Frederic Lepretre

List of Publications by Year
in descending order

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30 papers	2,443 citations	331670 21 h-index	477307 29 g-index
31 all docs	31 docs citations	31 times ranked	4039 citing authors

#	Article	IF	Citations
1	Detection of residual and chemoresistant leukemic cells in an immune-competent mouse model of acute myeloid leukemia: Potential for unravelling their interactions with immunity. PLoS ONE, 2022, 17, e0267508.	2.5	0
2	The stem cell-associated gene expression signature allows risk stratification in pediatric acute myeloid leukemia. Leukemia, 2019, 33, 348-357.	7.2	44
3	Functional Analysis of Somatic Mutations Affecting Receptor Tyrosine Kinase Family in Metastatic Colorectal Cancer. Molecular Cancer Therapeutics, 2019, 18, 1137-1148.	4.1	0
4	Optimization of Routine Testing for MET Exon 14 Splice Site Mutations in NSCLC Patients. Journal of Thoracic Oncology, 2018, 13, 1873-1883.	1.1	30
5	Isolation and characterization of two canine melanoma cell lines: new models for comparative oncology. BMC Cancer, 2018, 18, 1219.	2.6	11
6	BAP1 Is Altered by Copy Number Loss, Mutation, and/or Loss of Protein Expression in More Than 70% ofÂMalignant Peritoneal Mesotheliomas. Journal of Thoracic Oncology, 2017, 12, 724-733.	1.1	67
7	Sub-clonal analysis of the murine C1498 acute myeloid leukaemia cell line reveals genomic and immunogenic diversity. Immunology Letters, 2017, 192, 27-34.	2.5	1
8	RNA-binding disturbances as a continuum from spinocerebellar ataxia type 2 to Parkinson disease. Neurobiology of Disease, 2016, 96, 312-322.	4.4	14
9	Polymorphisms in the Mannose-Binding Lectin Gene are Associated with Defective Mannose-Binding Lectin Functional Activity in Crohnâ€™s Disease Patients. Scientific Reports, 2016, 6, 29636.	3.3	11
10	Genomic copy number alterations in 33 malignant peritoneal mesothelioma analyzed by comparative genomic hybridization array. Human Pathology, 2016, 55, 72-82.	2.0	40
11	Genomic Landscape of CXCR4 Mutations in WaldenstrÃ¶m Macroglobulinemia. Clinical Cancer Research, 2016, 22, 1480-1488.	7.0	102
12	Involvement of the immune system, endocytosis and EIF2 signaling in both genetically determined and sporadic forms of Parkinson's disease. Neurobiology of Disease, 2014, 63, 165-170.	4.4	53
13	Neurofibromatosisâ€1 gene deletions and mutations in de novo adult acute myeloid leukemia. American Journal of Hematology, 2013, 88, 306-311.	4.1	43
14	Transcriptional profile of Parkinson blood mononuclear cells with LRRK2 mutation. Neurobiology of Aging, 2011, 32, 1839-1848.	3.1	83
15	Translation Initiator EIF4G1 Mutations in Familial Parkinson Disease. American Journal of Human Genetics, 2011, 89, 398-406.	6.2	250
16	SNCA locus duplication carriers: from genetics to Parkinson disease phenotypes. Human Mutation, 2011, 32, E2079-90.	2.5	34
17	Expression of CD34 in hematopoietic cancer cell lines reflects tightly regulated stem/progenitorâ€like state. Journal of Cellular Biochemistry, 2011, 112, 1277-1285.	2.6	14
18	Genomic characterization of Imatinib resistance in CD34+ cell populations from chronic myeloid leukaemia patients. Leukemia Research, 2011, 35, 448-458.	0.8	17

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19	Deletion 18q21.2q21.32 involving TCF4 in a boy diagnosed by CGH-array. European Journal of Medical Genetics, 2008, 51, 172-177.	1.3	28
20	Characterization by array-CGH of an interstitial de novo tandem 6p21.2p22.1 duplication in a boy with epilepsy and developmental delay. European Journal of Medical Genetics, 2008, 51, 373-381.	1.3	9
21	EIF4A2 Is a Positional Candidate Gene at the 3q27 Locus Linked to Type 2 Diabetes in French Families. Diabetes, 2006, 55, 1171-1176.	0.6	23
22	Genetic Analysis of ADIPOR1 and ADIPOR2 Candidate Polymorphisms for Type 2 Diabetes in the Caucasian Population. Diabetes, 2006, 55, 856-861.	0.6	72
23	A Synonymous Coding Polymorphism in the Â2-Heremans-Schmid Glycoprotein Gene Is Associated With Type 2 Diabetes in French Caucasians. Diabetes, 2005, 54, 2477-2481.	0.6	83
24	A CD36 nonsense mutation associated with insulin resistance and familial type 2 diabetes. Human Mutation, 2004, 24, 104-104.	2.5	51
25	A promoter polymorphism in CD36 is associated with an atherogenic lipid profile in a French general population. Atherosclerosis, 2004, 173, 373-375.	0.8	23
26	VE-statin, an endothelial repressor of smooth muscle cell migration. EMBO Journal, 2003, 22, 5700-5711.	7.8	112
27	The genetics of adiponectin. Current Diabetes Reports, 2003, 3, 151-158.	4.2	84
28	Single-nucleotide polymorphism haplotypes in the both proximal promoter and exon 3 of the APM1 gene modulate adipocyte-secreted adiponectin hormone levels and contribute to the genetic risk for type 2 diabetes in French Caucasians. Human Molecular Genetics, 2002, 11, 2607-2614.	2.9	433
29	Cloning and Characterization of the Human and Rat Islet-specific Glucose-6-phosphatase Catalytic Subunit-related Protein (IGRP) Genes. Journal of Biological Chemistry, 2001, 276, 25197-25207.	3.4	68
30	Genomewide Search for Type 2 Diabetesâ€“Susceptibility Genes in French Whites: Evidence for a Novel Susceptibility Locus for Early-Onset Diabetes on Chromosome 3q27-qter and Independent Replication of a Type 2â€“Diabetes Locus on Chromosome 1q21â€“q24. American Journal of Human Genetics, 2000, 67, 1470-1480.	6.2	630