

Erika Pietrolongo

List of Publications by Year in descending order

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Version: 2024-02-01

46
papers

1,334
citations

331259

21
h-index

344852

36
g-index

46
all docs

46
docs citations

46
times ranked

1836
citing authors

#	ARTICLE	IF	CITATIONS
1	Predictors and dynamics of postpartum relapses in women with multiple sclerosis. <i>Multiple Sclerosis Journal</i> , 2014, 20, 739-746.	1.4	148
2	Sex as a determinant of relapse incidence and progressive course of multiple sclerosis. <i>Brain</i> , 2013, 136, 3609-3617.	3.7	140
3	Comparison of Switch to Fingolimod or Interferon Beta/Glatiramer Acetate in Active Multiple Sclerosis. <i>JAMA Neurology</i> , 2015, 72, 405.	4.5	100
4	The brief international cognitive assessment for multiple sclerosis (BICAMS): normative values with gender, age and education corrections in the Italian population. <i>BMC Neurology</i> , 2014, 14, 171.	0.8	99
5	Unmet Needs of People with Severe Multiple Sclerosis and Their Carers: Qualitative Findings for a Home-Based Intervention. <i>PLoS ONE</i> , 2014, 9, e109679.	1.1	67
6	Role Preferences of People with Multiple Sclerosis: Image-Revised, Computerized Self-Administered Version of the Control Preference Scale. <i>PLoS ONE</i> , 2013, 8, e66127.	1.1	52
7	Efficacy of fingolimod and interferon beta-1b on cognitive, MRI, and clinical outcomes in relapsingâ€“remitting multiple sclerosis: an 18-month, open-label, rater-blinded, randomised, multicentre study (the GOLDEN study). <i>Journal of Neurology</i> , 2017, 264, 2436-2449.	1.8	44
8	Decision-Making in Multiple Sclerosis Consultations in Italy: Third Observer and Patient Assessments. <i>PLoS ONE</i> , 2013, 8, e60721.	1.1	44
9	Illness Perception and Well-Being Among Persons with Multiple Sclerosis and Their Caregivers. <i>Journal of Clinical Psychology in Medical Settings</i> , 2016, 23, 33-52.	0.8	39
10	The Kurtzke EDSS rank stability increases 4â€“years after the onset of multiple sclerosis: results from the MSBase Registry. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2012, 83, 305-310.	0.9	37
11	The coexistence of well- and ill-being in persons with multiple sclerosis, their caregivers and health professionals. <i>Journal of the Neurological Sciences</i> , 2014, 337, 67-73.	0.3	37
12	Comparative effectiveness of glatiramer acetate and interferon beta formulations in relapsingâ€“remitting multiple sclerosis. <i>Multiple Sclerosis Journal</i> , 2015, 21, 1159-1171.	1.4	36
13	Increasing age at disability milestones among MS patients in the MSBase Registry. <i>Journal of the Neurological Sciences</i> , 2012, 318, 94-99.	0.3	35
14	Country, Sex, EDSS Change and Therapy Choice Independently Predict Treatment Discontinuation in Multiple Sclerosis and Clinically Isolated Syndrome. <i>PLoS ONE</i> , 2012, 7, e38661.	1.1	35
15	Prognostic indicators in pediatric clinically isolated syndrome. <i>Annals of Neurology</i> , 2017, 81, 729-739.	2.8	34
16	A comparison of the brief international cognitive assessment for multiple sclerosis and the brief repeatable battery in multiple sclerosis patients. <i>BMC Neurology</i> , 2015, 15, 204.	0.8	31
17	Patient Expression of Emotions and Neurologist Responses in First Multiple Sclerosis Consultations. <i>PLoS ONE</i> , 2015, 10, e0127734.	1.1	31
18	Beyond Disease: Happiness, Goals, and Meanings among Persons with Multiple Sclerosis and Their Caregivers. <i>Frontiers in Psychology</i> , 2017, 8, 2216.	1.1	30

#	ARTICLE	IF	CITATIONS
19	Risk-benefit considerations in the treatment of relapsing-remitting multiple sclerosis. <i>Neuropsychiatric Disease and Treatment</i> , 2013, 9, 893.	1.0	28
20	Home-based palliative approach for people with severe multiple sclerosis and their carers: study protocol for a randomized controlled trial. <i>Trials</i> , 2015, 16, 184.	0.7	28
21	Development and validation of a patient self-assessed questionnaire on satisfaction with communication of the multiple sclerosis diagnosis. <i>Multiple Sclerosis Journal</i> , 2010, 16, 1237-1247.	1.4	27
22	Conversion to Secondary Progressive Multiple Sclerosis: Patient Awareness and Needs. Results From an Online Survey in Italy and Germany. <i>Frontiers in Neurology</i> , 2019, 10, 916.	1.1	21
23	Alemtuzumab treatment of multiple sclerosis in real-world clinical practice: A report from a single Italian center. <i>Multiple Sclerosis and Related Disorders</i> , 2020, 38, 101504.	0.9	19
24	Effect of the disclosure of MS diagnosis on anxiety, mood and quality of life of patients: a prospective study. <i>International Journal of Clinical Practice</i> , 2012, 66, 504-514.	0.8	17
25	Illness perceptions and psychological adjustment among persons with multiple sclerosis: the mediating role of coping strategies and social support. <i>Disability and Rehabilitation</i> , 2020, 42, 3780-3792.	0.9	17
26	The caring experience in multiple sclerosis: Caregiving tasks, coping strategies and psychological well-being. <i>Health and Social Care in the Community</i> , 2020, 28, 236-246.	0.7	17
27	The Contribution of Illness Beliefs, Coping Strategies, and Social Support to Perceived Physical Health and Fatigue in Multiple Sclerosis. <i>Journal of Clinical Psychology in Medical Settings</i> , 2021, 28, 149-160.	0.8	17
28	Managing the transition (ManTra): a resource for persons with secondary progressive multiple sclerosis and their health professionals: protocol for a mixed-methods study in Italy. <i>BMJ Open</i> , 2017, 7, e017254.	0.8	16
29	Implementation of the "Sapere Migliora"™ information aid for newly diagnosed people with multiple sclerosis in routine clinical practice: a late-phase controlled trial. <i>Multiple Sclerosis Journal</i> , 2014, 20, 1234-1243.	1.4	14
30	Experience of an information aid for newly diagnosed multiple sclerosis patients: a qualitative study on the SIMS trial. <i>Health Expectations</i> , 2014, 17, 36-48.	1.1	12
31	Simultaneous early-onset severe autoimmune hemolytic anemia and albuminuria during alemtuzumab treatment for multiple sclerosis. <i>Multiple Sclerosis Journal</i> , 2018, 24, 813-815.	1.4	11
32	eMSQOL-29: Prospective validation of the abbreviated, electronic version of MSQOL-54. <i>Multiple Sclerosis Journal</i> , 2019, 25, 856-866.	1.4	11
33	Individualized quality of life of severely affected multiple sclerosis patients: practicability and value in comparison with standard inventories. <i>Quality of Life Research</i> , 2016, 25, 2755-2763.	1.5	10
34	Assessing subjective quality of life domains after multiple sclerosis diagnosis disclosure. <i>Health Expectations</i> , 2016, 19, 437-447.	1.1	10
35	Conversion to secondary progressive multiple sclerosis: Multistakeholder experiences and needs in Italy. <i>PLoS ONE</i> , 2020, 15, e0228587.	1.1	9
36	Assessing measurement invariance of MSQOL-54 across Italian and English versions. <i>Quality of Life Research</i> , 2020, 29, 783-791.	1.5	4

#	ARTICLE	IF	CITATIONS
37	The minimal neuropsychological assessment of MS patients (MACFIMS): normative data of the Italian population. <i>Neurological Sciences</i> , 2020, 41, 1489-1496.	0.9	3
38	Viability of a MSQOL-54 general health-related quality of life score using bifactor model. <i>Health and Quality of Life Outcomes</i> , 2021, 19, 224.	1.0	2
39	Risk attitude and personality in people with multiple sclerosis facing the choice of different disease-modifying therapy scenarios. <i>Journal of the Neurological Sciences</i> , 2020, 417, 117064.	0.3	1
40	Data of safety in a single-center alemtuzumab treated population. <i>Data in Brief</i> , 2020, 29, 105341.	0.5	1
41	Role preferences in treatment decisions of MS patients in Italy. <i>Journal of the Neurological Sciences</i> , 2015, 357, e316.	0.3	0
42	Italian validation of the caregiving tasks in multiple sclerosis scale (CTiMSS). <i>Neurological Sciences</i> , 2020, 41, 1881-1889.	0.9	0
43	Title is missing!. , 2020, 15, e0228587.		0
44	Title is missing!. , 2020, 15, e0228587.		0
45	Title is missing!. , 2020, 15, e0228587.		0
46	Title is missing!. , 2020, 15, e0228587.		0