

# Sandrine de Montgolfier

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/4414826/publications.pdf>

Version: 2024-02-01

14  
papers

377  
citations

1040056

9  
h-index

794594

19  
g-index

20  
all docs

20  
docs citations

20  
times ranked

686  
citing authors

#	ARTICLE	IF	CITATIONS
1	Study protocol comparing the ethical, psychological and socio-economic impact of personalised breast cancer screening to that of standard screening in the "My Personal Breast Screening"(MyPeBS) randomised clinical trial. BMC Cancer, 2022, 22, 507.	2.6	32
2	Personalized early detection and prevention of breast cancer: ENVISION consensus statement. Nature Reviews Clinical Oncology, 2020, 17, 687-705.	27.6	178
3	Editorial: When Data Science, Humanities and Social Sciences Meet: Cross-Talks and Insights in Public Health. Frontiers in Public Health, 2020, 8, 41.	2.7	0
4	COVID-19 and ethical considerations: Valuable decision-making tools from the leading medical societies in France. Anaesthesia, Critical Care & Pain Medicine, 2020, 39, 365-366.	1.4	4
5	Familial disclosure by genetic healthcare professionals: a useful but sparingly used legal provision in France. Journal of Medical Ethics, 2019, 45, 811-816.	1.8	8
6	Genetic diseases and information to relatives: practical and ethical issues for professionals after introduction of a legal framework in France. European Journal of Human Genetics, 2018, 26, 786-795.	2.8	22
7	Supporting disclosure of genetic information to family members: professional practice and timelines in cancer genetics. Familial Cancer, 2017, 16, 447-457.	1.9	18
8	Genetic Test Results and Disclosure to Family Members: Qualitative Interviews of Healthcare Professionals' Perceptions of Ethical and Professional Issues in France. Journal of Genetic Counseling, 2016, 25, 483-494.	1.6	20
9	Women's participation in breast cancer screening in France " an ethical approach. BMC Medical Ethics, 2014, 15, 64.	2.4	20
10	Evaluation of biobank constitution and use: multicentre analysis in France and propositions for formalising the activities of research ethics committees. European Journal of Medical Genetics, 2006, 49, 159-167.	1.3	10
11	Communication of pharmacogenetic research results to HIV-infected treated patients: standpoints of professionals and patients. European Journal of Human Genetics, 2005, 13, 1055-1062.	2.8	25
12	Enjeux Éthiques des collections d'échantillons humains dans le cadre de la recherche. Droit, Deontologie Et Soins, 2004, 4, 148-164.	0.0	3
13	Study of the involvement of research ethics committees in the constitution and use of biobanks in France. Pharmacogenetics and Genomics, 2004, 14, 195-198.	5.7	7
14	Ethical reflections on pharmacogenetics and DNA banking in a cohort of HIV-infected patients. Pharmacogenetics and Genomics, 2002, 12, 667-675.	5.7	11