

Sebastian M Waszak

List of Publications by Year in descending order

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Version: 2024-02-01

62
papers

9,761
citations

147726
31
h-index

197736
49
g-index

73
all docs

73
docs citations

73
times ranked

17930
citing authors

#	ARTICLE	IF	CITATIONS
1	Pan-cancer analysis of whole genomes. <i>Nature</i> , 2020, 578, 82-93.	13.7	1,966
2	The landscape of genomic alterations across childhood cancers. <i>Nature</i> , 2018, 555, 321-327.	13.7	1,068
3	The whole-genome landscape of medulloblastoma subtypes. <i>Nature</i> , 2017, 547, 311-317.	13.7	787
4	Patterns of somatic structural variation in human cancer genomes. <i>Nature</i> , 2020, 578, 112-121.	13.7	560
5	Variation in Transcription Factor Binding Among Humans. <i>Science</i> , 2010, 328, 232-235.	6.0	521
6	Analyses of non-coding somatic drivers in 2,658 cancer whole genomes. <i>Nature</i> , 2020, 578, 102-111.	13.7	424
7	A Dual Program for Translation Regulation in Cellular Proliferation and Differentiation. <i>Cell</i> , 2014, 158, 1281-1292.	13.5	414
8	Coordinated Effects of Sequence Variation on DNA Binding, Chromatin Structure, and Transcription. <i>Science</i> , 2013, 342, 744-747.	6.0	364
9	Pan-cancer analysis of somatic copy-number alterations implicates <i>IRS4</i> and <i>IGF2</i> in enhancer hijacking. <i>Nature Genetics</i> , 2017, 49, 65-74.	9.4	326
10	Active medulloblastoma enhancers reveal subgroup-specific cellular origins. <i>Nature</i> , 2016, 530, 57-62.	13.7	318
11	Genomic basis for RNA alterations in cancer. <i>Nature</i> , 2020, 578, 129-136.	13.7	280
12	Pan-cancer analysis of whole genomes identifies driver rearrangements promoted by LINE-1 retrotransposition. <i>Nature Genetics</i> , 2020, 52, 306-319.	9.4	275
13	Spectrum and prevalence of genetic predisposition in medulloblastoma: a retrospective genetic study and prospective validation in a clinical trial cohort. <i>Lancet Oncology</i> , 2018, 19, 785-798.	5.1	268
14	Population Variation and Genetic Control of Modular Chromatin Architecture in Humans. <i>Cell</i> , 2015, 162, 1039-1050.	13.5	210
15	Molecular Evolution of Early-Onset Prostate Cancer Identifies Molecular Risk Markers and Clinical Trajectories. <i>Cancer Cell</i> , 2018, 34, 996-1011.e8.	7.7	190
16	Risk-adapted therapy for young children with medulloblastoma (SJYC07): therapeutic and molecular outcomes from a multicentre, phase 2 trial. <i>Lancet Oncology</i> , 2018, 19, 768-784.	5.1	151
17	A cell-based model system links chromothripsis with hyperploidy. <i>Molecular Systems Biology</i> , 2015, 11, 828.	3.2	118
18	Relating CNVs to transcriptome data at fine resolution: Assessment of the effect of variant size, type, and overlap with functional regions. <i>Genome Research</i> , 2011, 21, 2004-2013.	2.4	109

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19	Genomic Variation and Its Impact on Gene Expression in <i>Drosophila melanogaster</i> . <i>PLoS Genetics</i> , 2012, 8, e1003055.	1.5	102
20	Identification of the transcription factor ZEB1 as a central component of the adipogenic gene regulatory network. <i>ELife</i> , 2014, 3, e03346.	2.8	101
21	Genomic and transcriptomic changes complement each other in the pathogenesis of sporadic Burkitt lymphoma. <i>Nature Communications</i> , 2019, 10, 1459.	5.8	99
22	Integrative Genomics Identifies the Corepressor SMRT as a Gatekeeper of Adipogenesis through the Transcription Factors C/EBP β and KAISO. <i>Molecular Cell</i> , 2012, 46, 335-350.	4.5	96
23	The molecular landscape of ETMR at diagnosis and relapse. <i>Nature</i> , 2019, 576, 274-280.	13.7	94
24	Germline Elongator mutations in Sonic Hedgehog medulloblastoma. <i>Nature</i> , 2020, 580, 396-401.	13.7	94
25	Personal receptor repertoires: olfaction as a model. <i>BMC Genomics</i> , 2012, 13, 414.	1.2	92
26	Antibiotics-induced monodominance of a novel gut bacterial order. <i>Gut</i> , 2019, 68, 1781-1790.	6.1	73
27	Chromosomal contacts connect loci associated with autism, BMI and head circumference phenotypes. <i>Molecular Psychiatry</i> , 2017, 22, 836-849.	4.1	68
28	Systematic Inference of Copy-Number Genotypes from Personal Genome Sequencing Data Reveals Extensive Olfactory Receptor Gene Content Diversity. <i>PLoS Computational Biology</i> , 2010, 6, e1000988.	1.5	56
29	Germline <i>GPR161</i> Mutations Predispose to Pediatric Medulloblastoma. <i>Journal of Clinical Oncology</i> , 2020, 38, 43-50.	0.8	50
30	Pharmaco-proteogenomic profiling of pediatric diffuse midline glioma to inform future treatment strategies. <i>Oncogene</i> , 2022, 41, 461-475.	2.6	39
31	<i>PDX</i> models recapitulate the genetic and epigenetic landscape of pediatric T-cell leukemia. <i>EMBO Molecular Medicine</i> , 2018, 10, .	3.3	38
32	Imipridones affect tumor bioenergetics and promote cell lineage differentiation in diffuse midline gliomas. <i>Neuro-Oncology</i> , 2022, 24, 1438-1451.	0.6	36
33	A yeast one-hybrid and microfluidics-based pipeline to map mammalian gene regulatory networks. <i>Molecular Systems Biology</i> , 2013, 9, 682.	3.2	35
34	Mutational mechanisms shaping the coding and noncoding genome of germinal center derived B-cell lymphomas. <i>Leukemia</i> , 2021, 35, 2002-2016.	3.3	34
35	Mechanisms of imipridones in targeting mitochondrial metabolism in cancer cells. <i>Neuro-Oncology</i> , 2021, 23, 542-556.	0.6	30
36	No correlation between NF1 mutation position and risk of optic pathway glioma in 77 unrelated NF1 patients. <i>Human Genetics</i> , 2016, 135, 469-475.	1.8	29

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37	Retrospective evaluation of whole exome and genome mutation calls in 746 cancer samples. <i>Nature Communications</i> , 2020, 11, 4748.	5.8	27
38	Genome-wide Screens Implicate Loss of Cullin Ring Ligase 3 in Persistent Proliferation and Genome Instability in TP53-Deficient Cells. <i>Cell Reports</i> , 2020, 31, 107465.	2.9	24
39	Identification and removal of low-complexity sites in allele-specific analysis of ChIP-seq data. <i>Bioinformatics</i> , 2014, 30, 165-171.	1.8	21
40	NCoR1: Putting the Brakes on the Dendritic Cell Immune Tolerance. <i>IScience</i> , 2019, 19, 996-1011.	1.9	20
41	Chromatin accessibility landscape of pediatric T-lymphoblastic leukemia and human T-cell precursors. <i>EMBO Molecular Medicine</i> , 2020, 12, e12104.	3.3	13
42	Butler enables rapid cloud-based analysis of thousands of human genomes. <i>Nature Biotechnology</i> , 2020, 38, 288-292.	9.4	11
43	Versatile workflow for cell type-resolved transcriptional and epigenetic profiles from cryopreserved human lung. <i>JCI Insight</i> , 2021, 6, .	2.3	8
44	A leukemia-protective germline variant mediates chromatin module formation via transcription factor nucleation. <i>Nature Communications</i> , 2022, 13, 2042.	5.8	6
45	Cancer risk and tumour spectrum in 172 patients with a germline <i>SUFU</i> pathogenic variation: a collaborative study of the SIOPE Host Genome Working Group. <i>Journal of Medical Genetics</i> , 2022, 59, 1123-1132.	1.5	4
46	Variation in transcription factor binding among humans. <i>New Biotechnology</i> , 2010, 27, S81.	2.4	3
47	A common microdeletion affecting a hippocampus- and amygdala-specific isoform of tryptophan hydroxylase 2 is not associated with affective disorders. <i>Bipolar Disorders</i> , 2014, 16, 764-768.	1.1	2
48	HGG-32. ONC201 AND ONC206 TARGET TUMOR CELL METABOLISM IN PEDIATRIC DIFFUSE MIDLINE GLIOMA PRECLINICAL MODELS. <i>Neuro-Oncology</i> , 2021, 23, i23-i24.	0.6	2
49	Rare Germline Variants Are Associated with Rapid Biochemical Recurrence After Radical Prostate Cancer Treatment: A Pan Prostate Cancer Group Study. <i>European Urology</i> , 2022, 82, 201-211.	0.9	2
50	Rounding Up Natural Gene Expression Variation during Development. <i>Developmental Cell</i> , 2013, 27, 601-603.	3.1	1
51	MEDB-14. Clinical outcome of pediatric medulloblastoma patients with Li-Fraumeni syndrome. <i>Neuro-Oncology</i> , 2022, 24, i107-i107.	0.6	1
52	EPIG-04 THE CHROMATIN LANDSCAPE OF MEDULLOBLASTOMA. <i>Neuro-Oncology</i> , 2015, 17, v86.4-v87.	0.6	0
53	MBCL-44. THE MOLECULAR AND CLINICAL LANDSCAPE OF INFANT MEDULLOBLASTOMA (iMB): RESULTS AND MOLECULAR ANALYSIS FROM A PROSPECTIVE, MULTICENTER PHASE II TRIAL (SJYC07). <i>Neuro-Oncology</i> , 2018, 20, i126-i127.	0.6	0
54	EMBR-10. GENOMIC COMPLEXITY AND EVOLUTION OF EMBRYONAL TUMORS WITH MULTILAYERED ROSETTES (ETMR). <i>Neuro-Oncology</i> , 2018, 20, i70-i70.	0.6	0

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55	MEDU-11. MOLECULAR CHARACTERIZATION OF ETMRs REVEALS A ROLE FOR R-LOOP MEDIATED CHROMOSOMAL INSTABILITY. <i>Neuro-Oncology</i> , 2019, 21, ii105-ii105.	0.6	0
56	Abstract LB-B23: Medulloblastoma regulatory circuitries reveal subgroup-specific cellular origins. , 2015, , .		0
57	NCoR1 is a master repressor of the tolerogenic program in dendritic cells. <i>Canadian Journal of Biotechnology</i> , 2017, 1, 161-161.	0.3	0
58	Abstract 3172: Targeting genomic instability in embryonal tumors with multilayered rosettes (ETMR). , 2018, , .		0
59	Pediatric T-ALLs Developing into a Type 2 Relapse Originate from Cells That Carry the Potential of Variable Maturation into Subclones with Distinct Chromatin Landscapes. <i>Blood</i> , 2018, 132, 1545-1545.	0.6	0
60	DIPG-64. INTERNATIONAL PRECLINICAL DRUG DISCOVERY AND BIOMARKER PROGRAM INFORMING AN ADOPTIVE COMBINATORIAL TRIAL FOR DIFFUSE MIDLINE GLIOMAS. <i>Neuro-Oncology</i> , 2020, 22, iii300-iii300.	0.6	0
61	MBCL-21. GERMLINE ELONGATOR MUTATIONS IN SONIC HEDGEHOG MEDULLOBLASTOMA. <i>Neuro-Oncology</i> , 2020, 22, iii392-iii393.	0.6	0
62	DIPG-07. Preclinical and case study results underpinning the phase II clinical trial testing the combination of ONC201 and paxalisib for the treatment of patients with diffuse midline glioma (NCT05009992). <i>Neuro-Oncology</i> , 2022, 24, i18-i19.	0.6	0