

Jolanta Grembecka

List of Publications by Year in descending order

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48
papers

2,475
citations

270111

25
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252626

46
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all docs

49
docs citations

49
times ranked

3458
citing authors

#	ARTICLE	IF	CITATIONS
1	EWS-FLI1 and Menin Converge to Regulate ATF4 Activity in Ewing Sarcoma. <i>Molecular Cancer Research</i> , 2021, 19, 1182-1195.	1.5	6
2	Discovery of first-in-class inhibitors of ASH1L histone methyltransferase with anti-leukemic activity. <i>Nature Communications</i> , 2021, 12, 2792.	5.8	17
3	Genome-scale CRISPR-Cas9 screen of Wnt/ β -catenin signaling identifies therapeutic targets for colorectal cancer. <i>Science Advances</i> , 2021, 7, .	4.7	28
4	Small-molecule inhibitors targeting Polycomb repressive complex 1 RING domain. <i>Nature Chemical Biology</i> , 2021, 17, 784-793.	3.9	31
5	Development of potent dimeric inhibitors of GAS41 YEATS domain. <i>Cell Chemical Biology</i> , 2021, 28, 1716-1727.e6.	2.5	10
6	Combinatorial treatment with menin and FLT3 inhibitors induces complete remission in AML models with activating FLT3 mutations. <i>Blood</i> , 2020, 136, 2958-2963.	0.6	20
7	Covalent and noncovalent constraints yield a figure eight-like conformation of a peptide inhibiting the menin-MLL interaction. <i>European Journal of Medicinal Chemistry</i> , 2020, 207, 112748.	2.6	4
8	Covalent inhibition of NSD1 histone methyltransferase. <i>Nature Chemical Biology</i> , 2020, 16, 1403-1410.	3.9	52
9	Targeting epigenetic protein-protein interactions with small-molecule inhibitors. <i>Future Medicinal Chemistry</i> , 2020, 12, 1305-1326.	1.1	12
10	Menin inhibitor MI-3454 induces remission in MLL1-rearranged and NPM1-mutated models of leukemia. <i>Journal of Clinical Investigation</i> , 2020, 130, 981-997.	3.9	146
11	CEBPA-mutated leukemia is sensitive to genetic and pharmacological targeting of the MLL1 complex. <i>Leukemia</i> , 2019, 33, 1608-1619.	3.3	19
12	Combined MAPK Pathway and HDAC Inhibition Breaks Melanoma. <i>Cancer Discovery</i> , 2019, 9, 469-471.	7.7	27
13	Distinct pathways affected by menin versus MLL1/MLL2 in MLL-rearranged acute myeloid leukemia. <i>Experimental Hematology</i> , 2019, 69, 37-42.	0.2	13
14	Menin regulates the serine biosynthetic pathway in Ewing sarcoma. <i>Journal of Pathology</i> , 2018, 245, 324-336.	2.1	35
15	A tool compound targeting the core binding factor Runt domain to disrupt binding to CBF β in leukemic cells. <i>Leukemia and Lymphoma</i> , 2018, 59, 2188-2200.	0.6	11
16	Pharmacologic Inhibition of the Menin-MLL Interaction Leads to Transcriptional Repression of PEG10 and Blocks Hepatocellular Carcinoma. <i>Molecular Cancer Therapeutics</i> , 2018, 17, 26-38.	1.9	40
17	Validation of approximate nonempirical scoring model for menin-mixed lineage leukemia inhibitors. <i>Theoretical Chemistry Accounts</i> , 2018, 137, 1.	0.5	2
18	GAS41 Recognizes Diacetylated Histone H3 through a Bivalent Binding Mode. <i>ACS Chemical Biology</i> , 2018, 13, 2739-2746.	1.6	29

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19	Complexity of Blocking Bivalent Protein-Protein Interactions: Development of a Highly Potent Inhibitor of the Menin-Mixed-Lineage Leukemia Interaction. <i>Journal of Medicinal Chemistry</i> , 2018, 61, 4832-4850.	2.9	45
20	Stabilizing the Mixed Lineage Leukemia Protein. <i>New England Journal of Medicine</i> , 2017, 376, 1688-1689.	13.9	3
21	Theoretical models of inhibitory activity for inhibitors of protein-protein interactions: targeting menin-mixed lineage leukemia with small molecules. <i>MedChemComm</i> , 2017, 8, 2216-2227.	3.5	7
22	Gastrin Induces Nuclear Export and Proteasome Degradation of Menin in Enteric Glial Cells. <i>Gastroenterology</i> , 2017, 153, 1555-1567.e15.	0.6	28
23	Tumorigenicity of Ewing sarcoma is critically dependent on the trithorax proteins MLL1 and menin. <i>Oncotarget</i> , 2017, 8, 458-471.	0.8	29
24	H3K36 methyltransferases as cancer drug targets: rationale and perspectives for inhibitor development. <i>Future Medicinal Chemistry</i> , 2016, 8, 1589-1607.	1.1	37
25	BMI1 regulates PRC1 architecture and activity through homo- and hetero-oligomerization. <i>Nature Communications</i> , 2016, 7, 13343.	5.8	52
26	Small Molecule Inhibitor of CBF β -RUNX Binding for RUNX Transcription Factor Driven Cancers. <i>EBioMedicine</i> , 2016, 8, 117-131.	2.7	84
27	Property Focused Structure-Based Optimization of Small Molecule Inhibitors of the Protein-Protein Interaction between Menin and Mixed Lineage Leukemia (MLL). <i>Journal of Medicinal Chemistry</i> , 2016, 59, 892-913.	2.9	56
28	Progress towards small molecule menin-mixed lineage leukemia (MLL) interaction inhibitors with in vivo utility. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2015, 25, 2720-2725.	1.0	10
29	A small-molecule inhibitor of the aberrant transcription factor CBF β -SMMHC delays leukemia in mice. <i>Science</i> , 2015, 347, 779-784.	6.0	104
30	Epigenetic regulation of IL-12-dependent T cell proliferation. <i>Journal of Leukocyte Biology</i> , 2015, 98, 601-613.	1.5	35
31	Targeting the MLL complex in castration-resistant prostate cancer. <i>Nature Medicine</i> , 2015, 21, 344-352.	15.2	165
32	Pharmacologic Inhibition of the Menin-MLL Interaction Blocks Progression of MLL Leukemia In Vivo. <i>Cancer Cell</i> , 2015, 27, 589-602.	7.7	290
33	Rational Design of Orthogonal Multipolar Interactions with Fluorine in Protein-Ligand Complexes. <i>Journal of Medicinal Chemistry</i> , 2015, 58, 7465-7474.	2.9	70
34	Two Loops Undergoing Concerted Dynamics Regulate the Activity of the ASH1L Histone Methyltransferase. <i>Biochemistry</i> , 2015, 54, 5401-5413.	1.2	18
35	Targeting protein-protein interactions in hematologic malignancies: still a challenge or a great opportunity for future therapies?. <i>Immunological Reviews</i> , 2015, 263, 279-301.	2.8	42
36	Ash1l controls quiescence and self-renewal potential in hematopoietic stem cells. <i>Journal of Clinical Investigation</i> , 2015, 125, 2007-2020.	3.9	57

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37	Trithorax group genes in hematopoiesis. <i>Oncotarget</i> , 2015, 6, 17855-17856.	0.8	0
38	Challenges and opportunities in targeting the menin-MLL interaction. <i>Future Medicinal Chemistry</i> , 2014, 6, 447-462.	1.1	63
39	High-Affinity Small-Molecule Inhibitors of the Menin-Mixed Lineage Leukemia (MLL) Interaction Closely Mimic a Natural Protein-Protein Interaction. <i>Journal of Medicinal Chemistry</i> , 2014, 57, 1543-1556.	2.9	79
40	The same site on the integrase-binding domain of lens epithelium-derived growth factor is a therapeutic target for MLL leukemia and HIV. <i>Blood</i> , 2014, 124, 3730-3737.	0.6	30
41	Structural insights into inhibition of the bivalent menin-MLL interaction by small molecules in leukemia. <i>Blood</i> , 2012, 120, 4461-4469.	0.6	160
42	Menin-MLL inhibitors reverse oncogenic activity of MLL fusion proteins in leukemia. <i>Nature Chemical Biology</i> , 2012, 8, 277-284.	3.9	349
43	A Small Molecule Inhibitor of the CBF β -SMMHC/RUNX Interaction Attenuates Inv(16) Leukemia in Vivo. <i>Blood</i> , 2012, 120, 286-286.	0.6	1
44	Crystal Structure of Menin Reveals Binding Site for Mixed Lineage Leukemia (MLL) Protein. <i>Journal of Biological Chemistry</i> , 2011, 286, 31742-31748.	1.6	83
45	Targeting Menin-MLL Interaction to Inhibit MLL Fusion Oncoproteins in Leukemia. <i>Blood</i> , 2011, 118, 2497-2497.	0.6	1
46	Targeting LEDGF Interactions in MLL Leukemia. <i>Blood</i> , 2011, 118, 2500-2500.	0.6	0
47	Molecular Basis of the Mixed Lineage Leukemia-Menin Interaction. <i>Journal of Biological Chemistry</i> , 2010, 285, 40690-40698.	1.6	73
48	Molecular Basis of Menin-MLL Interaction: Implication for Targeted Therapies in MLL Leukemias. <i>Blood</i> , 2009, 114, 3775-3775.	0.6	2