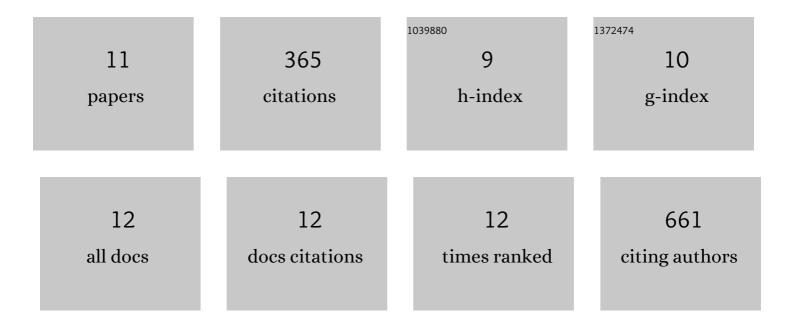
Pauline McCormack

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/2477147/publications.pdf Version: 2024-02-01



#	Article	IF	CITATIONS
1	International Charter of principles for sharing bio-specimens and data. European Journal of Human Genetics, 2015, 23, 721-728.	1.4	112
2	â€~You should at least ask'. The expectations, hopes and fears of rare disease patients on large-scale data and biomaterial sharing for genomics research. European Journal of Human Genetics, 2016, 24, 1403-1408.	1.4	70
3	Improving the informed consent process in international collaborative rare disease research: effective consent for effective research. European Journal of Human Genetics, 2016, 24, 1248-1254.	1.4	47
4	DISPUTING THE ETHICS OF RESEARCH: THE CHALLENGE FROM BIOETHICS AND PATIENT ACTIVISM TO THE INTERPRETATION OF THE DECLARATION OF HELSINKI IN CLINICAL TRIALS. Bioethics, 2013, 27, 243-250.	0.7	29
5	Precaution, governance and the failure of medical implants: the ASR(TM) hip in the UK. Life Sciences, Society and Policy, 2014, 10, 19.	3.1	24
6	Therapeutic Misconception: Hope, Trust and Misconception in Paediatric Research. Health Care Analysis, 2014, 22, 3-21.	1.4	24
7	The risks of therapeutic misconception and individual patient (n=1) "trials―in rare diseases such as Duchenne dystrophy. Neuromuscular Disorders, 2011, 21, 13-15.	0.3	19
8	Delivering genomic medicine in the UnitedÂKingdom National Health Service: a systematic review and narrative synthesis. Genetics in Medicine, 2019, 21, 2667-2675.	1.1	17
9	Guidance in Social and Ethical Issues Related to Clinical, Diagnostic Care and Novel Therapies for Hereditary Neuromuscular Rare Diseases: "Translating" the Translational. PLOS Currents, 2013, 5, .	1.4	15
10	Setting up strategies: patient inclusion in biobank and genomics research in Europe. Orphanet Journal of Rare Diseases, 2014, 9, P2.	1.2	8
11	New Recommendation on Biological Materials Could Hamper Muscular Dystrophy Research. PLOS	1.4	0