

Jill I Cameron

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/9173616/publications.pdf>

Version: 2024-02-01

92
papers

5,115
citations

134610

34
h-index

104191

69
g-index

93
all docs

93
docs citations

93
times ranked

6625
citing authors

#	ARTICLE	IF	CITATIONS
1	Conceptualization of frailty in rehabilitation interventions with adults: a scoping review. <i>Disability and Rehabilitation</i> , 2023, 45, 117-153.	0.9	2
2	A qualitative study exploring family caregivers' support needs in the context of medical assistance in dying. <i>Palliative and Supportive Care</i> , 2023, 21, 254-260.	0.6	5
3	A grounded theory study to identify caregiving phases and support needs across the Alzheimer's disease trajectory. <i>Disability and Rehabilitation</i> , 2022, 44, 1050-1059.	0.9	21
4	Perceptions of Digital Technology Experiences and Development Among Family Caregivers and Technology Researchers: Qualitative Study. <i>JMIR Formative Research</i> , 2022, 6, e19967.	0.7	5
5	Brothers and sisters sharing in the care of a parent with dementia. <i>Dementia</i> , 2022, 21, 765-780.	1.0	3
6	Community-based exercise programs incorporating healthcare-community partnerships to improve function post-stroke: feasibility of a 2-group randomized controlled trial. <i>Pilot and Feasibility Studies</i> , 2022, 8, 88.	0.5	4
7	Exploring the poststroke experiences and unmet needs of South Asian communities in high-income countries: a scoping review protocol. <i>BMJ Open</i> , 2022, 12, e059017.	0.8	4
8	Daughters' experiences of shared caregiving to a parent with dementia. <i>Scandinavian Journal of Caring Sciences</i> , 2021, 35, 853-859.	1.0	4
9	Exploring the Emergence of Self-Directed Home Care in Ontario: A Qualitative Case Study on Home Care Services. <i>Home Health Care Management and Practice</i> , 2021, 33, 28-36.	0.4	1
10	Key informants' perspectives on implementing caregiver programs in an organized system of stroke care. <i>Disability and Rehabilitation</i> , 2021, 43, 1145-1152.	0.9	5
11	A qualitative exploration of how gender and relationship shape family caregivers' experiences across the Alzheimer's disease trajectory. <i>Dementia</i> , 2021, 20, 2851-2866.	1.0	13
12	Caregivers' decision-making for health service utilisation across the Alzheimer's disease trajectory. <i>Health and Social Care in the Community</i> , 2021, , .	0.7	8
13	Impacts of the COVID-19 pandemic on health, financial worries, and perceived organizational support among people living with disabilities in Canada. <i>Disability and Health Journal</i> , 2021, 14, 101161.	1.6	23
14	Family caregiving research: Reflecting on the past to inform the future. <i>Journal of Spinal Cord Medicine</i> , 2021, 44, S19-S22.	0.7	2
15	A Qualitative Study of Caregiving to Older Adults with Depression: "Not the Person I Used to Know". <i>Canadian Journal of Community Mental Health</i> , 2021, 40, 55-71.	0.1	0
16	Facilitators and barriers to supporting individuals with spinal cord injury in the community: experiences of family caregivers and care recipients. <i>Disability and Rehabilitation</i> , 2020, 42, 1844-1854.	0.9	17
17	Experiences of people with stroke and multiple sclerosis and caregivers of a community exercise programme involving a healthcare-recreation partnership. <i>Disability and Rehabilitation</i> , 2020, 42, 1220-1226.	0.9	13
18	"Caregiving is a full-time job" impacting stroke caregivers' health and well-being: A qualitative meta-synthesis. <i>Health and Social Care in the Community</i> , 2020, 28, 325-340.	0.7	51

#	ARTICLE	IF	CITATIONS
19	Longitudinal mixed methods study assessing caregivers of seniors across diverse populations: research protocol. BMC Health Services Research, 2020, 20, 371.	0.9	0
20	Sex and gender differences in technology needs and preferences among informal caregivers of persons with dementia. BMC Geriatrics, 2020, 20, 176.	1.1	12
21	A Qualitative study assessing organisational readiness to implement caregiver support programmes in Ontario, Canada. BMJ Open, 2020, 10, e035559.	0.8	3
22	Support needs and health-related quality of life of family caregivers of patients requiring prolonged mechanical ventilation and admission to a specialised weaning centre: A qualitative longitudinal interview study. Intensive and Critical Care Nursing, 2020, 58, 102808.	1.4	18
23	Canadian Stroke Best Practice Recommendations: Rehabilitation, Recovery, and Community Participation following Stroke. Part One: Rehabilitation and Recovery Following Stroke; 6th Edition Update 2019. International Journal of Stroke, 2020, 15, 763-788.	2.9	194
24	Canadian Stroke Best Practice Recommendations: Rehabilitation, Recovery, and Community Participation following Stroke. Part Two: Transitions and Community Participation Following Stroke. International Journal of Stroke, 2020, 15, 789-806.	2.9	75
25	Sex and gender differences in caregiving burden experienced by family caregivers of persons with dementia: A systematic review. PLoS ONE, 2020, 15, e0231848.	1.1	109
26	Priority Outcomes in Critically Ill Children: A Patient and Parent Perspective. American Journal of Critical Care, 2020, 29, e94-e103.	0.8	16
27	Implementing Caregiver Support Programs in a Regional Stroke System. Stroke, 2019, 50, 3585-3591.	1.0	8
28	Towards a universal model of family centered care: a scoping review. BMC Health Services Research, 2019, 19, 564.	0.9	205
29	Re-building relationships after a spinal cord injury: experiences of family caregivers and care recipients. BMC Neurology, 2019, 19, 117.	0.8	29
30	Determinants of Depressive Symptoms at 1 Year Following ICU Discharge in Survivors of 7 Days of Mechanical Ventilation. Chest, 2019, 156, 466-476.	0.4	14
31	Frailty in the context of rehabilitation interventions for adults: protocol for a scoping review. BMJ Open, 2019, 9, e024838.	0.8	4
32	Identifying Required Skills to Enhance Family Caregiver Competency in Caring for Individuals With Spinal Cord Injury Living in the Community. Topics in Spinal Cord Injury Rehabilitation, 2019, 25, 290-302.	0.8	7
33	Comment: Improving the costing of informal caregiving in neurologic populations. Neurology, 2018, 90, 35-35.	1.5	0
34	Quantitative Evaluation of Muscle Function, Gait, and Postural Control in People Experiencing Critical Illness After Discharge From the Intensive Care Unit. Physical Therapy, 2018, 98, 8-15.	1.1	14
35	How does peer similarity influence adult children caregivers' perceptions of support from peers? A mixed-method study. Ageing and Society, 2018, 38, 2280-2303.	1.2	3
36	A longitudinal view of factors that influence the emotional well-being of family caregivers to individuals with heart failure. Aging and Mental Health, 2017, 21, 844-850.	1.5	29

#	ARTICLE	IF	CITATIONS
37	Web-based health interventions for family caregivers of elderly individuals: A Scoping Review. <i>International Journal of Medical Informatics</i> , 2017, 103, 109-138.	1.6	99
38	Sustaining care for a parent with dementia: an indefinite and intertwined process. <i>International Journal of Qualitative Studies on Health and Well-being</i> , 2017, 12, 1389578.	0.6	14
39	Mechanisms of Chronic Muscle Wasting and Dysfunction after an Intensive Care Unit Stay. A Pilot Study. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2016, 194, 821-830.	2.5	176
40	One-Year Outcomes in Caregivers of Critically Ill Patients. <i>New England Journal of Medicine</i> , 2016, 374, 1831-1841.	13.9	301
41	Adult children caregivers' experiences with online and in-person peer support. <i>Computers in Human Behavior</i> , 2016, 65, 14-22.	5.1	7
42	Canadian Stroke Best Practice Recommendations: Managing transitions of care following Stroke, Guidelines Update 2016. <i>International Journal of Stroke</i> , 2016, 11, 807-822.	2.9	86
43	Characterising community exercise programmes delivered by fitness instructors for people with neurological conditions: a scoping review. <i>Health and Social Care in the Community</i> , 2016, 24, e101-e116.	0.7	32
44	The RECOVER Program: Disability Risk Groups and 1-Year Outcome after 7 or More Days of Mechanical Ventilation. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2016, 194, 831-844.	2.5	272
45	Validation of the Evidence-Based Practice Confidence (EPIC) Scale With Occupational Therapists. <i>American Journal of Occupational Therapy</i> , 2016, 70, 7002280010p1-7002280010p9.	0.1	13
46	Roles and Coping Strategies of Sons Caring for a Parent With Dementia. <i>American Journal of Occupational Therapy</i> , 2016, 70, 7001260020p1-7001260020p9.	0.1	19
47	Factors that contribute to adult children caregivers' well-being: a scoping review. <i>Health and Social Care in the Community</i> , 2015, 23, 449-466.	0.7	35
48	The experiences and needs of Chinese-Canadian stroke survivors and family caregivers as they re-integrate into the community. <i>Health and Social Care in the Community</i> , 2015, 23, 523-531.	0.7	25
49	Co-Designing Ambient Assisted Living (AAL) Environments: Unravelling the Situated Context of Informal Dementia Care. <i>BioMed Research International</i> , 2015, 2015, 1-12.	0.9	30
50	Adult daughters providing post-stroke care to a parent: a qualitative study of the impact that role overload has on lifestyle, participation and family relationships. <i>Clinical Rehabilitation</i> , 2015, 29, 592-600.	1.0	36
51	Changing support needs of survivors of complex critical illness and their family caregivers across the care continuum: A qualitative pilot study of Towards RECOVER. <i>Journal of Critical Care</i> , 2015, 30, 242-249.	1.0	71
52	Getting on with the rest of your life following stroke: a randomized trial of a complex intervention aimed at enhancing life participation post stroke. <i>Clinical Rehabilitation</i> , 2015, 29, 1198-1211.	1.0	67
53	Restricted Participation in Stroke Caregivers: Who's at Risk?. <i>Archives of Physical Medicine and Rehabilitation</i> , 2015, 96, 1284-1290.	0.5	26
54	Studying long-term caregiver health outcomes with methodologic rigor. <i>Neurology</i> , 2015, 84, 1292-1293.	1.5	2

#	ARTICLE	IF	CITATIONS
55	Factors Influencing Physical Therapists' Use of Standardized Measures of Walking Capacity Poststroke Across the Care Continuum. <i>Physical Therapy</i> , 2015, 95, 1507-1517.	1.1	38
56	A feasibility and pilot randomized controlled trial of the "Timing it Right Stroke Family Support Program". <i>Clinical Rehabilitation</i> , 2015, 29, 1129-1140.	1.0	33
57	Stroke survivors' , caregivers' , and health care professionals' perspectives on the weekend pass to facilitate transition home. <i>Journal of Rehabilitation Medicine</i> , 2014, 46, 858-863.	0.8	12
58	Daughters Providing Poststroke Care. <i>Qualitative Health Research</i> , 2014, 24, 1527-1539.	1.0	4
59	Randomized clinical trial of the timing it right stroke family support program: research protocol. <i>BMC Health Services Research</i> , 2014, 14, 18.	0.9	27
60	What Makes Family Caregivers Happy During the First 2 Years Post Stroke?. <i>Stroke</i> , 2014, 45, 1084-1089.	1.0	43
61	Disability after Critical Illness. <i>New England Journal of Medicine</i> , 2013, 369, 1367-1369.	13.9	30
62	Intensive Care Unit-acquired Weakness. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2013, 187, 238-246.	2.5	193
63	Stroke family caregivers' support needs change across the care continuum: a qualitative study using the timing it right framework. <i>Disability and Rehabilitation</i> , 2013, 35, 315-324.	0.9	142
64	Lung-Injured Patients Do Not Need a Specialized Rehabilitation Program: ICUAW as a Case Study. <i>Seminars in Respiratory and Critical Care Medicine</i> , 2013, 34, 522-528.	0.8	2
65	Stroke Survivor Depressive Symptoms Are Associated With Family Caregiver Depression During the First 2 Years Poststroke. <i>Stroke</i> , 2011, 42, 302-306.	1.0	83
66	Determining the Factors Associated With Health Research Participation. <i>Journal of Ambulatory Care Management</i> , 2010, 33, 272-284.	0.5	2
67	Exploring the Feasibility of Videoconference Delivery of a Self-Management Program to Rural Participants with Stroke. <i>Telemedicine Journal and E-Health</i> , 2009, 15, 646-654.	1.6	39
68	A Longitudinal View of Apathy and Its Impact After Stroke. <i>Stroke</i> , 2009, 40, 3299-3307.	1.0	176
69	Education and support needs during recovery in acute respiratory distress syndrome survivors. <i>Critical Care</i> , 2009, 13, R153.	2.5	43
70	Cultural syndromes and age moderate the emotional impact of illness intrusiveness in rheumatoid arthritis. <i>Rehabilitation Psychology</i> , 2009, 54, 33-44.	0.7	16
71	"Timing It Right": A conceptual framework for addressing the support needs of family caregivers to stroke survivors from the hospital to the home. <i>Patient Education and Counseling</i> , 2008, 70, 305-314.	1.0	230
72	Differences in the Experiences and Support Needs of Family Caregivers to Stroke Survivors: Does Age Matter?. <i>Topics in Stroke Rehabilitation</i> , 2008, 15, 593-601.	1.0	22

#	ARTICLE	IF	CITATIONS
73	Factor Structure and Reliability of the Brain Impairment Behavior Scale. <i>Journal of Neuroscience Nursing</i> , 2008, 40, 40-47.	0.7	16
74	Optimizing Stroke Systems of Care by Enhancing Transitions Across Care Environments. <i>Stroke</i> , 2008, 39, 2637-2643.	1.0	92
75	One-Year Outcomes and Health Care Utilization in Survivors of Severe Acute Respiratory Syndrome. <i>Archives of Internal Medicine</i> , 2007, 167, 1312.	4.3	244
76	Program design features that can improve participation in health education interventions. <i>BMC Medical Research Methodology</i> , 2007, 7, 47.	1.4	18
77	Development and implementation of the Ontario Stroke System: the use of evidence. <i>International Journal of Integrated Care</i> , 2007, 7, e30.	0.1	7
78	Stroke Survivorsâ€™ Behavioral and Psychologic Symptoms Are Associated With Informal Caregiversâ€™ Experiences of Depression. <i>Archives of Physical Medicine and Rehabilitation</i> , 2006, 87, 177-183.	0.5	87
79	Well-being in informal caregivers of survivors of acute respiratory distress syndrome*. <i>Critical Care Medicine</i> , 2006, 34, 81-86.	0.4	123
80	Technology in the Lives of Women Who Live With Memory Impairment as a Result of a Traumatic Brain Injury. <i>Assistive Technology</i> , 2006, 18, 170-180.	1.2	9
81	A tailored intervention to promote breast cancer screening among South Asian immigrant women. <i>Social Science and Medicine</i> , 2005, 60, 575-586.	1.8	72
82	A brief problem-solving intervention for family caregivers to individuals with advanced cancer. <i>Journal of Psychosomatic Research</i> , 2004, 57, 137-143.	1.2	62
83	Medical decision-making and information needs in end-stage renal disease patients. <i>General Hospital Psychiatry</i> , 2003, 25, 324-331.	1.2	54
84	Physicians and Pap Testing in the Chinese and Vietnamese Communities in Toronto. <i>Journal of Health Care for the Poor and Underserved</i> , 2003, 14, 489-502.	0.4	7
85	THE EXPERIENCE OF PROVIDING CARE TO RELATIVES WITH CHRONIC MENTAL ILLNESS. <i>Journal of Nervous and Mental Disease</i> , 2002, 190, 108-114.	0.5	117
86	Promotion of Healthy Eating: Among New Immigrant Women in Ontario. <i>Canadian Journal of Dietetic Practice and Research</i> , 2002, 63, 125-129.	0.5	13
87	Lifestyle interference and emotional distress in family caregivers of advanced cancer patients. <i>Cancer</i> , 2002, 94, 521-527.	2.0	275
88	Rural Physicians' Perspectives on Cervical and Breast Cancer Screening: A Gender-Based Analysis. <i>Journal of Women's Health and Gender-Based Medicine</i> , 2001, 10, 201-208.	1.7	40
89	Information Needs and Decisional Preferences among Women with Ovarian Cancer. <i>Gynecologic Oncology</i> , 2000, 77, 357-361.	0.6	141
90	Sex Inequality in Kidney Transplantation Rates. <i>Archives of Internal Medicine</i> , 2000, 160, 2349.	4.3	78

#	ARTICLE	IF	CITATIONS
91	Differences in quality of life across renal replacement therapies: A meta-analytic comparison. American Journal of Kidney Diseases, 2000, 35, 629-637.	2.1	288
92	Behavioral Medicine and Women. Psychosomatic Medicine, 1998, 60, 785-786.	1.3	5