

# Nils Homer

## 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.

15 papers	33,302 citations	13 h-index	16 g-index
16 ext. papers	45,939 ext. citations	9.7 avg, IF	6.64 L-index

#	Paper	IF	Citations
15	The Sequence Alignment/Map format and SAMtools. <i>Bioinformatics</i> , <b>2009</b> , 25, 2078-9	7.2	30805
14	Resolving individuals contributing trace amounts of DNA to highly complex mixtures using high-density SNP genotyping microarrays. <i>PLoS Genetics</i> , <b>2008</b> , 4, e1000167	6	686
13	A survey of sequence alignment algorithms for next-generation sequencing. <i>Briefings in Bioinformatics</i> , <b>2010</b> , 11, 473-83	13.4	632
12	BFAST: an alignment tool for large scale genome resequencing. <i>PLoS ONE</i> , <b>2009</b> , 4, e7767	3.7	394
11	U87MG decoded: the genomic sequence of a cytogenetically aberrant human cancer cell line. <i>PLoS Genetics</i> , <b>2010</b> , 6, e1000832	6	195
10	Common sequence variants on 20q11.22 confer melanoma susceptibility. <i>Nature Genetics</i> , <b>2008</b> , 40, 838-40	34.3	188
9	Identification of the genetic basis for complex disorders by use of pooling-based genomewide single-nucleotide-polymorphism association studies. <i>American Journal of Human Genetics</i> , <b>2007</b> , 80, 126-39	11	129
8	An international effort towards developing standards for best practices in analysis, interpretation and reporting of clinical genome sequencing results in the CLARITY Challenge. <i>Genome Biology</i> , <b>2014</b> , 15, R53	18.3	86
7	Improved variant discovery through local re-alignment of short-read next-generation sequencing data using SRMA. <i>Genome Biology</i> , <b>2010</b> , 11, R99	18.3	57
6	A genome-wide analysis identifies genetic variants in the RELN gene associated with otosclerosis. <i>American Journal of Human Genetics</i> , <b>2009</b> , 84, 328-38	11	52
5	Improving the efficiency of genomic loci capture using oligonucleotide arrays for high throughput resequencing. <i>BMC Genomics</i> , <b>2009</b> , 10, 646	4.5	32
4	Local alignment of two-base encoded DNA sequence. <i>BMC Bioinformatics</i> , <b>2009</b> , 10, 175	3.6	30
3	Multimarker analysis and imputation of multiple platform pooling-based genome-wide association studies. <i>Bioinformatics</i> , <b>2008</b> , 24, 1896-902	7.2	15
2	Local alignment of generalized k-base encoded DNA sequence. <i>BMC Bioinformatics</i> , <b>2010</b> , 11, 347	3.6	1
1	Statistical comparison framework and visualization scheme for ranking-based algorithms in high-throughput genome-wide studies. <i>Journal of Computational Biology</i> , <b>2009</b> , 16, 565-77	1.7	