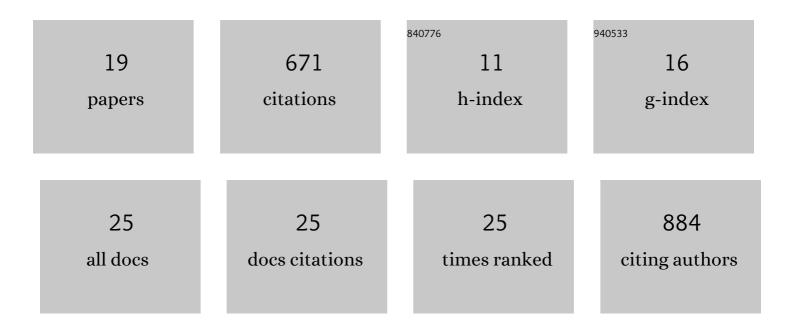
Susan E Mathers

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/1650725/publications.pdf Version: 2024-02-01



#	Article	IF	CITATIONS
1	MiNDAUS partnership: a roadmap for the cure and management of motor Neurone disease. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2022, 23, 321-328.	1.7	4
2	Functional characterisation of the amyotrophic lateral sclerosis risk locus GPX3/TNIP1. Genome Medicine, 2022, 14, 7.	8.2	12
3	Genome-wide study of DNA methylation shows alterations in metabolic, inflammatory, and cholesterol pathways in ALS. Science Translational Medicine, 2022, 14, eabj0264.	12.4	38
4	Public Policy in MND Care: The Australian Perspective. , 2021, , 29-49.		4
5	Common and rare variant association analyses in amyotrophic lateral sclerosis identify 15 risk loci with distinct genetic architectures and neuron-specific biology. Nature Genetics, 2021, 53, 1636-1648.	21.4	223
6	Genome-wide Meta-analysis Finds the ACSL5-ZDHHC6 Locus Is Associated with ALS and Links Weight Loss to the Disease Genetics. Cell Reports, 2020, 33, 108323.	6.4	41
7	Serial assessment of iron in the motor cortex in limb-onset amyotrophic lateral sclerosis using quantitative susceptibility mapping. Quantitative Imaging in Medicine and Surgery, 2020, 10, 1465-1476.	2.0	17
8	Phase 2 randomized placebo controlled double blind study to assess the efficacy and safety of tecfidera in patients with amyotrophic lateral sclerosis (TEALS Study). Medicine (United States), 2020, 99, e18904.	1.0	23
9	A self-care, problem-solving and mindfulness intervention for informal caregivers of people with motor neurone disease: A pilot study. Palliative Medicine, 2018, 32, 726-732.	3.1	7
10	Evaluating the Utility of a Structured Clinical Protocol for Reducing the Impact of Behavioural and Psychological Symptoms of Dementia in Progressive Neurological Diseases: A Pilot Study. Behavioural Neurology, 2018, 2018, 1-9.	2.1	0
11	Factors to consider for motor neurone disease carer intervention research: A narrative literature review. Palliative and Supportive Care, 2017, 15, 600-608.	1.0	13
12	Motor neurone disease: progress and challenges. Medical Journal of Australia, 2017, 206, 357-362.	1.7	28
13	Identification and outcomes of clinical phenotypes in amyotrophic lateral sclerosis/motor neuron disease: Australian National Motor Neuron Disease observational cohort. BMJ Open, 2016, 6, e012054.	1.9	48
14	Identifying who will benefit from non-invasive ventilation in amyotrophic lateral sclerosis/motor neurone disease in a clinical cohort. Journal of Neurology, Neurosurgery and Psychiatry, 2016, 87, 280-286.	1.9	100
15	End of Life Care in Progressive Neurological Disease: Australia. , 2013, , 205-212.		0
16	Home or Residential Care? The Role of Behavioral and Psychosocial Factors in Determining Discharge Outcomes for Inpatients with Huntington's Disease. Journal of Huntington's Disease, 2012, 1, 187-193.	1.9	9
17	Clinical phenotypes and natural progression for motor neuron disease: Analysis from an Australian database. Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders, 2009, 10, 79-84.	2.1	47
18	Multidisciplinary care for adults with amyotrophic lateral sclerosis or motor neuron disease. The Cochrane Library, 2009, , CD007425.	2.8	45

#	Article	IF	CITATIONS
19	Faecal incontinence. International Disability Studies, 1988, 10, 164-168.	0.4	Ο