

Philippe arnaud

List of Publications by Year in Descending Order

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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.

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|-------------------|-------------------------|----------------|-----------------|
| 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 | 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 |
| 46 | 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 |
| 45 | 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 |
| 44 | Transcriptional alterations in glioma result primarily from DNA methylation-independent mechanisms. <i>Genome Research</i> , 2019 , 29, 1605-1621 | 9.7 | 17 |
| 43 | 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 |
| 42 | CLIFinder: identification of LINE-1 chimeric transcripts in RNA-seq data. <i>Bioinformatics</i> , 2018 , 34, 688-690 | 9.2 | 6 |
| 41 | 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 |
| 40 | 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 |
| 39 | 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 |
| 38 | 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 |
| 37 | 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 |
| 36 | The tumoral A genotype of the MGMT rs34180180 single-nucleotide polymorphism in aggressive gliomas is associated with shorter patients survival. <i>Carcinogenesis</i> , 2016 , 37, 169-176 | 4.6 | 10 |
| 35 | 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 |
| 34 | 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 |
| 33 | Liver x receptors protect from development of prostatic intra-epithelial neoplasia in mice. <i>PLoS Genetics</i> , 2013 , 9, e1003483 | 6 | 33 |
| 32 | Epigenetic Reprogramming in the Mammalian Germline 2013 , 3-34 | | |
| 31 | 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 |

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| 30 | Transcription and histone methylation changes correlate with imprint acquisition in male germ cells. <i>EMBO Journal</i> , 2012 , 31, 606-15 | 13 | 62 |
| 29 | 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 |
| 28 | 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 |
| 27 | 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 |
| 26 | Genome-wide identification of new imprinted genes. <i>Briefings in Functional Genomics</i> , 2010 , 9, 304-14 | 4.9 | 40 |
| 25 | Genomic imprinting in germ cells: imprints are under control. <i>Reproduction</i> , 2010 , 140, 411-23 | 3.8 | 71 |
| 24 | 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 |
| 23 | 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 |
| 22 | A mono-allelic bivalent chromatin domain controls tissue-specific imprinting at Grb10. <i>EMBO Journal</i> , 2008 , 27, 2523-32 | 13 | 63 |
| 21 | 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 |
| 20 | Early mouse embryo development: could epigenetics influence cell fate determination?. <i>BioEssays</i> , 2007 , 29, 520-4 | 4.1 | 3 |
| 19 | Identification of the imprinted KLF14 transcription factor undergoing human-specific accelerated evolution. <i>PLoS Genetics</i> , 2007 , 3, e65 | 6 | 70 |
| 18 | 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 |
| 17 | PCR-Based Analysis of Immunoprecipitated Chromatin. <i>Cold Spring Harbor Protocols</i> , 2007 , 2007, pdb.prot4768 | 1.2 | 23 |
| 16 | Stochastic imprinting in the progeny of Dnmt3L ^{-/-} females. <i>Human Molecular Genetics</i> , 2006 , 15, 589-98 | 5.6 | 55 |
| 15 | 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 |
| 14 | 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 |
| 13 | MEDEA takes control of its own imprinting. <i>Cell</i> , 2006 , 124, 468-70 | 56.2 | 13 |

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| 12 | 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 |
| 11 | Identification and properties of imprinted genes and their control elements. <i>Cytogenetic and Genome Research</i> , 2004 , 105, 335-45 | 1.9 | 15 |
| 10 | 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 |
| 9 | 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 |
| 8 | 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 |
| 7 | The mouse Zac1 locus: basis for imprinting and comparison with human ZAC. <i>Gene</i> , 2002 , 292, 101-12 | 3.8 | 46 |
| 6 | 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 |
| 5 | 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 |
| 4 | 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 |
| 3 | Plant S1 SINEs as a model to study retroposition. <i>Genetica</i> , 1997 , 100, 155-160 | 1.5 | 7 |
| 2 | 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 |
| 1 | A transcriptional analysis of the S1Bn (Brassica napus) family of SINE retroposons. <i>Plant Molecular Biology</i> , 1996 , 32, 869-78 | 4.6 | 25 |