

Philippe arnaud

List of Publications by Citations

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

47 papers	2,144 citations	25 h-index	46 g-index
53 ext. papers	2,404 ext. citations	7.7 avg, IF	4.46 L-index

#	Paper	IF	Citations
47	Limited evolutionary conservation of imprinting in the human placenta. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2006 , 103, 6623-8	11.5	208
46	Imprinting of IGF2 P0 transcript and novel alternatively spliced INS-IGF2 isoforms show differences between mouse and human. <i>Human Molecular Genetics</i> , 2006 , 15, 1259-69	5.6	123
45	Conserved methylation imprints in the human and mouse GRB10 genes with divergent allelic expression suggests differential reading of the same mark. <i>Human Molecular Genetics</i> , 2003 , 12, 1005-19	5.6	121
44	Deep sequencing and de novo assembly of the mouse oocyte transcriptome define the contribution of transcription to the DNA methylation landscape. <i>Genome Biology</i> , 2015 , 16, 209	18.3	117
43	Histone methylation is mechanistically linked to DNA methylation at imprinting control regions in mammals. <i>Human Molecular Genetics</i> , 2009 , 18, 3375-83	5.6	105
42	Epigenetic properties and identification of an imprint mark in the Nesp-Gnasxl domain of the mouse Gnas imprinted locus. <i>Molecular and Cellular Biology</i> , 2003 , 23, 5475-88	4.8	102
41	Improved thermostability of the North American firefly luciferase: saturation mutagenesis at position 354. <i>Biochemical Journal</i> , 1996 , 319 (Pt 2), 343-50	3.8	98
40	Ring1B and Suv39h1 delineate distinct chromatin states at bivalent genes during early mouse lineage commitment. <i>Development (Cambridge)</i> , 2010 , 137, 2483-92	6.6	90
39	Epigenetic deregulation of genomic imprinting in human disorders and following assisted reproduction. <i>Birth Defects Research Part C: Embryo Today Reviews</i> , 2005 , 75, 81-97		79
38	SINE retroposons can be used in vivo as nucleation centers for de novo methylation. <i>Molecular and Cellular Biology</i> , 2000 , 20, 3434-41	4.8	79
37	Genomic imprinting in germ cells: imprints are under control. <i>Reproduction</i> , 2010 , 140, 411-23	3.8	71
36	Reciprocal imprinting of human GRB10 in placental trophoblast and brain: evolutionary conservation of reversed allelic expression. <i>Human Molecular Genetics</i> , 2009 , 18, 3066-74	5.6	70
35	Identification of the imprinted KLF14 transcription factor undergoing human-specific accelerated evolution. <i>PLoS Genetics</i> , 2007 , 3, e65	6	70
34	Comparative analysis of human chromosome 7q21 and mouse proximal chromosome 6 reveals a placental-specific imprinted gene, TFPI2/Tfpi2, which requires EHMT2 and EED for allelic-silencing. <i>Genome Research</i> , 2008 , 18, 1270-81	9.7	69
33	A mono-allelic bivalent chromatin domain controls tissue-specific imprinting at Grb10. <i>EMBO Journal</i> , 2008 , 27, 2523-32	13	63
32	Transcription and histone methylation changes correlate with imprint acquisition in male germ cells. <i>EMBO Journal</i> , 2012 , 31, 606-15	13	62
31	Stochastic imprinting in the progeny of Dnmt3L ^{-/-} females. <i>Human Molecular Genetics</i> , 2006 , 15, 589-98	5.6	55

30	ICR noncoding RNA expression controls imprinting and DNA replication at the Dlk1-Dio3 domain. <i>Developmental Cell</i> , 2014 , 31, 19-33	10.2	53
29	The mouse Zac1 locus: basis for imprinting and comparison with human ZAC. <i>Gene</i> , 2002 , 292, 101-12	3.8	46
28	Genome-wide identification of new imprinted genes. <i>Briefings in Functional Genomics</i> , 2010 , 9, 304-14	4.9	40
27	Synergic reprogramming of mammalian cells by combined exposure to mitotic <i>Xenopus</i> egg extracts and transcription factors. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2011 , 108, 17331-6	11.5	36
26	Liver x receptors protect from development of prostatic intra-epithelial neoplasia in mice. <i>PLoS Genetics</i> , 2013 , 9, e1003483	6	33
25	Imprinting control regions (ICRs) are marked by mono-allelic bivalent chromatin when transcriptionally inactive. <i>Nucleic Acids Research</i> , 2016 , 44, 621-35	20.1	29
24	Analysis of the SINE S1 Pol III promoter from Brassica; impact of methylation and influence of external sequences. <i>Plant Journal</i> , 2001 , 26, 295-305	6.9	27
23	S1 SINE retroposons are methylated at symmetrical and non-symmetrical positions in Brassica napus: identification of a preferred target site for asymmetrical methylation. <i>Plant Molecular Biology</i> , 1999 , 39, 243-55	4.6	27
22	A transcriptional analysis of the S1Bn (Brassica napus) family of SINE retroposons. <i>Plant Molecular Biology</i> , 1996 , 32, 869-78	4.6	25
21	An annotated list of bivalent chromatin regions in human ES cells: a new tool for cancer epigenetic research. <i>Oncotarget</i> , 2017 , 8, 4110-4124	3.3	24
20	Chromatin Immunoprecipitation (ChIP) on Unfixed Chromatin from Cells and Tissues to Analyze Histone Modifications. <i>Cold Spring Harbor Protocols</i> , 2007 , 2007, pdb.prot4767	1.2	22
19	Characterization of novel paternal ncRNAs at the Plagl1 locus, including Hymai, predicted to interact with regulators of active chromatin. <i>PLoS ONE</i> , 2012 , 7, e38907	3.7	21
18	Human imprinted retrogenes exhibit non-canonical imprint chromatin signatures and reside in non-imprinted host genes. <i>Nucleic Acids Research</i> , 2011 , 39, 4577-86	20.1	18
17	Transcriptional alterations in glioma result primarily from DNA methylation-independent mechanisms. <i>Genome Research</i> , 2019 , 29, 1605-1621	9.7	17
16	DNA methylation profiling reveals a pathological signature that contributes to transcriptional defects of CD34 CD15 cells in early chronic-phase chronic myeloid leukemia. <i>Molecular Oncology</i> , 2018 , 12, 814-829	7.9	17
15	The long non-coding RNA is transcriptionally activated by HOXA9 and is an independent prognostic marker in patients with malignant glioma. <i>Oncotarget</i> , 2018 , 9, 15740-15756	3.3	17
14	Identification and properties of imprinted genes and their control elements. <i>Cytogenetic and Genome Research</i> , 2004 , 105, 335-45	1.9	15
13	Detection of the alternative lengthening of telomeres pathway in malignant gliomas for improved molecular diagnosis. <i>Journal of Neuro-Oncology</i> , 2017 , 135, 381-390	4.8	14

12	MEDEA takes control of its own imprinting. <i>Cell</i> , 2006 , 124, 468-70	56.2	13
11	Imprinted methylation profiles for proximal mouse chromosomes 11 and 7 as revealed by methylation-sensitive representational difference analysis. <i>Mammalian Genome</i> , 2003 , 14, 805-16	3.2	13
10	In Vitro Corticogenesis from Embryonic Stem Cells Recapitulates the In Vivo Epigenetic Control of Imprinted Gene Expression. <i>Cerebral Cortex</i> , 2017 , 27, 2418-2433	5.1	11
9	HOX gene cluster (de)regulation in brain: from neurodevelopment to malignant glial tumours. <i>Cellular and Molecular Life Sciences</i> , 2020 , 77, 3797-3821	10.3	10
8	The tumoral A genotype of the MGMT rs34180180 single-nucleotide polymorphism in aggressive gliomas is associated with shorter patientsTsurvival. <i>Carcinogenesis</i> , 2016 , 37, 169-176	4.6	10
7	Plant S1 SINEs as a model to study retroposition. <i>Genetica</i> , 1997 , 100, 155-160	1.5	7
6	CLIFinder: identification of LINE-1 chimeric transcripts in RNA-seq data. <i>Bioinformatics</i> , 2018 , 34, 688-690	7.2	6
5	Early mouse embryo development: could epigenetics influence cell fate determination?. <i>BioEssays</i> , 2007 , 29, 520-4	4.1	3
4	PCR-Based Analysis of Immunoprecipitated Chromatin. <i>Cold Spring Harbor Protocols</i> , 2007 , 2007, pdb.prot47683	1.4	3
3	TET3 controls the expression of the H3K27me3 demethylase Kdm6b during neural commitment. <i>Cellular and Molecular Life Sciences</i> , 2021 , 78, 757-768	10.3	3
2	Widespread overexpression from the four DNA hypermethylated HOX clusters in aggressive (IDHwt) glioma is associated with H3K27me3 depletion and alternative promoter usage. <i>Molecular Oncology</i> , 2021 , 15, 1995-2010	7.9	1
1	Epigenetic Reprogramming in the Mammalian Germline 2013 , 3-34		