> , 2014, 289, 31361-31372.	3.4	24
70	CBP binding outside of promoters and enhancers in <i>Drosophila melanogaster</i> . <i>Epigenetics and Chromatin</i> , 2015, 8, 48.	3.9	24
71	Histone H2B Deacetylation Selectivity: Exploring Chromatin's Dark Matter with an Engineered Sortase. <i>Journal of the American Chemical Society</i> , 2022, 144, 3360-3364.	13.7	24
72	Comparative analysis of the catalytic regulation of NEDD4-1 and WWP2 ubiquitin ligases. <i>Journal of Biological Chemistry</i> , 2019, 294, 17421-17436.	3.4	23

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73	Measurement of nanoscale DNA translocation by uracil DNA glycosylase in human cells. <i>Nucleic Acids Research</i> , 2017, 45, 12413-12424.	14.5	21
74	The structural basis of PTEN regulation by multi-site phosphorylation. <i>Nature Structural and Molecular Biology</i> , 2021, 28, 858-868.	8.2	20
75	Analysis of p300/CBP Histone Acetyltransferase Regulation Using Circular Permutation and Semisynthesis. <i>Journal of the American Chemical Society</i> , 2010, 132, 1222-1223.	13.7	19
76	Regulation of Myf5 Early Enhancer by Histone Acetyltransferase P300 during Stem Cell Differentiation. <i>Molecular Biology (Los Angeles, Calif)</i> , 2012, 01, .	0.0	19
77	Disordered N-Terminal Domain of Human Uracil DNA Glycosylase (hUNG2) Enhances DNA Translocation. <i>ACS Chemical Biology</i> , 2017, 12, 2260-2263.	3.4	18
78	Mechanistic analysis of ghrelin-O-acyltransferase using substrate analogs. <i>Bioorganic Chemistry</i> , 2015, 62, 64-73.	4.1	17
79	Interaction with the DNA Repair Protein Thymine DNA Glycosylase Regulates Histone Acetylation by p300. <i>Biochemistry</i> , 2016, 55, 6766-6775.	2.5	17
80	Analysis of Site-Specific Phosphorylation of PTEN by Using Enzyme-Catalyzed Expressed Protein Ligation. <i>ChemBioChem</i> , 2020, 21, 64-68.	2.6	17
81	An Fc Domain Protein-Small Molecule Conjugate as an Enhanced Immunomodulator. <i>Journal of the American Chemical Society</i> , 2014, 136, 3370-3373.	13.7	14
82	Allosteric regulation of epigenetic modifying enzymes. <i>Current Opinion in Chemical Biology</i> , 2017, 39, 109-115.	6.1	14
83	Getting the Most Out of Your Crystals: Data Collection at the New High-Flux, Microfocus MX Beamlines at NSLS-II. <i>Molecules</i> , 2019, 24, 496.	3.8	13
84	The protein kinase Akt acts as a coat adaptor in endocytic recycling. <i>Nature Cell Biology</i> , 2020, 22, 927-933.	10.3	13
85	Selective protein N-terminal labeling with N-hydroxysuccinimide esters. <i>Methods in Enzymology</i> , 2020, 639, 333-353.	1.0	12
86	N-Terminal Protein Labeling with N-Hydroxysuccinimide Esters and Microscale Thermophoresis Measurements of Protein-Protein Interactions Using Labeled Protein. <i>Current Protocols</i> , 2021, 1, e14.	2.9	12
87	Chemical Screen Identifies Diverse and Novel Histone Deacetylase Inhibitors as Repressors of NUT Function: Implications for NUT Carcinoma Pathogenesis and Treatment. <i>Molecular Cancer Research</i> , 2021, 19, 1818-1830.	3.4	12
88	Investigation into the use of histone deacetylase inhibitor MS-275 as a topical agent for the prevention and treatment of cutaneous squamous cell carcinoma in an SKH-1 hairless mouse model. <i>PLoS ONE</i> , 2019, 14, e0213095.	2.5	10
89	HDAC2 targeting stabilizes the CoREST complex in renal tubular cells and protects against renal ischemia/reperfusion injury. <i>Scientific Reports</i> , 2021, 11, 9018.	3.3	10
90	Complementary Roles of GCN5 and PCAF in Foxp3+ T-Regulatory Cells. <i>Cancers</i> , 2019, 11, 554.	3.7	9

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91	Discovery of spirohydantoins as selective, orally bioavailable inhibitors of p300/CBP histone acetyltransferases. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2021, 39, 127854.	2.2	9
92	Methods and Applications of Expressed Protein Ligation. <i>Methods in Molecular Biology</i> , 2020, 2133, 1-13.	0.9	8
93	Site-Specific 5-Formyl Cytosine Mediated DNA-Histone Cross-Links: Synthesis and Polymerase Bypass by Human DNA Polymerase β . <i>Angewandte Chemie - International Edition</i> , 2021, 60, 26489-26494.	13.8	7
94	Multifaceted Regulation of Akt by Diverse C-Terminal Post-translational Modifications. <i>ACS Chemical Biology</i> , 2022, 17, 68-76.	3.4	7
95	Ubiquitin Ligase Activities of WWP1 Germline Variants K740N and N745S. <i>Biochemistry</i> , 2021, 60, 357-364.	2.5	6
96	Enzymatic analysis of WWP2 E3 ubiquitin ligase using protein microarrays identifies autophagy-related substrates. <i>Journal of Biological Chemistry</i> , 2022, 298, 101854.	3.4	6
97	Protein Chemical Approaches to Understanding PTEN Lipid Phosphatase Regulation. <i>Methods in Enzymology</i> , 2018, 607, 405-422.	1.0	5
98	Distinct biochemical properties of the class I histone deacetylase complexes. <i>Current Opinion in Chemical Biology</i> , 2022, 70, 102179.	6.1	5
99	Analysis of Cellular Tyrosine Phosphorylation via Chemical Rescue of Conditionally Active Abl Kinase. <i>Biochemistry</i> , 2018, 57, 1390-1398.	2.5	4
100	Site-Specific 5-Formyl Cytosine Mediated DNA-Histone Cross-Links: Synthesis and Polymerase Bypass by Human DNA Polymerase β . <i>Angewandte Chemie</i> , 2021, 133, 26693-26698.	2.0	3
101	Open questions: two challenges in chemical biology - chemical engineering and the science of diet. <i>BMC Biology</i> , 2013, 11, 87.	3.8	1
102	An Fc-Small Molecule Conjugate for Targeted Inhibition of the Adenosine 2A Receptor. <i>ChemBioChem</i> , 2016, 17, 1951-1960.	2.6	1
103	Connectivity Mapping of BCL6 Targeted Therapy Guides Rational Design of Potent and Specific Non-Chemotherapy Combinatorial Regimens in DLBCL. <i>Blood</i> , 2007, 110, 523-523.	1.4	1
104	Switching immune signals on and off. <i>ELife</i> , 2015, 4, .	6.0	1
105	GENE-22. RE-PROGRAMING CHROMATIN WITH A BIFUNCTIONAL LSD1/HDAC INHIBITOR INDUCES THERAPEUTIC DIFFERENTIATION IN DIPG. <i>Neuro-Oncology</i> , 2018, 20, vi107-vi108.	1.2	0
106	Editorial overview: Biological catalysis at the cross-roads of signaling and metabolism. <i>Current Opinion in Structural Biology</i> , 2019, 59, iii-v.	5.7	0
107	Histone Acetyltransferase Activity of p300 Is Required for Transcriptional Repression by the Promyelocytic Leukemia Zinc Finger Protein. <i>Blood</i> , 2004, 104, 359-359.	1.4	0
108	Structure and chemistry of the human p300/CBP and yeast Rtt109 histone acetyltransferase. <i>FASEB Journal</i> , 2009, 23, 89.2.	0.5	0

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109	Tackling Targets in Epigenetics. FASEB Journal, 2012, 26, 230.3.	0.5	0
110	The Epigenetic Regulators CBP and p300 Facilitate Leukemogenesis and Represent Therapeutic Targets In Acute Myeloid Leukemia (AML). Blood, 2013, 122, 3732-3732.	1.4	0
111	The Interplay of Phosphorylation and Ubiquitylation in the Regulation of PTEN. FASEB Journal, 2015, 29, 570.2.	0.5	0
112	Targeting Reversible Lysine Modifications. FASEB Journal, 2015, 29, 107.3.	0.5	0