

Anne Cambon-Thomsen

List of Publications by Year in descending order

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Version: 2024-02-01

62
papers

4,444
citations

257450

24
h-index

133252

59
g-index

69
all docs

69
docs citations

69
times ranked

9626
citing authors

#	ARTICLE	IF	CITATIONS
1	Genome-Wide Meta-Analysis Identifies Variants in DSCAM and PDLIM3 That Correlate with Efficacy Outcomes in Metastatic Renal Cell Carcinoma Patients Treated with Sunitinib. <i>Cancers</i> , 2022, 14, 2838.	3.7	1
2	International transfers of personal data for health research following Schrems II: a problem in need of a solution. <i>European Journal of Human Genetics</i> , 2021, 29, 1502-1509.	2.8	12
3	Fostering global data sharing: highlighting the recommendations of the Research Data Alliance COVID-19 working group. <i>Wellcome Open Research</i> , 2020, 5, 267.	1.8	11
4	Fostering global data sharing: highlighting the recommendations of the Research Data Alliance COVID-19 working group. <i>Wellcome Open Research</i> , 2020, 5, 267.	1.8	6
5	FAIRness Literacy: The Achillesâ€™ Heel of Applying FAIR Principles. <i>Data Science Journal</i> , 2020, 19, .	1.3	19
6	Attitudes of French populations towards the disclosure of unsolicited findings in medical genetics. <i>Journal of Health Psychology</i> , 2019, 26, 135910531988662.	2.3	1
7	Is it research or is it clinical? Revisiting an old frontier through the lens of next-generation sequencing technologies. <i>European Journal of Medical Genetics</i> , 2018, 61, 634-641.	1.3	7
8	How to responsibly acknowledge research work in the era of big data and biobanks: ethical aspects of the Bioresource Research Impact Factor (BRIF). <i>Journal of Community Genetics</i> , 2018, 9, 169-176.	1.2	15
9	Academic Valorization of Biobanks. , 2018, , 113-125.		0
10	Including all voices in international data-sharing governance. <i>Human Genomics</i> , 2018, 12, 13.	2.9	50
11	Guidelines for reporting secondary findings of genome sequencing in cancer genes: the SFMPP recommendations. <i>European Journal of Human Genetics</i> , 2018, 26, 1732-1742.	2.8	44
12	French Peopleâ€™s Views on the Appropriateness of Disclosing an Unsolicited Finding in Medical Genetics: A Preliminary Study. <i>Universitas Psychologica</i> , 2018, 17, 1-11.	0.6	1
13	Mechanisms of the Development of Allergy (MeDALL): Introducing novel concepts in allergy phenotypes. <i>Journal of Allergy and Clinical Immunology</i> , 2017, 139, 388-399.	2.9	145
14	Description of the EuroTARGET cohort: A European collaborative project on TArgeted therapy in renal cell cancerâ€™ GENetic- and tumor-related biomarkers for response and toxicity. <i>Urologic Oncology: Seminars and Original Investigations</i> , 2017, 35, 529.e9-529.e16.	1.6	9
15	Critical points for an accurate human genome analysis. <i>Human Mutation</i> , 2017, 38, 912-921.	2.5	5
16	Towards new tools for bioresource use andÂ’sharing. <i>Information Services and Use</i> , 2017, 36, 133-146.	0.2	6
17	Biobankers: Treat the Poison of Invisibility with CoBRA, a Systematic Way of Citing Bioresources in Journal Articles. <i>Biopreservation and Biobanking</i> , 2016, 14, 350-352.	1.0	11
18	Judging health risk as a function of risk factors and type of illness: Do people weight risk factors in a flexible way?. <i>Journal of Health Psychology</i> , 2016, 21, 832-843.	2.3	2

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19	From the arcane to the mundane: engaging French publics in discussing clinical applications of genomic technology. <i>New Genetics and Society</i> , 2016, 35, 1-28.	1.2	4
20	Quand l'anticipation devient plurielle: la complexité des données génomiques à l'épreuve des pratiques professionnelles. <i>Revue Française D'Ethique Appliquée</i> , 2016, N° 2, 19-28.	0.2	4
21	A new classification of HLA-DRB1 alleles based on acid-base properties of the amino acids located at positions 13, 70 and 71: impact on ACPA status or structural progression, and meta-analysis on 1235 patients with rheumatoid from two cohorts (ESPOIR and EAC cohort). <i>RMD Open</i> , 2015, 1, e000099.	3.8	8
22	Genetic markers as a predictive tool based on statistics in medical practice: ethical considerations through the analysis of the use of HLA-B*27 in rheumatology in France. <i>Frontiers in Genetics</i> , 2015, 6, 299.	2.3	1
23	Towards a European consensus for reporting incidental findings during clinical NGS testing. <i>European Journal of Human Genetics</i> , 2015, 23, 1601-1606.	2.8	85
24	Genome Editing and Dialogic Responsibility: "What's in a Name?" <i>American Journal of Bioethics</i> , 2015, 15, 54-57.	0.9	1
25	The impact of European embryonic stem cell patent decisions on research strategies. <i>Nature Biotechnology</i> , 2015, 33, 41-43.	17.5	10
26	Developing a guideline to standardize the citation of bioresources in journal articles (CoBRA). <i>BMC Medicine</i> , 2015, 13, 33.	5.5	43
27	Governing Biobanks Through a European Infrastructure. <i>The International Library of Ethics, Law and Technology</i> , 2015, , 139-151.	0.4	5
28	Beyond public health genomics: proposals from an international working group. <i>European Journal of Public Health</i> , 2014, 24, 877-879.	0.3	22
29	Professionals' Attitudes Regarding Large-Scale Genetic Information Generated Through Next Generation Sequencing in Research. <i>Journal of Empirical Research on Human Research Ethics</i> , 2014, 9, 56-58.	1.3	2
30	Variation in genomic landscape of clear cell renal cell carcinoma across Europe. <i>Nature Communications</i> , 2014, 5, 5135.	12.8	158
31	Association of IL-2RA and IL-2RB genes with erosive status in early rheumatoid arthritis patients (ESPOIR and RMP cohorts). <i>Joint Bone Spine</i> , 2014, 81, 228-234.	1.6	17
32	Systems Medicine Approaches for the Definition of Complex Phenotypes in Chronic Diseases and Ageing. From Concept to Implementation and Policies. <i>Current Pharmaceutical Design</i> , 2014, 20, 5928-5944.	1.9	63
33	Quantifying the use of bioresources for promoting their sharing in scientific research. <i>GigaScience</i> , 2013, 2, 7.	6.4	38
34	Mapping the translational science policy "valley of death". <i>Clinical and Translational Medicine</i> , 2013, 2, 14.	4.0	58
35	Producing "Human Elements Based Medical Technologies" in Biotech Companies: Some Ethical and Organisational Ingredients for Innovative Cooking. <i>Science, Technology and Society</i> , 2013, 18, 93-114.	1.9	0
36	Measuring the Contribution of Tumor Biobanks to Research in Oncology: Surrogate Indicators and Bibliographic Output. <i>Biopreservation and Biobanking</i> , 2013, 11, 235-244.	1.0	29

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37	Genomic Incidental Findings: Reducing the Burden to Be Fair. American Journal of Bioethics, 2013, 13, 52-54.	0.9	13
38	Whole-genome sequencing in health care. European Journal of Human Genetics, 2013, 21, 580-584.	2.8	330
39	Direct-to-consumer health genetic testing services: What commercial strategies for which socio-ethical issues?. Health Sociology Review, 2013, 22, 75-87.	2.8	8
40	Patients' motives for consenting or refusing to participate in a clinical trial in organ transplantation. Clinical Transplantation, 2013, 27, 724-731.	1.6	7
41	Communication of results and disclosure of incidental findings in longitudinal paediatric research. Pediatric Allergy and Immunology, 2013, 24, 389-394.	2.6	17
42	Developing a policy for paediatric biobanks: principles for good practice. European Journal of Human Genetics, 2013, 21, 2-7.	2.8	63
43	Open Data Sharing in the Context of Bioresources. Acta Informatica Medica, 2013, 21, 291.	1.1	15
44	ELSI 2.0 for Genomics and Society. Science, 2012, 336, 673-674.	12.6	39
45	Disclosing Results to Genomic Research Participants: Differences That Matter. American Journal of Bioethics, 2012, 12, 20-22.	0.9	7
46	Understanding the complexity of IgE-related phenotypes from childhood to young adulthood: A Mechanisms of the Development of Allergy (MeDALL) Seminar. Journal of Allergy and Clinical Immunology, 2012, 129, 943-954.e4.	2.9	68
47	Link between traditional cardiovascular risk factors and inflammation in patients with early arthritis: Results from a French Multicenter Cohort. Arthritis Care and Research, 2012, 64, 872-880.	3.4	20
48	The Emergence of Biobanks in the Legal Landscape: Towards a New Model of Governance. Journal of Law and Society, 2012, 39, 113-130.	0.7	14
49	Systems medicine and integrated care to combat chronic noncommunicable diseases. Genome Medicine, 2011, 3, 43.	8.2	181
50	Clarity and claims in variation/mutation databasing. Nature Biotechnology, 2011, 29, 790-792.	17.5	0
51	Comprehensive catalog of European biobanks. Nature Biotechnology, 2011, 29, 795-797.	17.5	83
52	BRIF Workshop Introduction. Nature Precedings, 2011, , .	0.1	0
53	The BRIF (Bioresource Research Impact Factor) as a tool for improving bioresource sharing in biomedical research. Nature Precedings, 2011, , .	0.1	0
54	The role of a bioresource research impact factor as an incentive to share human bioresources. Nature Genetics, 2011, 43, 503-504.	21.4	66

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55	Genetic testing and common disorders in a public health framework: how to assess relevance and possibilities. European Journal of Human Genetics, 2011, 19, S6-S44.	2.8	75
56	Linkage disequilibrium organization of the human KIR superlocus: implications for KIR data analyses. Immunogenetics, 2010, 62, 729-740.	2.4	52
57	Practical guidelines addressing ethical issues pertaining to the curation of human locus-specific variation databases (LSDBs). Human Mutation, 2010, 31, 1179-1184.	2.5	36
58	Italian appeal court: a genetic predisposition to commit murder?. European Journal of Human Genetics, 2010, 18, 519-521.	2.8	39
59	International network of cancer genome projects. Nature, 2010, 464, 993-998.	27.8	2,114
60	2004-2009: l'évolution de la loi de bioéthique en France, quels enjeux, quels défis? Assistance médicale à la procréation, gestation pour autrui, transplantation. Medecine Et Droit, 2010, 2010, 42-48.	0.1	0
61	Prepublication data sharing. Nature, 2009, 461, 168-170.	27.8	243
62	International Transfers of Health Research Data Following Schrems II: A Problem in Need of a Solution. SSRN Electronic Journal, 0, , .	0.4	1