

# Chris Gunter

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/2093069/publications.pdf>

Version: 2024-02-01

42  
papers

21,648  
citations

623734

14  
h-index

265206

42  
g-index

67  
all docs

67  
docs citations

67  
times ranked

43936  
citing authors

#	ARTICLE	IF	CITATIONS
1	Heritability of social behavioral phenotypes and preliminary associations with autism spectrum disorder risk genes in rhesus macaques: A whole exome sequencing study. Autism Research, 2022, 15, 447-463.	3.8	14
2	Validation of the Social Responsiveness Scale (SRS) to screen for atypical social behaviors in juvenile macaques. PLoS ONE, 2021, 16, e0235946.	2.5	11
3	Mini-Review: Genetic Literacy and Engagement With Genetic Testing for Autism Spectrum Disorder. Frontiers in Genetics, 2021, 12, 693158.	2.3	2
4	Strategic vision for improving human health at The Forefront of Genomics. Nature, 2020, 586, 683-692.	27.8	192
5	Public Discussion Affects Question Asking at Academic Conferences. American Journal of Human Genetics, 2019, 105, 189-197.	6.2	17
6	Ethical principles for the use of human cellular biotechnologies. Nature Biotechnology, 2017, 35, 1050-1058.	17.5	15
7	Diversity in Clinical and Biomedical Research: A Promise Yet to Be Fulfilled. PLoS Medicine, 2015, 12, e1001918.	8.4	424
8	Toward innovative, cost-effective, and systemic solutions to improve outcomes and well-being of military families affected by autism spectrum disorder. Yale Journal of Biology and Medicine, 2015, 88, 73-9.	0.2	6
9	Guidelines for investigating causality of sequence variants in human disease. Nature, 2014, 508, 469-476.	27.8	1,130
10	Science: it's a role model thing. Genome Biology, 2013, 14, 105.	9.6	6
11	What it's like to be an editor at a conference. Genome Biology, 2013, 14, 136.	9.6	0
12	Conference Scene: Accelerating public awareness in the age of personal genetics. Personalized Medicine, 2013, 10, 535-538.	1.5	0
13	Educational Issues and Strategies for Genomic Medicine. , 2013, , 415-421.		1
14	Constructing a 'cancerpaedia'. Nature Reviews Drug Discovery, 2012, 11, 353-353.	46.4	1
15	An integrated encyclopedia of DNA elements in the human genome. Nature, 2012, 489, 57-74.	27.8	15,516
16	Constructing a 'cancerpaedia'. Nature Reviews Genetics, 2012, 13, 300-300.	16.3	1
17	Constructing a 'cancerpaedia'. Nature Reviews Cancer, 2012, 12, 315-315.	28.4	0
18	A modest proposal for an outreach section in scientific publications. Genome Biology, 2012, 13, 168.	9.6	0

#	ARTICLE	IF	CITATIONS
19	Known and unknown. Nature Reviews Genetics, 2011, 12, 521-521.	16.3	3
20	International network of cancer genome projects. Nature, 2010, 464, 993-998.	27.8	2,114
21	A metamorphosis in synthesis. Nature Reviews Genetics, 2010, 11, 5-5.	16.3	0
22	Get in LINE for silencing. Nature Reviews Genetics, 2010, 11, 528-529.	16.3	2
23	A picture worth 1000 Genomes. Nature Reviews Genetics, 2010, 11, 814-814.	16.3	8
24	Education and personalized genomics: deciphering the public's genetic health report. Personalized Medicine, 2009, 6, 681-690.	1.5	5
25	Prepublication data sharing. Nature, 2009, 461, 168-170.	27.8	243
26	Quantitative genetics. Nature, 2008, 456, 719-719.	27.8	6
27	Chipping away at psychiatric disorders. Nature Reviews Genetics, 2008, 9, 654-654.	16.3	2
28	Vive les différences. Nature Reviews Genetics, 2007, 8, S19-S19.	16.3	0
29	Replicating genotype-phenotype associations. Nature, 2007, 447, 655-660.	27.8	1,509
30	Genome labours bear fruit. Nature, 2007, 450, 183-183.	27.8	0
31	She moves in mysterious ways. Nature, 2005, 434, 279-280.	27.8	28
32	The chimpanzee genome. Nature, 2005, 437, 47-47.	27.8	6
33	The X factor. Nature Reviews Molecular Cell Biology, 2005, 6, S6-S6.	37.0	0
34	Human genomics and medicine. Nature, 2004, 429, 439-439.	27.8	9
35	Secrets of a porkier porker. Nature, 2003, 425, 777-777.	27.8	3
36	Stick it in the family album. Nature, 2002, 418, 30-30.	27.8	2

#	ARTICLE	IF	CITATIONS
37	Human biology by proxy. Nature, 2002, 420, 509-509.	27.8	29
38	The molecular genetics of cancer: down the rabbit hole. Human Molecular Genetics, 2001, 10, 655-656.	2.9	5
39	Survey of the Fragile X Syndrome CGG Repeat and the Short-Tandem-Repeat and Single-Nucleotide-Polymorphism Haplotypes in an African American Population. American Journal of Human Genetics, 2000, 66, 480-493.	6.2	45
40	Polymorphism in the FMR1 gene. Human Genetics, 1998, 103, 365-365.	3.8	2
41	Purified Recombinant Fmrp Exhibits Selective RNA Binding as an Intrinsic Property of the Fragile X Mental Retardation Protein. Journal of Biological Chemistry, 1998, 273, 15521-15527.	3.4	148
42	Re-examination of factors associated with expansion of CGG repeats using a single nucleotide polymorphism in FMR1. Human Molecular Genetics, 1998, 7, 1935-1946.	2.9	67