

Ina Oehme

List of Publications by Year in descending order

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Version: 2024-02-01

45
papers

7,681
citations

331670

21
h-index

377865

34
g-index

47
all docs

47
docs citations

47
times ranked

17601
citing authors

#	ARTICLE	IF	CITATIONS
1	Guidelines for the use and interpretation of assays for monitoring autophagy (3rd edition). <i>Autophagy</i> , 2016, 12, 1-222.	9.1	4,701
2	HDAC family: What are the cancer relevant targets?. <i>Cancer Letters</i> , 2009, 277, 8-21.	7.2	893
3	Histone Deacetylase 8 in Neuroblastoma Tumorigenesis. <i>Clinical Cancer Research</i> , 2009, 15, 91-99.	7.0	335
4	DNMT and HDAC inhibitors induce cryptic transcription start sites encoded in long terminal repeats. <i>Nature Genetics</i> , 2017, 49, 1052-1060.	21.4	235
5	HDAC8: a multifaceted target for therapeutic interventions. <i>Trends in Pharmacological Sciences</i> , 2015, 36, 481-492.	8.7	210
6	HDAC5 and HDAC9 in Medulloblastoma: Novel Markers for Risk Stratification and Role in Tumor Cell Growth. <i>Clinical Cancer Research</i> , 2010, 16, 3240-3252.	7.0	175
7	Histone deacetylase 10 promotes autophagy-mediated cell survival. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2013, 110, E2592-601.	7.1	168
8	Synthesis and Biological Investigation of Oxazole Hydroxamates as Highly Selective Histone Deacetylase 6 (HDAC6) Inhibitors. <i>Journal of Medicinal Chemistry</i> , 2016, 59, 1545-1555.	6.4	90
9	Targeting histone deacetylase 8 as a therapeutic approach to cancer and neurodegenerative diseases. <i>Future Medicinal Chemistry</i> , 2016, 8, 1609-1634.	2.3	79
10	HDAC Family Members Intertwined in the Regulation of Autophagy: A Druggable Vulnerability in Aggressive Tumor Entities. <i>Cells</i> , 2015, 4, 135-168.	4.1	71
11	Targeting class I histone deacetylase 2 in MYC amplified group 3 medulloblastoma. <i>Acta Neuropathologica Communications</i> , 2015, 3, 22.	5.2	66
12	Targeting of HDAC8 and investigational inhibitors in neuroblastoma. <i>Expert Opinion on Investigational Drugs</i> , 2009, 18, 1605-1617.	4.1	64
13	Structure-Based Design and Biological Characterization of Selective Histone Deacetylase 8 (HDAC8) Inhibitors with Anti-Neuroblastoma Activity. <i>Journal of Medicinal Chemistry</i> , 2017, 60, 10188-10204.	6.4	56
14	Selective Inhibition of Histone Deacetylase 10: Hydrogen Bonding to the Gatekeeper Residue is Implicated. <i>Journal of Medicinal Chemistry</i> , 2019, 62, 4426-4443.	6.4	56
15	<i>GRHL1</i> Acts as Tumor Suppressor in Neuroblastoma and Is Negatively Regulated by MYCN and HDAC3. <i>Cancer Research</i> , 2014, 74, 2604-2616.	0.9	54
16	Three-dimensional tumor cell growth stimulates autophagic flux and recapitulates chemotherapy resistance. <i>Cell Death and Disease</i> , 2017, 8, e3013-e3013.	6.3	43
17	Establishment and application of a novel patient-derived KIAA1549:BRAF-driven pediatric pilocytic astrocytoma model for preclinical drug testing. <i>Oncotarget</i> , 2017, 8, 11460-11479.	1.8	43
18	The enzyme activity of histone deacetylase 8 is modulated by a redox-switch. <i>Redox Biology</i> , 2019, 20, 60-67.	9.0	37

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19	Inhibition of Histone Deacetylases Permits Lipopolysaccharide-Mediated Secretion of Bioactive IL-1 ^β via a Caspase-1-Independent Mechanism. <i>Journal of Immunology</i> , 2015, 195, 5421-5431.	0.8	36
20	Dual role of HDAC10 in lysosomal exocytosis and DNA repair promotes neuroblastoma chemoresistance. <i>Scientific Reports</i> , 2018, 8, 10039.	3.3	36
21	The HDAC6/8/10 inhibitor TH34 induces DNA damage-mediated cell death in human high-grade neuroblastoma cell lines. <i>Archives of Toxicology</i> , 2018, 92, 2649-2664.	4.2	28
22	Histone deacetylase 10-promoted autophagy as a druggable point of interference to improve the treatment response of advanced neuroblastomas. <i>Autophagy</i> , 2013, 9, 2163-2165.	9.1	22
23	A kinome-wide RNAi screen identifies ALK as a target to sensitize neuroblastoma cells for HDAC8-inhibitor treatment. <i>Cell Death and Differentiation</i> , 2018, 25, 2053-2070.	11.2	22
24	Design and Synthesis of Dihydroxamic Acids as HDAC6/8/10 Inhibitors. <i>ChemMedChem</i> , 2020, 15, 1163-1174.	3.2	21
25	Rapid In Vivo Validation of HDAC Inhibitor-Based Treatments in Neuroblastoma Zebrafish Xenografts. <i>Pharmaceuticals</i> , 2020, 13, 345.	3.8	19
26	Synthesis and structure activity relationship of 1, 3-benzo-thiazine-2-thiones as selective HDAC8 inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2019, 184, 111756.	5.5	17
27	Broad-Spectrum HDAC Inhibitors Promote Autophagy through FOXO Transcription Factors in Neuroblastoma. <i>Cells</i> , 2021, 10, 1001.	4.1	17
28	Identification of histone deacetylase 10 (HDAC10) inhibitors that modulate autophagy in transformed cells. <i>European Journal of Medicinal Chemistry</i> , 2022, 234, 114272.	5.5	15
29	Design, Synthesis and Biological Characterization of Histone Deacetylase 8 (HDAC8) Proteolysis Targeting Chimeras (PROTACs) with Anti-Neuroblastoma Activity. <i>International Journal of Molecular Sciences</i> , 2022, 23, 7535.	4.1	15
30	Pediatric Targeted Therapy: Clinical Feasibility of Personalized Diagnostics in Children with Relapsed and Progressive Tumors. <i>Brain Pathology</i> , 2016, 26, 506-516.	4.1	14
31	Functional Therapeutic Target Validation Using Pediatric Zebrafish Xenograft Models. <i>Cancers</i> , 2022, 14, 849.	3.7	13
32	iTReX: Interactive exploration of mono- and combination therapy dose response profiling data. <i>Pharmacological Research</i> , 2022, 175, 105996.	7.1	11
33	First Fluorescent Acetylspermidine Deacetylation Assay for HDAC10 Identifies Selective Inhibitors with Cellular Target Engagement**. <i>ChemBioChem</i> , 2022, 23, .	2.6	9
34	Combining APR-246 and HDAC-Inhibitors: A Novel Targeted Treatment Option for Neuroblastoma. <i>Cancers</i> , 2021, 13, 4476.	3.7	8
35	MODL-15. High-throughput combination drug screening identifies synergism between retinoic acid treatment and BCL-XL-inhibition in MYC(N)-driven medulloblastoma and neuroblastoma models. <i>Neuro-Oncology</i> , 2022, 24, i171-i172.	1.2	1
36	EMBR-01. CLASS I HDAC INHIBITORS AND PLK1 INHIBITORS SYNERGIZE IN MYC-AMPLIFIED MEDULLOBLASTOMA. <i>Neuro-Oncology</i> , 2021, 23, i5-i5.	1.2	0

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37	EMBR-11. SYNERGISTIC DRUG COMBINATIONS FOR THE TREATMENT OF MYC AMPLIFIED GROUP 3 MEDULLOBLASTOMA. <i>Neuro-Oncology</i> , 2021, 23, i7-i8.	1.2	0
38	TMOD-04. IMAGE-BASED DRUG RESPONSE PROFILING FROM PEDIATRIC TUMOR CELL SPHEROIDS USING PATIENT-BY-PATIENT DEEP TRANSFER LEARNING. <i>Neuro-Oncology</i> , 2021, 23, i36-i36.	1.2	0
39	LGG-11. BH3-MIMETICS TARGETING BCL-XL SELECTIVELY IMPACT THE SENESCENT COMPARTMENT OF PILOCYTIC ASTROCYTOMA. <i>Neuro-Oncology</i> , 2021, 23, i33-i34.	1.2	0
40	MBRS-19. SYNERGISM OF HDAC AND PARP INHIBITORS IN MYC-DRIVEN GROUP 3 MEDULLOBLASTOMA CELLS. <i>Neuro-Oncology</i> , 2020, 22, iii401-iii401.	1.2	0
41	MODL-04. Drug screening in Disorders with Abnormal DNA Damage Response/Repair (DADDR) and <i>in vivo</i> validation. <i>Neuro-Oncology</i> , 2022, 24, i168-i169.	1.2	0
42	LGG-18. Inhibition of Bcl-xL targets the senescent compartment of pilocytic astrocytoma. <i>Neuro-Oncology</i> , 2022, 24, i91-i92.	1.2	0
43	LGG-25. The first-in-class ERK inhibitor ulixertinib (BVD-523) shows activity in MAPK-driven pediatric low-grade glioma models as single agent and in combination with MEK inhibitors or senolytics. <i>Neuro-Oncology</i> , 2022, 24, i93-i93.	1.2	0
44	DDEL-01. The role of key pharmacodynamic and pharmacokinetic parameters in drug response prediction of pediatric tumors in the precision oncology study INFORM. <i>Neuro-Oncology</i> , 2022, 24, i33-i34.	1.2	0
45	Multiomics analysis of pediatric solid tumors within the INFORM precision oncology study: From functional drug profiling to biomarker identification.. <i>Journal of Clinical Oncology</i> , 2022, 40, 10036-10036.	1.6	0