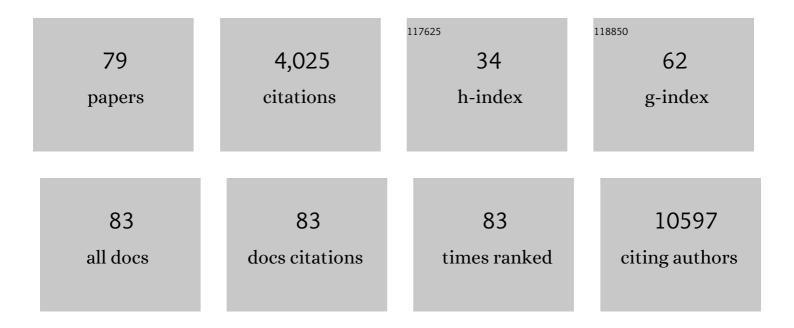
## Alexandre Puissant

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Targeting MYCN in Neuroblastoma by BET Bromodomain Inhibition. Cancer Discovery, 2013, 3, 308-323.	9.4	549
2	Resveratrol Promotes Autophagic Cell Death in Chronic Myelogenous Leukemia Cells via JNK-Mediated p62/SQSTM1 Expression and AMPK Activation. Cancer Research, 2010, 70, 1042-1052.	0.9	335
3	Metformin inhibits melanoma development through autophagy and apoptosis mechanisms. Cell Death and Disease, 2011, 2, e199-e199.	6.3	250
4	InÂVivo RNAi Screening Identifies a Leukemia-Specific Dependence on Integrin Beta 3 Signaling. Cancer Cell, 2013, 24, 45-58.	16.8	144
5	When autophagy meets cancer through p62/SQSTM1. American Journal of Cancer Research, 2012, 2, 397-413.	1.4	139
6	The dual mTORC1 and mTORC2 inhibitor AZD8055 has anti-tumor activity in acute myeloid leukemia. Leukemia, 2012, 26, 1195-1202.	7.2	138
7	Sestrin2 integrates Akt and mTOR signaling to protect cells against energetic stress-induced death. Cell Death and Differentiation, 2013, 20, 611-619.	11.2	137
8	SYK Is a Critical Regulator of FLT3 in Acute Myeloid Leukemia. Cancer Cell, 2014, 25, 226-242.	16.8	126
9	Autophagy, a key mechanism of oncogenesis and resistance in leukemia. Blood, 2017, 129, 547-552.	1.4	121
10	Targeting MTHFD2 in acute myeloid leukemia. Journal of Experimental Medicine, 2016, 213, 1285-1306.	8.5	118
11	Aberrant activation of the PI3K/mTOR pathway promotes resistance to sorafenib in AML. Oncogene, 2016, 35, 5119-5131.	5.9	96
12	Autophagy is an important event for megakaryocytic differentiation of the chronic myelogenous leukemia K562 cell line. Autophagy, 2009, 5, 1092-1098.	9.1	92
13	The creatine kinase pathway is a metabolic vulnerability in EVI1-positive acute myeloid leukemia. Nature Medicine, 2017, 23, 301-313.	30.7	79
14	Acadesine Kills Chronic Myelogenous Leukemia (CML) Cells through PKC-Dependent Induction of Autophagic Cell Death. PLoS ONE, 2009, 4, e7889.	2.5	79
15	BCL2L10 is a predictive factor for resistance to Azacitidine in MDS and AML patients. Oncotarget, 2012, 3, 490-501.	1.8	75
16	Gene expression profiling of imatinib and PD166326-resistant CML cell lines identifies Fyn as a gene associated with resistance to BCR-ABL inhibitors. Molecular Cancer Therapeutics, 2009, 8, 1924-1933.	4.1	71
17	Targeting autophagy to fight hematopoietic malignancies. Cell Cycle, 2010, 9, 3470-3478.	2.6	70
18	Exploiting an Asp-Glu "switch―in glycogen synthase kinase 3 to design paralog-selective inhibitors for use in acute myeloid leukemia. Science Translational Medicine, 2018, 10, .	12.4	69

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19	AMPK- and p62/SQSTM1-dependent autophagy mediate Resveratrol-induced cell death in chronic myelogenous leukemia. Autophagy, 2010, 6, 655-657.	9.1	63
20	SPARC functions as an anti-stress factor by inactivating p53 through Akt-mediated MDM2 phosphorylation to promote melanoma cell survival. Oncogene, 2011, 30, 4887-4900.	5.9	60
21	Imatinib mesylateâ€resistant human chronic myelogenous leukemia cell lines exhibit high sensitivity to the phytoalexin resveratrol. FASEB Journal, 2008, 22, 1894-1904.	0.5	59
22	Persistent Activation of the Fyn/ERK Kinase Signaling Axis Mediates Imatinib Resistance in Chronic Myelogenous Leukemia Cells through Upregulation of Intracellular SPARC. Cancer Research, 2010, 70, 9659-9670.	0.9	56
23	Ultrasound-assisted one-pot synthesis of anti-CML nucleosides featuring 1,2,3-triazole nucleobase under iron-copper catalysis. Ultrasonics Sonochemistry, 2012, 19, 1132-1138.	8.2	56
24	Tumor suppressor function of miR-483-3p on squamous cell carcinomas due to its pro-apoptotic properties. Cell Cycle, 2013, 12, 2183-2193.	2.6	52
25	In Vitro and In Vivo Anti-Melanoma Effects of Ciglitazone. Journal of Investigative Dermatology, 2009, 129, 1208-1218.	0.7	51
26	Using antagonistic pleiotropy to design a chemotherapy-induced evolutionary trap to target drug resistance in cancer. Nature Genetics, 2020, 52, 408-417.	21.4	47
27	CDK6 is an essential direct target of NUP98 fusion proteins in acute myeloid leukemia. Blood, 2020, 136, 387-400.	1.4	46
28	Apoptosis and erythroid differentiation triggered by Bcr-Abl inhibitors in CML cell lines are fully distinguishable processes that exhibit different sensitivity to caspase inhibition. Oncogene, 2007, 26, 2445-2458.	5.9	45
29	The anti-apoptotic Bcl-B protein inhibits BECN1-dependent autophagic cell death. Autophagy, 2012, 8, 637-649.	9.1	45
30	SYK regulates mTOR signaling in AML. Leukemia, 2013, 27, 2118-2128.	7.2	44
31	Cathepsin B release after imatinib-mediated lysosomal membrane permeabilization triggers BCR–ABL cleavage and elimination of chronic myelogenous leukemia cells. Leukemia, 2010, 24, 115-124.	7.2	43
32	The small heat shock protein B8 (HSPB8) confers resistance to bortezomib by promoting autophagic removal of misfolded proteins in multiple myeloma cells. Oncotarget, 2014, 5, 6252-6266.	1.8	43
33	Azacitidine-resistant SKM1 myeloid cells are defective for AZA-induced mitochondrial apoptosis and autophagy. Cell Cycle, 2011, 10, 2339-2343.	2.6	37
34	The p53/p21 <sup>Cip1/ Waf1</sup> pathway mediates the effects of SPARC on melanoma cell cycle progression. Pigment Cell and Melanoma Research, 2011, 24, 219-232.	3.3	36
35	Imatinib triggers mesenchymal-like conversion of CML cells associated with increased aggressiveness. Journal of Molecular Cell Biology, 2012, 4, 207-220.	3.3	32
36	Ciglitazone negatively regulates CXCL1 signaling through MITF to suppress melanoma growth. Cell Death and Differentiation, 2011, 18, 109-121.	11.2	31

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37	All tyrosine kinase inhibitor-resistant chronic myelogenous cells are highly sensitive to Ponatinib. Oncotarget, 2012, 3, 1557-1565.	1.8	30
38	Targeting acute myeloid leukemia dependency on VCP-mediated DNA repair through a selective second-generation small-molecule inhibitor. Science Translational Medicine, 2021, 13, .	12.4	29
39	Dual Role of Sp3 Transcription Factor as an Inducer of Apoptosis and a Marker of Tumour Aggressiveness. PLoS ONE, 2009, 4, e4478.	2.5	29
40	Mechanism of action of the multikinase inhibitor Foretinib. Cell Cycle, 2011, 10, 4138-4148.	2.6	28
41	BCL-B (BCL2L10) is overexpressed in patients suffering from multiple myeloma (MM) and drives an MM-like disease in transgenic mice. Journal of Experimental Medicine, 2016, 213, 1705-1722.	8.5	24
42	Cystine uptake inhibition potentiates front-line therapies in acute myeloid leukemia. Leukemia, 2022, 36, 1585-1595.	7.2	24
43	Inhibition of imatinib-mediated apoptosis by the caspase-cleaved form of the tyrosine kinase Lyn in chronic myelogenous leukemia cells. Leukemia, 2009, 23, 1500-1506.	7.2	23
44	Targeting serine hydroxymethyltransferases 1 and 2 for T-cell acute lymphoblastic leukemia therapy. Leukemia, 2022, 36, 348-360.	7.2	23
45	Drug Resistance in Hematological Malignancies. International Journal of Molecular Sciences, 2020, 21, 6091.	4.1	21
46	Increased SYK activity is associated with unfavorable outcome among patients with acute myeloid leukemia. Oncotarget, 2015, 6, 25575-25587.	1.8	20
47	The caspase 6 derived N-terminal fragment of DJ-1 promotes apoptosis via increased ROS production. Cell Death and Differentiation, 2012, 19, 1769-1778.	11.2	19
48	Characterization of midostaurin as a dual inhibitor of FLT3 and SYK and potentiation of FLT3 inhibition against FLT3-ITD-driven leukemia harboring activated SYK kinase. Oncotarget, 2017, 8, 52026-52044.	1.8	19
49	Tumor Lysis Syndrome and AKI: Beyond Crystal Mechanisms. Journal of the American Society of Nephrology: JASN, 2022, 33, 1154-1171.	6.1	18
50	Targeting lysosomes to eradicate imatinib-resistant chronic myelogenous leukemia cells. Leukemia, 2010, 24, 1099-1101.	7.2	17
51	MUC1-C oncoprotein promotes FLT3 receptor activation in acute myeloid leukemia cells. Blood, 2014, 123, 734-742.	1.4	16
52	Matched Targeted Therapy for Pediatric Patients with Relapsed, Refractory, or High-Risk Leukemias: A Report from the LEAP Consortium. Cancer Discovery, 2021, 11, 1424-1439.	9.4	16
53	Structure elucidation of the new citharoxazole from the Mediterranean deepâ€sea sponge <i>Latrunculia (Biannulata) citharistae</i> . Magnetic Resonance in Chemistry, 2011, 49, 533-536.	1.9	13
54	The Folate Cycle Enzyme MTHFR Is a Critical Regulator of Cell Response to MYC-Targeting Therapies. Cancer Discovery, 2020, 10, 1894-1911.	9.4	13

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55	Creatine kinase pathway inhibition alters GSK3 and WNT signaling in EVI1-positive AML. Leukemia, 2019, 33, 800-804.	7.2	10
56	BET inhibitors impair leukemic stem cell function only in defined oncogenic subgroups of acute myeloid leukaemias. Leukemia Research, 2019, 87, 106269.	0.8	9
57	Endothelial cells: major players in acute myeloid leukaemia. Blood Reviews, 2022, 54, 100932.	5.7	9
58	Screening of ETO2-GLIS2–induced Super Enhancers identifies targetable cooperative dependencies in acute megakaryoblastic leukemia. Science Advances, 2022, 8, eabg9455.	10.3	9
59	P2RY2-AKT activation is a therapeutically actionable consequence of XPO1 inhibition in acute myeloid leukemia. Nature Cancer, 2022, 3, 837-851.	13.2	9
60	HER2 Signaling Hijacks the Creatine Shuttle to Fuel Breast Cancer Cell Growth. Cell Metabolism, 2018, 28, 805-807.	16.2	8
61	Granulomonocytic progenitors are key target cells of azacytidine in higher risk myelodysplastic syndromes and acute myeloid leukemia. Leukemia, 2018, 32, 1856-1860.	7.2	7
62	A multiparametric niche-like drug screening platform in acute myeloid leukemia. Blood Cancer Journal, 2022, 12, .	6.2	6
63	A new posttranslational regulation of REDD1/DDIT4 through cleavage by caspase 3 modifies its cellular function. Cell Death and Disease, 2014, 5, e1349-e1349.	6.3	5
64	Niche-like Ex Vivo High Throughput (NEXT) Drug Screening Platform in Acute Myeloid Leukemia. Blood, 2020, 136, 12-13.	1.4	4
65	BCL2L10 (Bcl-B) Is Associated with Resistance to Azacitidine (AZA) in MDS and AML, and Is a Possible Therapeutic Target in AZA Resistant Patients. Blood, 2012, 120, 701-701.	1.4	2
66	Targeting MTHFD2 in Acute Myeloid Leukemia. Blood, 2015, 126, 443-443.	1.4	2
67	The Folate Cycle Enzyme MTHFR Is a Critical Regulator of Cell Response to MYC-Targeting Therapies. Blood, 2019, 134, 877-877.	1.4	1
68	Induction of Autophagic Cell Death Circumvents Azacitidine-Resistance In Myelodysplastic Syndrome-Derived Cell Lines. Blood, 2010, 116, 1817-1817.	1.4	1
69	Targeting Folate Metabolism In Acute Myelogenous Leukemia. Blood, 2013, 122, 3798-3798.	1.4	1
70	Trials in Progress: A Phase I Study to Evaluate the Safety and Pharmacokinetic Profiles of CB-5339 in Participants with Relapsed/Refractory Acute Myeloid Leukemia or Relapsed/Refractory Intermediate or High-Risk Myelodysplastic Syndrome. Blood, 2020, 136, 21-21.	1.4	1
71	Autophagy and blood diseases. Hematologie, 2015, 21, 107-116.	0.0	0
72	In Vivo RNA Interference Screening Identifies a Leukemia-Specific Dependence on Integrin Beta 3 Signaling. Blood, 2011, 118, 758-758.	1.4	0

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73	Involvement of autophagy in cellular development and differentiation. Hematologie, 2015, 21, 212-220.	0.0	Ο
74	Aberrant Activation of the PI3K/mTOR Pathway Promotes Resistance to Sorafenib in AML. Blood, 2015, 126, 2472-2472.	1.4	0
75	Identification of CKMT1B As a New Target in EVI1-Positive AML. Blood, 2015, 126, 3674-3674.	1.4	Ο
76	Implication of the Anti-Apoptotic Protein Bcl-B (BCL2L10) in the Pathogenesis of Multiple Myeloma. Blood, 2015, 126, 2958-2958.	1.4	0
77	Targeting MTHFD2 in acute myeloid leukemia. Journal of Cell Biology, 2016, 214, 2141OIA135.	5.2	Ο
78	Targeting the Creatine Kinase Pathway in EVI1-Positive Acute Myeloid Leukemia. Blood, 2016, 128, 523-523.	1.4	0
79	Vcp-Regulated Homologous Recombination Represents a New Druggable Vulnerability in Acute Myeloid Leukemia. Blood, 2017, 130, 880-880.	1.4	0