

# Sebastian M Waszak

## 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.

62

papers

6,046

citations

34

h-index

73

g-index

73

ext. papers

8,712

ext. citations

17.8

avg, IF

7.1

L-index

#	Paper	IF	Citations
62	Pan-cancer analysis of whole genomes. <i>Nature</i> , <b>2020</b> , 578, 82-93	50.4	840
61	The landscape of genomic alterations across childhood cancers. <i>Nature</i> , <b>2018</b> , 555, 321-327	50.4	603
60	The whole-genome landscape of medulloblastoma subtypes. <i>Nature</i> , <b>2017</b> , 547, 311-317	50.4	472
59	Variation in transcription factor binding among humans. <i>Science</i> , <b>2010</b> , 328, 232-5	33.3	447
58	A dual program for translation regulation in cellular proliferation and differentiation. <i>Cell</i> , <b>2014</b> , 158, 1281-1292	56.2	278
57	Coordinated effects of sequence variation on DNA binding, chromatin structure, and transcription. <i>Science</i> , <b>2013</b> , 342, 744-7	33.3	278
56	Active medulloblastoma enhancers reveal subgroup-specific cellular origins. <i>Nature</i> , <b>2016</b> , 530, 57-62	50.4	234
55	Patterns of somatic structural variation in human cancer genomes. <i>Nature</i> , <b>2020</b> , 578, 112-121	50.4	232
54	Analyses of non-coding somatic drivers in 2,658 cancer whole genomes. <i>Nature</i> , <b>2020</b> , 578, 102-111	50.4	220
53	Pan-cancer analysis of somatic copy-number alterations implicates IRS4 and IGF2 in enhancer hijacking. <i>Nature Genetics</i> , <b>2017</b> , 49, 65-74	36.3	220
52	Spectrum and prevalence of genetic predisposition in medulloblastoma: a retrospective genetic study and prospective validation in a clinical trial cohort. <i>Lancet Oncology, The</i> , <b>2018</b> , 19, 785-798	21.7	159
51	Population Variation and Genetic Control of Modular Chromatin Architecture in Humans. <i>Cell</i> , <b>2015</b> , 162, 1039-50	56.2	156
50	Genomic basis for RNA alterations in cancer. <i>Nature</i> , <b>2020</b> , 578, 129-136	50.4	148
49	Pan-cancer analysis of whole genomes identifies driver rearrangements promoted by LINE-1 retrotransposition. <i>Nature Genetics</i> , <b>2020</b> , 52, 306-319	36.3	122
48	Risk-adapted therapy for young children with medulloblastoma (SJYC07): therapeutic and molecular outcomes from a multicentre, phase 2 trial. <i>Lancet Oncology, The</i> , <b>2018</b> , 19, 768-784	21.7	95
47	Molecular Evolution of Early-Onset Prostate Cancer Identifies Molecular Risk Markers and Clinical Trajectories. <i>Cancer Cell</i> , <b>2018</b> , 34, 996-1011.e8	24.3	89
46	A cell-based model system links chromothripsis with hyperploidy. <i>Molecular Systems Biology</i> , <b>2015</b> , 11, 828	12.2	88

45	Genomic variation and its impact on gene expression in <i>Drosophila melanogaster</i> . <i>PLoS Genetics</i> , <b>2012</b> , 8, e1003055	6	85
44	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> , <b>2011</b> , 21, 2004-13	9.7	80
43	EMBR-10. GENOMIC COMPLEXITY AND EVOLUTION OF EMBRYONAL TUMORS WITH MULTILAYERED ROSETTES (ETMR). <i>Neuro-Oncology</i> , <b>2018</b> , 20, i70-i70	1	78
42	MEDU-11. MOLECULAR CHARACTERIZATION OF ETMRs REVEALS A ROLE FOR R-LOOP MEDIATED CHROMOSOMAL INSTABILITY. <i>Neuro-Oncology</i> , <b>2019</b> , 21, ii105-ii105	1	78
41	EPIG-04 THE CHROMATIN LANDSCAPE OF MEDULLOBLASTOMA. <i>Neuro-Oncology</i> , <b>2015</b> , 17, v86.4-v87	1	78
40	DIPG-64. INTERNATIONAL PRECLINICAL DRUG DISCOVERY AND BIOMARKER PROGRAM INFORMING AN ADOPTIVE COMBINATORIAL TRIAL FOR DIFFUSE MIDLINE GLIOMAS. <i>Neuro-Oncology</i> , <b>2020</b> , 22, iii300-iii300	1	78
39	MBCL-21. GERMLINE ELONGATOR MUTATIONS IN SONIC HEDGEHOG MEDULLOBLASTOMA. <i>Neuro-Oncology</i> , <b>2020</b> , 22, iii392-iii393	1	78
38	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> , <b>2018</b> , 20, i126-i127	1	78
37	Personal receptor repertoires: olfaction as a model. <i>BMC Genomics</i> , <b>2012</b> , 13, 414	4.5	74
36	Integrative genomics identifies the corepressor SMRT as a gatekeeper of adipogenesis through the transcription factors C/EBP $\beta$ and KAISO. <i>Molecular Cell</i> , <b>2012</b> , 46, 335-50	17.6	62
35	Identification of the transcription factor ZEB1 as a central component of the adipogenic gene regulatory network. <i>ELife</i> , <b>2014</b> , 3, e03346	8.9	60
34	Systematic inference of copy-number genotypes from personal genome sequencing data reveals extensive olfactory receptor gene content diversity. <i>PLoS Computational Biology</i> , <b>2010</b> , 6, e1000988	5	54
33	Genomic and transcriptomic changes complement each other in the pathogenesis of sporadic Burkitt lymphoma. <i>Nature Communications</i> , <b>2019</b> , 10, 1459	17.4	49
32	Germline Elongator mutations in Sonic Hedgehog medulloblastoma. <i>Nature</i> , <b>2020</b> , 580, 396-401	50.4	47
31	The molecular landscape of ETMR at diagnosis and relapse. <i>Nature</i> , <b>2019</b> , 576, 274-280	50.4	46
30	Chromosomal contacts connect loci associated with autism, BMI and head circumference phenotypes. <i>Molecular Psychiatry</i> , <b>2017</b> , 22, 836-849	15.1	43
29	The whole-genome panorama of cancer drivers		38
28	Antibiotics-induced monodominance of a novel gut bacterial order. <i>Gut</i> , <b>2019</b> , 68, 1781-1790	19.2	33

27	A yeast one-hybrid and microfluidics-based pipeline to map mammalian gene regulatory networks. <i>Molecular Systems Biology</i> , <b>2013</b> , 9, 682	12.2	31
26	Germline Mutations Predispose to Pediatric Medulloblastoma. <i>Journal of Clinical Oncology</i> , <b>2020</b> , 38, 43-50	2.2	28
25	No correlation between NF1 mutation position and risk of optic pathway glioma in 77 unrelated NF1 patients. <i>Human Genetics</i> , <b>2016</b> , 135, 469-475	6.3	21
24	PDX models recapitulate the genetic and epigenetic landscape of pediatric T-cell leukemia. <i>EMBO Molecular Medicine</i> , <b>2018</b> , 10,	12	20
23	Identification and removal of low-complexity sites in allele-specific analysis of CHIP-seq data. <i>Bioinformatics</i> , <b>2014</b> , 30, 165-71	7.2	17
22	Germline determinants of the somatic mutation landscape in 2,642 cancer genomes		13
21	Genome-wide Screens Implicate Loss of Cullin Ring Ligase 3 in Persistent Proliferation and Genome Instability in TP53-Deficient Cells. <i>Cell Reports</i> , <b>2020</b> , 31, 107465	10.6	11
20	Mechanisms of imipridones in targeting mitochondrial metabolism in cancer cells. <i>Neuro-Oncology</i> , <b>2021</b> , 23, 542-556	1	11
19	NCoR1: Putting the Brakes on the Dendritic Cell Immune Tolerance. <i>iScience</i> , <b>2019</b> , 19, 996-1011	6.1	10
18	Pan-cancer analysis of whole genomes reveals driver rearrangements promoted by LINE-1 retrotransposition in human tumours		10
17	Genomic basis for RNA alterations revealed by whole-genome analyses of 27 cancer types		10
16	Retrospective evaluation of whole exome and genome mutation calls in 746 cancer samples. <i>Nature Communications</i> , <b>2020</b> , 11, 4748	17.4	10
15	Butler enables rapid cloud-based analysis of thousands of human genomes. <i>Nature Biotechnology</i> , <b>2020</b> , 38, 288-292	44.5	9
14	Pharmaco-proteogenomic profiling of pediatric diffuse midline glioma to inform future treatment strategies. <i>Oncogene</i> , <b>2021</b> ,	9.2	5
13	Assessing the Gene Regulatory Landscape in 1,188 Human Tumors		3
12	Chromatin accessibility landscape of pediatric T-lymphoblastic leukemia and human T-cell precursors. <i>EMBO Molecular Medicine</i> , <b>2020</b> , 12, e12104	12	3
11	Mutational mechanisms shaping the coding and noncoding genome of germinal center derived B-cell lymphomas. <i>Leukemia</i> , <b>2021</b> , 35, 2002-2016	10.7	3
10	A common microdeletion affecting a hippocampus- and amygdala-specific isoform of tryptophan hydroxylase 2 is not associated with affective disorders. <i>Bipolar Disorders</i> , <b>2014</b> , 16, 764-8	3.8	2

9	Imipridones affect tumor bioenergetics and promote cell lineage differentiation in diffuse midline gliomas.. <i>Neuro-Oncology</i> , <b>2022</b> ,	1	2
8	Enabling rapid cloud-based analysis of thousands of human genomes via Butler		2
7	Versatile workflow for cell type-resolved transcriptional and epigenetic profiles from cryopreserved human lung. <i>JCI Insight</i> , <b>2021</b> , 6,	9.9	2
6	A leukemia-protective germline variant mediates chromatin module formation via transcription factor nucleation.. <i>Nature Communications</i> , <b>2022</b> , 13, 2042	17.4	1
5	Rounding up natural gene expression variation during development. <i>Developmental Cell</i> , <b>2013</b> , 27, 601-310.2	10.2	0
4	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> , <b>2018</b> , 132, 1545-1545	2.2	
3	HGG-32. ONC201 AND ONC206 TARGET TUMOR CELL METABOLISM IN PEDIATRIC DIFFUSE MIDLINE GLIOMA PRECLINICAL MODELS. <i>Neuro-Oncology</i> , <b>2021</b> , 23, i23-i24	1	
2	MEDB-14. Clinical outcome of pediatric medulloblastoma patients with Li-Fraumeni syndrome. <i>Neuro-Oncology</i> , <b>2022</b> , 24, i107-i107	1	
1	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> , <b>2022</b> , 24, i18-i19	1	