

# Antonello Mai

## List of Publications by Year in descending order

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351  
papers

15,744  
citations

16411

64  
h-index

31759

101  
g-index

375  
all docs

375  
docs citations

375  
times ranked

22616  
citing authors

#	ARTICLE	IF	CITATIONS
1	The Nitrobenzoxadiazole Derivative NBDHEX Behaves as Plasmodium falciparum Gametocyte Selective Inhibitor with Malaria Parasite Transmission Blocking Activity. <i>Pharmaceuticals</i> , 2022, 15, 168.	1.7	3
2	A potent HDAC inhibitor blocks Toxoplasma gondii tachyzoite growth and profoundly disrupts parasite gene expression. <i>International Journal of Antimicrobial Agents</i> , 2022, 59, 106526.	1.1	12
3	Transcriptomic and genomic studies classify NKL54 as a histone deacetylase inhibitor with indirect influence on MEF2-dependent transcription. <i>Nucleic Acids Research</i> , 2022, 50, 2566-2586.	6.5	12
4	Targeting the anti-apoptotic Bcl-2 family proteins: machine learning virtual screening and biological evaluation of new small molecules. <i>Theranostics</i> , 2022, 12, 2427-2444.	4.6	12
5	Cytoplasmic HDAC4 regulates the membrane repair mechanism in Duchenne muscular dystrophy. <i>Journal of Cachexia, Sarcopenia and Muscle</i> , 2022, 13, 1339-1359.	2.9	11
6	Inhibition of PKC $\delta$ Improves Dystrophic Heart Phenotype and Function in a Novel Model of DMD Cardiomyopathy. <i>International Journal of Molecular Sciences</i> , 2022, 23, 2256.	1.8	1
7	Heterocycle-containing tranilcypromine derivatives endowed with high anti-LSD1 activity. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2022, 37, 973-985.	2.5	2
8	Biochemical Functions and Clinical Characterizations of the Sirtuins in Diabetes-Induced Retinal Pathologies. <i>International Journal of Molecular Sciences</i> , 2022, 23, 4048.	1.8	11
9	Determinants of epigenetic resistance to HDAC inhibitors in dystrophic fibroâ€œadipogenic progenitors. <i>EMBO Reports</i> , 2022, 23, e54721.	2.0	7
10	Novel non-covalent LSD1 inhibitors endowed with anticancer effects in leukemia and solid tumor cellular models. <i>European Journal of Medicinal Chemistry</i> , 2022, 237, 114410.	2.6	15
11	Sirtuin modulators: past, present, and future perspectives. <i>Future Medicinal Chemistry</i> , 2022, 14, 915-939.	1.1	24
12	Novel Targeting of DNA Methyltransferase Activity Inhibits Ewing Sarcoma Cell Proliferation and Enhances Tumor Cell Sensitivity to DNA Damaging Drugs by Activating the DNA Damage Response. <i>Frontiers in Endocrinology</i> , 2022, 13, .	1.5	4
13	Effects of Structurally Different HDAC Inhibitors against <i>Trypanosoma cruzi</i> , <i>Leishmania</i> , and <i>Schistosoma mansoni</i> . <i>ACS Infectious Diseases</i> , 2022, 8, 1356-1366.	1.8	13
14	Therapeutic Potential and Activity Modulation of the Protein Lysine Deacylase Sirtuin 5. <i>Journal of Medicinal Chemistry</i> , 2022, 65, 9580-9606.	2.9	21
15	Downregulation of miRâ€œ326 and its host gene <i>Arrestin1</i> induces proâ€œsurvival activity of E2F1 and promotes medulloblastoma growth. <i>Molecular Oncology</i> , 2021, 15, 523-542.	2.1	8
16	Novel Pyridineâ€œBased Hydroxamates and 2â€œAminoanilides as Histone Deacetylase Inhibitors: Biochemical Profile and Anticancer Activity. <i>ChemMedChem</i> , 2021, 16, 989-999.	1.6	8
17	Sirtuins. , 2021, , 1-15.		0
18	Amphetamine Modulation of Long-Term Object Recognition Memory in Rats: Influence of Stress. <i>Frontiers in Pharmacology</i> , 2021, 12, 644521.	1.6	1

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19	Regulatory Interplay between miR-181a-5p and Estrogen Receptor Signaling Cascade in Breast Cancer. <i>Cancers</i> , 2021, 13, 543.	1.7	10
20	The Two-Faced Role of SIRT6 in Cancer. <i>Cancers</i> , 2021, 13, 1156.	1.7	33
21	SIRT5 Inhibition Induces Brown Fat-Like Phenotype in 3T3-L1 Preadipocytes. <i>Cells</i> , 2021, 10, 1126.	1.8	16
22	Emerging Therapeutic Potential of SIRT6 Modulators. <i>Journal of Medicinal Chemistry</i> , 2021, 64, 9732-9758.	2.9	38
23	Metabolic Rewiring by Loss of Sirt5 Promotes Kras-Induced Pancreatic Cancer Progression. <i>Gastroenterology</i> , 2021, 161, 1584-1600.	0.6	50
24	Histone deacetylase 8 drives the immune response and the growth of glioma. <i>Glia</i> , 2021, 69, 2682-2698.	2.5	14
25	Polycomb Repressive Complex 2 Modulation through the Development of EZH2-EED Interaction Inhibitors and EED Binders. <i>Journal of Medicinal Chemistry</i> , 2021, 64, 11774-11797.	2.9	25
26	Anti-influenza A virus activity and structure-activity relationship of a series of nitrobenzoxadiazole derivatives. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2021, 36, 2128-2138.	2.5	5
27	Mass spectrometry enables the discovery of inhibitors of an LPS transport assembly via disruption of protein-protein interactions. <i>Chemical Communications</i> , 2021, 57, 10747-10750.	2.2	6
28	First-in-Class Inhibitors of the Ribosomal Oxygenase MINA53. <i>Journal of Medicinal Chemistry</i> , 2021, 64, 17031-17050.	2.9	7
29	Sirtuins. , 2021, , 1422-1435.		0
30	Sex-dependent Effects of the Drugs of Abuse Amphetamine and the Smart Drug 3,4-Methylenedioxypyrovalerone on Fear Memory Generalization in Rats. <i>Neuroscience</i> , 2021, , .	1.1	2
31	Epigenetic polypharmacology: A new frontier for epigenetic drug discovery. <i>Medicinal Research Reviews</i> , 2020, 40, 190-244.	5.0	74
32	Inhibition of class I HDACs imprints adipogenesis toward oxidative and brown-like phenotype. <i>Biochimica Et Biophysica Acta - Molecular and Cell Biology of Lipids</i> , 2020, 1865, 158594.	1.2	11
33	Recent advances in epigenetic proteolysis targeting chimeras (Epi-PROTACs). <i>European Journal of Medicinal Chemistry</i> , 2020, 207, 112750.	2.6	12
34	From PARP1 to TNKS2 Inhibition: A Structure-Based Approach. <i>ACS Medicinal Chemistry Letters</i> , 2020, 11, 862-868.	1.3	8
35	Properly Substituted Cyclic Bis-(2-bromobenzylidene) Compounds Behaved as Dual p300/CARM1 Inhibitors and Induced Apoptosis in Cancer Cells. <i>Molecules</i> , 2020, 25, 3122.	1.7	4
36	Lysine Acetyltransferase Inhibitors From Natural Sources. <i>Frontiers in Pharmacology</i> , 2020, 11, 1243.	1.6	22

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37	CDK9 as a Valuable Target in Cancer: From Natural Compounds Inhibitors to Current Treatment in Pediatric Soft Tissue Sarcomas. <i>Frontiers in Pharmacology</i> , 2020, 11, 1230.	1.6	20
38	Diphenylene Iodonium Is a Noncovalent MAO Inhibitor: A Biochemical and Structural Analysis. <i>ChemMedChem</i> , 2020, 15, 1394-1397.	1.6	4
39	Detrimental effects of the $\alpha$ -bath salt™ methylenedioxyprovalerone on social play behavior in male rats. <i>Neuropsychopharmacology</i> , 2020, 45, 2012-2019.	2.8	5
40	Identification of Inhibitors to Trypanosoma cruzi Sirtuins Based on Compounds Developed to Human Enzymes. <i>International Journal of Molecular Sciences</i> , 2020, 21, 3659.	1.8	8
41	The Pan-Sirtuin Inhibitor MC2494 Regulates Mitochondrial Function in a Leukemia Cell Line. <i>Frontiers in Oncology</i> , 2020, 10, 820.	1.3	8
42	Design of First-in-Class Dual EZH2/HDAC Inhibitor: Biochemical Activity and Biological Evaluation in Cancer Cells. <i>ACS Medicinal Chemistry Letters</i> , 2020, 11, 977-983.	1.3	49
43	Sirtuin modulators: where are we now? A review of patents from 2015 to 2019. <i>Expert Opinion on Therapeutic Patents</i> , 2020, 30, 389-407.	2.4	39
44	The Innovative Potential of Statins in Cancer: New Targets for New Therapies. <i>Frontiers in Chemistry</i> , 2020, 8, 516.	1.8	73
45	Pharmacological inhibition of lysine-specific demethylase 1 (LSD1) induces global transcriptional deregulation and ultrastructural alterations that impair viability in Schistosoma mansoni. <i>PLoS Neglected Tropical Diseases</i> , 2020, 14, e0008332.	1.3	11
46	Targeting histone acetylation/deacetylation in parasites: an update (2017–2020). <i>Current Opinion in Chemical Biology</i> , 2020, 57, 65-74.	2.8	35
47	Novel Quinoline Compounds Active in Cancer Cells through Coupled DNA Methyltransferase Inhibition and Degradation. <i>Cancers</i> , 2020, 12, 447.	1.7	8
48	A closer look into NADPH oxidase inhibitors: Validation and insight into their mechanism of action. <i>Redox Biology</i> , 2020, 32, 101466.	3.9	62
49	Tranylcypromine-Based LSD1 Inhibitors: Structure-Activity Relationships, Antiproliferative Effects in Leukemia, and Gene Target Modulation. <i>ChemMedChem</i> , 2020, 15, 643-658.	1.6	18
50	Altered mitochondrial function in cells carrying a premutation or unmethylated full mutation of the FMR1 gene. <i>Human Genetics</i> , 2020, 139, 227-245.	1.8	16
51	Sirt4: A Multifaceted Enzyme at the Crossroads of Mitochondrial Metabolism and Cancer. <i>Frontiers in Oncology</i> , 2020, 10, 474.	1.3	49
52	Targeting the scaffolding role of LSD1 (KDM1A) poises acute myeloid leukemia cells for retinoic acid-induced differentiation. <i>Science Advances</i> , 2020, 6, eaax2746.	4.7	56
53	Discovery of the First Human Arylsulfatase A Reversible Inhibitor Impairing Mouse Oocyte Fertilization. <i>ACS Chemical Biology</i> , 2020, 15, 1349-1357.	1.6	4
54	HAT inhibitors in cancer therapy. , 2020, , 51-80.		2

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55	Selective class II HDAC inhibitors impair myogenesis by modulating the stability and activity of HDAC-MEF2 complexes. <i>EMBO Reports</i> , 2020, 21, e51028.	2.0	1
56	Title is missing!. , 2020, 14, e0008332.		0
57	Title is missing!. , 2020, 14, e0008332.		0
58	Title is missing!. , 2020, 14, e0008332.		0
59	Title is missing!. , 2020, 14, e0008332.		0
60	H19-Dependent Transcriptional Regulation of $\beta$ 3 and $\beta$ 4 Integrins Upon Estrogen and Hypoxia Favors Metastatic Potential in Prostate Cancer. <i>International Journal of Molecular Sciences</i> , 2019, 20, 4012.	1.8	22
61	The Protein Arginine Methyltransferases 1 and 5 affect Myc properties in glioblastoma stem cells. <i>Scientific Reports</i> , 2019, 9, 15925.	1.6	35
62	Structure-Reactivity Relationships on Substrates and Inhibitors of the Lysine Deacetylase Sirtuin 2 from <i>Schistosoma mansoni</i> ( <i>Sirt2</i> ). <i>Journal of Medicinal Chemistry</i> , 2019, 62, 8733-8759.	2.9	18
63	Histone deacetylase inhibitors exert anti-tumor effects on human adherent and stem-like glioma cells. <i>Clinical Epigenetics</i> , 2019, 11, 11.	1.8	55
64	Epigenetic Pharmacology in Regenerative Medicine (Epi-Drugs). , 2019, , 405-444.		0
65	Histone Deacetylases Contribute to Excitotoxicity-Triggered Degeneration of Retinal Ganglion Cells In Vivo. <i>Molecular Neurobiology</i> , 2019, 56, 8018-8034.	1.9	20
66	Identification of a novel quinoline-based DNA demethylating compound highly potent in cancer cells. <i>Clinical Epigenetics</i> , 2019, 11, 68.	1.8	30
67	Histone deacetylases as an epigenetic pillar for the development of hybrid inhibitors in cancer. <i>Current Opinion in Chemical Biology</i> , 2019, 50, 89-100.	2.8	23
68	Comparison of the effects of synthetic and plant-derived mTOR regulators on healthy human ovarian cells. <i>European Journal of Pharmacology</i> , 2019, 854, 70-78.	1.7	3
69	Statins and Histone Deacetylase Inhibitors Affect Lamin A/C - Histone Deacetylase 2 Interaction in Human Cells. <i>Frontiers in Cell and Developmental Biology</i> , 2019, 7, 6.	1.8	23
70	The emerging role of epigenetics in human autoimmune disorders. <i>Clinical Epigenetics</i> , 2019, 11, 34.	1.8	200
71	$\beta$ -Diketocarboxylic Acids and Their Esters Act as Carbonic Anhydrase IX and XII Selective Inhibitors. <i>ACS Medicinal Chemistry Letters</i> , 2019, 10, 661-665.	1.3	18
72	Enzymatic and Biological Characterization of Novel Sirtuin Modulators against Cancer. <i>International Journal of Molecular Sciences</i> , 2019, 20, 5654.	1.8	16

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73	Inhibition of Histone Demethylases LSD1 and UTX Regulates ER $\alpha$ Signaling in Breast Cancer. <i>Cancers</i> , 2019, 11, 2027.	1.7	34
74	Amphetamine and the Smart Drug 3,4-Methylenedioxypropylamphetamine (MDPV) Induce Generalization of Fear Memory in Rats. <i>Frontiers in Molecular Neuroscience</i> , 2019, 12, 292.	1.4	9
75	Development of alkyl glycerone phosphate synthase inhibitors: Structure-activity relationship and effects on ether lipids and epithelial-mesenchymal transition in cancer cells. <i>European Journal of Medicinal Chemistry</i> , 2019, 163, 722-735.	2.6	15
76	Intergenerational inheritance of high fat diet-induced cardiac lipotoxicity in <i>Drosophila</i> . <i>Nature Communications</i> , 2019, 10, 193.	5.8	49
77	Acute and chronic neurobehavioral effects of the designer drug and bath salt constituent 3,4-methylenedioxypropylamphetamine in the rat. <i>Journal of Psychopharmacology</i> , 2019, 33, 392-405.	2.0	21
78	Identification of novel quinazoline derivatives as potent antiplasmodial agents. <i>European Journal of Medicinal Chemistry</i> , 2019, 161, 277-291.	2.6	44
79	Effect of $\beta$ -Methoxy Substitution on the Anti-HIV Activity of Dihydropyrimidin-4(3 <i>H</i> )-ones. <i>Journal of Medicinal Chemistry</i> , 2019, 62, 604-621.	2.9	14
80	Epigenetic Metalloenzymes. <i>Current Medicinal Chemistry</i> , 2019, 26, 2748-2785.	1.2	12
81	DNA Methylation: Biological Implications and Modulation of Its Aberrant Dysregulation. <i>RNA Technologies</i> , 2019, , 295-331.	0.2	1
82	Lysine acetyltransferase inhibitors: structure-activity relationships and potential therapeutic implications. <i>Future Medicinal Chemistry</i> , 2018, 10, 1067-1091.	1.1	24
83	Pyrazole-based inhibitors of enhancer of zeste homologue 2 induce apoptosis and autophagy in cancer cells. <i>Philosophical Transactions of the Royal Society B: Biological Sciences</i> , 2018, 373, 20170150.	1.8	13
84	Disruptor of telomeric silencing 1-like (DOT1L): disclosing a new class of non-nucleoside inhibitors by means of ligand-based and structure-based approaches. <i>Journal of Computer-Aided Molecular Design</i> , 2018, 32, 435-458.	1.3	15
85	Targeting the CoREST complex with dual histone deacetylase and demethylase inhibitors. <i>Nature Communications</i> , 2018, 9, 53.	5.8	175
86	KAT3B-p300 and H3AcK18/H3AcK14 levels are prognostic markers for kidney ccRCC tumor aggressiveness and target of KAT inhibitor CPTH2. <i>Clinical Epigenetics</i> , 2018, 10, 44.	1.8	12
87	RIP1-HAT1-SIRT Complex Identification and Targeting in Treatment and Prevention of Cancer. <i>Clinical Cancer Research</i> , 2018, 24, 2886-2900.	3.2	40
88	Effects of Class II-Selective Histone Deacetylase Inhibitor on Neuromuscular Function and Disease Progression in SOD1-ALS Mice. <i>Neuroscience</i> , 2018, 379, 228-238.	1.1	22
89	Multi-omics profiling reveals a distinctive epigenome signature for high-risk acute promyelocytic leukemia. <i>Oncotarget</i> , 2018, 9, 25647-25660.	0.8	13
90	EZH2, HIF-1, and Their Inhibitors: An Overview on Pediatric Cancers. <i>Frontiers in Pediatrics</i> , 2018, 6, 328.	0.9	14

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91	Application of Small Epigenetic Modulators in Pediatric Medulloblastoma. <i>Frontiers in Pediatrics</i> , 2018, 6, 370.	0.9	12
92	Pharmacological activation of SIRT6 triggers lethal autophagy in human cancer cells. <i>Cell Death and Disease</i> , 2018, 9, 996.	2.7	75
93	Six Years (2012–2018) of Researches on Catalytic EZH2 Inhibitors: The Boom of the 2-Pyridone Compounds. <i>Chemical Record</i> , 2018, 18, 1818-1832.	2.9	76
94	Trends of LSD1 inhibitors in viral infections. <i>Future Medicinal Chemistry</i> , 2018, 10, 1133-1136.	1.1	10
95	HDAC1 inhibition by MS-275 in mesothelial cells limits cellular invasion and promotes MMT reversal. <i>Scientific Reports</i> , 2018, 8, 8492.	1.6	23
96	A Quinoline-Based DNA Methyltransferase Inhibitor as a Possible Adjuvant in Osteosarcoma Therapy. <i>Molecular Cancer Therapeutics</i> , 2018, 17, 1881-1892.	1.9	33
97	Altered modulation of lamin A/C–HDAC2 interaction and p21 expression during oxidative stress response in HGPS. <i>Aging Cell</i> , 2018, 17, e12824.	3.0	39
98	Nucleocytoplasmic export of HDAC5 and SIRT2 downregulation: two epigenetic mechanisms by which antidepressants enhance synaptic plasticity markers. <i>Psychopharmacology</i> , 2018, 235, 2831-2846.	1.5	11
99	Combined HAT/EZH2 modulation leads to cancer-selective cell death. <i>Oncotarget</i> , 2018, 9, 25630-25646.	0.8	5
100	A new nitrobenzoxadiazole-based GSTP1-1 inhibitor with a previously unheard of mechanism of action and high stability. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2017, 32, 240-247.	2.5	17
101	Selective targeting of HDAC1/2 elicits anticancer effects through Gli1 acetylation in preclinical models of SHH Medulloblastoma. <i>Scientific Reports</i> , 2017, 7, 44079.	1.6	57
102	Metabolite profiling of ascidian <i>Styela plicata</i> using LC–MS with multivariate statistical analysis and their antitumor activity. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2017, 32, 614-623.	2.5	17
103	Lysine Deacetylase Inhibitors in Parasites: Past, Present, and Future Perspectives. <i>Journal of Medicinal Chemistry</i> , 2017, 60, 4780-4804.	2.9	71
104	Development of 1,2,4-Oxadiazoles as Potent and Selective Inhibitors of the Human Deacetylase Sirtuin 2: Structure–Activity Relationship, X-ray Crystal Structure, and Anticancer Activity. <i>Journal of Medicinal Chemistry</i> , 2017, 60, 2344-2360.	2.9	82
105	Novel coumarin- and quinolinone-based polycycles as cell division cycle 25-A and -C phosphatases inhibitors induce proliferation arrest and apoptosis in cancer cells. <i>European Journal of Medicinal Chemistry</i> , 2017, 134, 316-333.	2.6	24
106	EZH2 inhibitors: a patent review (2014-2016). <i>Expert Opinion on Therapeutic Patents</i> , 2017, 27, 797-813.	2.4	40
107	The relevance of K <sub>i</sub> calculation for bi-substrate enzymes illustrated by kinetic evaluation of a novel lysine (K) acetyltransferase 8 inhibitor. <i>European Journal of Medicinal Chemistry</i> , 2017, 136, 480-486.	2.6	7
108	Structure–Activity Relationships on Cinnamoyl Derivatives as Inhibitors of p300 Histone Acetyltransferase. <i>ChemMedChem</i> , 2017, 12, 1359-1368.	1.6	11

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109	Towards the development of activity-based probes for detection of lysine-specific demethylase-1 activity. <i>Bioorganic and Medicinal Chemistry</i> , 2017, 25, 847-856.	1.4	11
110	Structural Basis of Sirtuin 6 Activation by Synthetic Small Molecules. <i>Angewandte Chemie - International Edition</i> , 2017, 56, 1007-1011.	7.2	125
111	Structural Basis of Sirtuin 6 Activation by Synthetic Small Molecules. <i>Angewandte Chemie</i> , 2017, 129, 1027-1031.	1.6	4
112	Effect of Class II HDAC inhibition on glutamate transporter expression and survival in SOD1-ALS mice. <i>Neuroscience Letters</i> , 2017, 656, 120-125.	1.0	22
113	Crystal structures of the mitochondrial deacylase Sirtuin 4 reveal isoform-specific acyl recognition and regulation features. <i>Nature Communications</i> , 2017, 8, 1513.	5.8	80
114	Epi-drugs in combination with immunotherapy: a new avenue to improve anticancer efficacy. <i>Clinical Epigenetics</i> , 2017, 9, 59.	1.8	118
115	Attenuation of diet-induced obesity and induction of white fat browning with a chemical inhibitor of histone deacetylases. <i>International Journal of Obesity</i> , 2017, 41, 289-298.	1.6	41
116	Essential Oil Extraction, Chemical Analysis and Anti-Candida Activity of <i>Calamintha nepeta</i> (L.) Savi subsp. <i>glandulosa</i> (Req.) Ball. New Approaches. <i>Molecules</i> , 2017, 22, 203.	1.7	30
117	The histone methyltransferase EZH2 as a druggable target in SHH medulloblastoma cancer stem cells. <i>Oncotarget</i> , 2017, 8, 68557-68570.	0.8	49
118	LSD1 inhibitors: a patent review (2010-2015). <i>Expert Opinion on Therapeutic Patents</i> , 2016, 26, 565-580.	2.4	46
119	Sirtuin functions and modulation: from chemistry to the clinic. <i>Clinical Epigenetics</i> , 2016, 8, 61.	1.8	291
120	Chemical epigenetics to assess the role of HDAC1 <sup>Δ3</sup> inhibition in macrophage pro-inflammatory gene expression. <i>MedChemComm</i> , 2016, 7, 2184-2190.	3.5	7
121	The emerging role of lysine methyltransferase SETD8 in human diseases. <i>Clinical Epigenetics</i> , 2016, 8, 102.	1.8	77
122	Progress in the Development of Lysine Methyltransferase SETD8 Inhibitors. <i>ChemMedChem</i> , 2016, 11, 1680-1685.	1.6	18
123	MC1568 inhibits HDAC6/8 activity and influenza A virus replication in lung epithelial cells: role of Hsp90 acetylation. <i>Future Medicinal Chemistry</i> , 2016, 8, 2017-2031.	1.1	33
124	The International Human Epigenome Consortium: A Blueprint for Scientific Collaboration and Discovery. <i>Cell</i> , 2016, 167, 1145-1149.	13.5	404
125	Polymyxins and quinazolines are LSD1/KDM1A inhibitors with unusual structural features. <i>Science Advances</i> , 2016, 2, e1601017.	4.7	61
126	Histone deacetylase inhibitors restore IL-10 expression in lipopolysaccharide-induced cell inflammation and reduce IL-1 $\beta$ and IL-6 production in breast silicone implant in C57BL/6J wild-type murine model. <i>Autoimmunity</i> , 2016, 49, 155-165.	1.2	20



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127	The histone acetyltransferase p300 inhibitor C646 reduces pro-inflammatory gene expression and inhibits histone deacetylases. <i>Biochemical Pharmacology</i> , 2016, 102, 130-140.	2.0	46
128	1,4-Dihydropyridines Active on the SIRT1/AMPK Pathway Ameliorate Skin Repair and Mitochondrial Function and Exhibit Inhibition of Proliferation in Cancer Cells. <i>Journal of Medicinal Chemistry</i> , 2016, 59, 1471-1491.	2.9	60
129	Discovery of a Novel Inhibitor of Histone Lysine-Specific Demethylase 1A (KDM1A/LSD1) as Orally Active Antitumor Agent. <i>Journal of Medicinal Chemistry</i> , 2016, 59, 1501-1517.	2.9	70
130	Histone acetyltransferase inhibitor CPTH6 preferentially targets lung cancer stem-like cells. <i>Oncotarget</i> , 2016, 7, 11332-11348.	0.8	49
131	Tumour-specific metabolic adaptation to acidosis is coupled to epigenetic stability in osteosarcoma cells. <i>American Journal of Cancer Research</i> , 2016, 6, 859-75.	1.4	18
132	Targeting Lysine Deacetylases (KDACs) in Parasites. <i>PLoS Neglected Tropical Diseases</i> , 2015, 9, e0004026.	1.3	47
133	Pure enantiomers of benzoylamino-tranylcypromine: LSD1 inhibition, gene modulation in human leukemia cells and effects on clonogenic potential of murine promyelocytic blasts. <i>European Journal of Medicinal Chemistry</i> , 2015, 94, 163-174.	2.6	28
134	Enzyme kinetics and inhibition of histone acetyltransferase KAT8. <i>European Journal of Medicinal Chemistry</i> , 2015, 105, 289-296.	2.6	31
135	Sirtuin modulators: an updated patent review (2012 – 2014). <i>Expert Opinion on Therapeutic Patents</i> , 2015, 25, 5-15.	2.4	46
136	Sirtuin function in aging heart and vessels. <i>Journal of Molecular and Cellular Cardiology</i> , 2015, 83, 55-61.	0.9	83
137	Pyrrole- and indole-containing tranylcypromine derivatives as novel lysine-specific demethylase 1 inhibitors active on cancer cells. <i>MedChemComm</i> , 2015, 6, 665-670.	3.5	24
138	Interplay among nucleosomal DNA, histone tails, and corepressor CoREST underlies LSD1-mediated H3 demethylation. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2015, 112, 2752-2757.	3.3	71
139	SIRT5 regulation of ammonia-induced autophagy and mitophagy. <i>Autophagy</i> , 2015, 11, 253-270.	4.3	223
140	New Insights on the Mechanism of Quinoline-based DNA Methyltransferase Inhibitors. <i>Journal of Biological Chemistry</i> , 2015, 290, 6293-6302.	1.6	50
141	Novel Histone Deacetylase Inhibitors Induce Growth Arrest, Apoptosis, and Differentiation in Sarcoma Cancer Stem Cells. <i>Journal of Medicinal Chemistry</i> , 2015, 58, 4073-4079.	2.9	55
142	Emerging approaches for histone deacetylase inhibitor drug discovery. <i>Expert Opinion on Drug Discovery</i> , 2015, 10, 599-613.	2.5	63
143	A new water soluble MAPK activator exerts antitumor activity in melanoma cells resistant to the BRAF inhibitor vemurafenib. <i>Biochemical Pharmacology</i> , 2015, 95, 16-27.	2.0	29
144	A Novel Cell-Permeable, Selective, and Noncompetitive Inhibitor of KAT3 Histone Acetyltransferases from a Combined Molecular Pruning/Classical Isosterism Approach. <i>Journal of Medicinal Chemistry</i> , 2015, 58, 2779-2798.	2.9	48

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145	Chronic stress and antidepressant induced changes in Hdac5 and Sirt2 affect synaptic plasticity. <i>European Neuropsychopharmacology</i> , 2015, 25, 2036-2048.	0.3	51
146	Discovery of Inhibitors for the Ether Lipid-Generating Enzyme AGPS as Anti-Cancer Agents. <i>ACS Chemical Biology</i> , 2015, 10, 2589-2597.	1.6	54
147	Synthesis and structure-activity relationship of new cytotoxic agents targeting human glutathione-S-transferases. <i>European Journal of Medicinal Chemistry</i> , 2015, 89, 156-171.	2.6	32
148	Identification of Structural Features of $\alpha,\beta$ -Dicarbonyl Derivatives that Induce Inhibition and/or Activation of Histone Acetyltransferases KAT3B/p300 and KAT2B/PCAF. <i>ChemMedChem</i> , 2015, 10, 144-157.	1.6	21
149	Pure Diastereomers of a Tranylcyproline-Based LSD1 Inhibitor: Enzyme Selectivity and In-Cell Studies. <i>ACS Medicinal Chemistry Letters</i> , 2015, 6, 173-177.	1.3	16
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