Catherine Bourgain

List of Publications by Year
in descending order

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42 papers	1,477 citations	430874 18 h-index	315739 38 g-index
44 all docs	44 docs citations	44 times ranked	2051 citing authors

#	Article	IF	Citations
1	Genomic susceptibility in practice: The regulatory trajectory of non-rare thrombophilia (NRT) genetic tests in the clinical management of venous thrombo-embolism (VTE). Social Science and Medicine, 2022, 304, 112903.	3.8	7
2	Les fÃ©licitÃ©s du capital en santÃ©. Revue FranÃ§aise De Socio-Ã‰conomie, 2021, nÂ° 26, 127-147.	0.2	1
3	Appraising screening, making risk in/visible. The medical debate over Nonâ€Rare Thrombophilia (NRT) testing before prescribing the pill. Sociology of Health and Illness, 2021, 43, 1627-1642.	2.1	5
4	Imperfect biomarkers for adjuvant chemotherapy in early stage breast cancer with good prognosis. Social Science and Medicine, 2020, 246, 112735.	3.8	5
5	Geneva Statement on Heritable Human Genome Editing: The Need for Course Correction. Trends in Biotechnology, 2020, 38, 351-354.	9.3	37
6	Adopt a moratorium on heritable genome editing. Nature, 2019, 567, 165-168.	27.8	314
7	Searching for secondary findings: considering actionability and preserving the right not to know. European Journal of Human Genetics, 2019, 27, 1481-1484.	2.8	13
8	â€˜It has to become true geneticsâ€™: tumour genetics and the division of diagnostic labour in the clinic. Sociology of Health and Illness, 2019, 41, 643-657.	2.1	7
9	Nouveaux usages de lâ€™ADN en matiÃ¨re pÃ©naleÂ: promesses, savoirs et pratiques. Cahiers Droit Sciences & Technologies, 2019, , 83-93.	0.1	1
10	De la gÃ©nÃ©tique clinique Ã la mÃ©decine gÃ©nomique. Cahiers Droit Sciences & Technologies, 2019, , 15-29.	0.1	2
11	Iron reinforcements in Beauvais and Metz Cathedrals: from bloomery or finery? The use of logistic regression for differentiating smelting processes. Journal of Archaeological Science, 2014, 42, 315-333.	2.4	55
12	Quantitative damage-benefit evaluation of drug effects: major discrepancies between the general population, users and experts. Journal of Psychopharmacology, 2013, 27, 590-599.	4.0	11
13	Genetic and Environmental Factors Influencing the Placental Growth Factor (PGF) Variation in Two Populations. PLoS ONE, 2012, 7, e42537.	2.5	11
14	Association between a polymorphism in the promoter of a glutamate receptor subunit gene (GRIN2A) and alcoholism. Addiction Biology, 2012, 17, 783-785.	2.6	31
15	A damage/benefit evaluation of addictive product use. Addiction, 2012, 107, 441-450.	3.3	24
16	Genetics of VEGF Serum Variation in Human Isolated Populations of Cilento: Importance of VEGF Polymorphisms. PLoS ONE, 2011, 6, e16982.	2.5	68
17	Family History of Thyroid Cancer and the Risk of Differentiated Thyroid Cancer in French Polynesia. Thyroid, 2010, 20, 393-400.	4.5	26
18	A common SNP near BMP2 is associated with severity of the iron burden in HFE p.C282Y homozygous patients: A follow-up study. Blood Cells, Molecules, and Diseases, 2010, 44, 34-37.	1.4	32

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19	Linkage Analysis with Dense SNP Maps in Isolated Populations. Human Heredity, 2009, 68, 87-97.	0.8	16
20	Comprehensive Linkage and Association Analyses Identify Haplotype, Near to the TNFSF15 Gene, Significantly Associated with Spondyloarthritis. PLoS Genetics, 2009, 5, e1000528.	3.5	55
21	Epidemiology, Pathology, and Genetics of Histiocytic Sarcoma in the Bernese Mountain Dog Breed. Journal of Heredity, 2009, 100, S19-S27.	2.4	84
22	A multiple splitting approach to linkage analysis in large pedigrees identifies a linkage to asthma on chromosome 12. Genetic Epidemiology, 2009, 33, 207-216.	1.3	13
23	PEL: an unbiased method for estimating ageâ€dependent genetic disease risk from pedigree data unselected for family history. Genetic Epidemiology, 2009, 33, 379-385.	1.3	17
24	Progressive Retinal Atrophy in the Border Collie: A new XLPRA. BMC Veterinary Research, 2008, 4, 10.	1.9	18
25	Identification and Replication of a Novel Obesity Locus on Chromosome 1q24 in Isolated Populations of Cilento. Diabetes, 2008, 57, 783-790.	0.6	16
26	Facteurs acquis et gÃ©nÃ©tiques de modulation de la pÃ©nÃ©trance de lâ€™hÃ©mochromatose HFE. Bulletin De L'Academie Nationale De Medecine, 2008, 192, 873-881.	0.0	0
27	Inheritance of arterial lesions in renal fibromuscular dysplasia. Journal of Human Hypertension, 2007, 21, 393-400.	2.2	99
28	Common Variants in the BMP2, BMP4, and HJV Genes of the Hepcidin Regulation Pathway Modulate HFE Hemochromatosis Penetrance. American Journal of Human Genetics, 2007, 81, 799-807.	6.2	120
29	A mixture model approach to multiple testing for the genetic analysis of gene expression. BMC Proceedings, 2007, 1, S141.	1.6	5
30	Power of genome-wide association studies in the presence of interacting loci. Genetic Epidemiology, 2007, 31, 748-762.	1.3	17
31	Are genome-wide association studies all that we need to dissect the genetic component of complex human diseases?. European Journal of Human Genetics, 2007, 15, 260-263.	2.8	25
32	New susceptibility locus for hypertension on chromosome 8q by efficient pedigree-breaking in an Italian isolate. Human Molecular Genetics, 2006, 15, 1735-1743.	2.9	39
33	Complex trait mapping in isolated populations: Are specific statistical methods required?. European Journal of Human Genetics, 2005, 13, 698-706.	2.8	28
34	Impact of the diagnosis definition on linkage detection. BMC Genetics, 2005, 6, S140.	2.7	1
35	Detection of susceptibility loci by genome-wide linkage analysis. BMC Genetics, 2005, 6, S18.	2.7	5
36	Modeling the effect of a genetic factor for a complex trait in a simulated population. BMC Genetics, 2005, 6, S87.	2.7	1

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37	Comparing strategies for association mapping in samples with related individuals. BMC Genetics, 2005, 6, S98.	2.7	9
38	Testing for Hardy-Weinberg Equilibrium in Samples With Related Individuals. Genetics, 2004, 168, 2349-2361.	2.9	59
39	Are common disease susceptibility alleles the same in outbred and founder populations?. European Journal of Human Genetics, 2004, 12, 584-590.	2.8	31
40	Novel Case-Control Test in a Founder Population Identifies P-Selectin as an Atopy-Susceptibility Locus. American Journal of Human Genetics, 2003, 73, 612-626.	6.2	148
41	Comparison of family based haplotype methods using intragenic SNPs in candidate genes. European Journal of Human Genetics, 2002, 10, 313-319.	2.8	2
42	Maximum Identity Length Contrast: A Powerful Method For Susceptibility Gene Detection in Isolated Populations. Genetic Epidemiology, 2001, 21, S560-4.	1.3	15