Francis Meire FÃ¡vero

List of Publications by Year
in descending order

58 papers	447 citations	759233 12 h-index	794594 19 g-index
59 all docs	59 docs citations	59 times ranked	636 citing authors

#	Article	IF	Citations
1	Virtual reality in multiple sclerosis â€“ A systematic review. Multiple Sclerosis and Related Disorders, 2016, 8, 107-112.	2.0	83
2	Effects of Exercises on Bell's Palsy. Otology and Neurotology, 2008, 29, 557-560.	1.3	39
3	Achievement of Virtual and Real Objects Using a Short-Term Motor Learning Protocol in People with Duchenne Muscular Dystrophy: A Crossover Randomized Controlled Trial. Games for Health Journal, 2018, 7, 107-115.	2.0	18
4	The relevance of trunk evaluation in Duchenne muscular dystrophy: the segmental assessment of trunk control. Arquivos De Neuro-Psiquiatria, 2016, 74, 791-795.	0.8	17
5	Motor learning from virtual reality to natural environments in individuals with Duchenne muscular dystrophy. Disability and Rehabilitation: Assistive Technology, 2019, 14, 12-20.	2.2	17
6	Relationship between muscle strength and motor function in Duchenne muscular dystrophy. Arquivos De Neuro-Psiquiatria, 2016, 74, 530-535.	0.8	16
7	Perfil clÃ¬nico e funcional dos pacientes com Distrofia Muscular de Duchenne assistidos na AssociaÃ§Ã£o Brasileira de Distrofia Muscular (ABDIM). Revista Neurociencias, 2006, 14, 15-22.	0.0	16
8	Motor learning through virtual reality in cerebral palsy - a literature review. Medical Express, 2014, 1, .	0.2	15
9	Autonomic Modulation in Duchenne Muscular Dystrophy during a Computer Task: A Prospective Control Trial. PLoS ONE, 2017, 12, e0169633.	2.5	15
10	Matching pairs difficulty in children with spinal muscular atrophy type I. Neuromuscular Disorders, 2017, 27, 419-427.	0.6	14
11	Computer task performance by subjects with Duchenne muscular dystrophy. Neuropsychiatric Disease and Treatment, 2016, 12, 41.	2.2	13
12	Pain characterization in Duchenne muscular dystrophy. Arquivos De Neuro-Psiquiatria, 2016, 74, 767-774.	0.8	13
13	Reliability, validity and description of timed performance of the Jebsenâ€“Taylor Test in patients with muscular dystrophies. Brazilian Journal of Physical Therapy, 2018, 22, 190-197.	2.5	12
14	Efficacy of different interaction devices using non-immersive virtual tasks in individuals with Amyotrophic Lateral Sclerosis: a cross-sectional randomized trial. BMC Neurology, 2018, 18, 209.	1.8	12
15	Functional performance and muscular strength in symptomatic female carriers of Duchenne muscular dystrophy. Arquivos De Neuro-Psiquiatria, 2020, 78, 143-148.	0.8	12
16	Mortality rates due to amyotrophic lateral sclerosis in SÃ£o Paulo City from 2002 to 2006. Arquivos De Neuro-Psiquiatria, 2011, 69, 861-866.	0.8	11
17	The clinical relevance of timed motor performance in children with Duchenne muscular dystrophy. Physiotherapy Theory and Practice, 2015, 31, 173-181.	1.3	11
18	Epidemiological and clinical factors impact on the benefit of riluzole in the survival rates of patients with ALS. Arquivos De Neuro-Psiquiatria, 2017, 75, 515-522.	0.8	10

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19	Qualidade de vida: anÃ¡lise comparativa entre crianÃ§as com distrofia muscular de Duchenne e seus cuidadores. Revista Neurociencias, 2008, 16, 275-279.	0.0	10
20	Patients with Duchenne and Becker muscular dystrophies are not more asymmetrical than healthy controls on timed performance of upper limb tasks. Brazilian Journal of Medical and Biological Research, 2017, 50, e6031.	1.5	9
21	Improvements in motor tasks through the use of smartphone technology for individuals with Duchenne muscular dystrophy. Neuropsychiatric Disease and Treatment, 2017, Volume 13, 2209-2217.	2.2	8
22	Prolonged sitting and physical discomfort in university students. Acta FisiÃ¡trica, 2015, 22, .	0.1	8
23	Hand Function in Muscular Dystrophies. Perceptual and Motor Skills, 2017, 124, 441-451.	1.3	6
24	VersÃ£o brasileira da Segmental Assessment of Trunk Control (SATCo). Fisioterapia E Pesquisa, 2017, 24, 89-99.	0.1	6
25	Boys With Duchenne Muscular Dystrophy: 1-Year Locomotor Changes in Relation to a Control Group. Perceptual and Motor Skills, 2018, 125, 40-56.	1.3	6
26	Technological Tools for Observational Evaluation - the Experience with the Software for Functional Evaluation Scale for Duchenne Muscular Dystrophy â€“ A Pilot Study Software for Observational Evaluation. British Journal of Medicine and Medical Research, 2016, 16, 1-7.	0.2	6
27	Cardiac Autonomic Modulation in Subjects with Amyotrophic Lateral Sclerosis (ALS) during an Upper Limb Virtual Reality Task: A Prospective Control Trial. BioMed Research International, 2022, 2022, 1-11.	1.9	6
28	Trunk Control and Upper Limb Function of Walking and Non-walking Duchenne Muscular Dystrophy Individuals. Developmental Neurorehabilitation, 2021, 24, 435-441.	1.1	5
29	Principais instrumentos para a anÃ¡lise da marcha de pacientes com distrofia muscular de Duchenne. Revista Neurociencias, 2007, 15, 153-159.	0.0	5
30	O efeito do treinamento muscular respiratÃ³rio na miastenia grave. Revista Neurociencias, 2009, 17, 37-45.	0.0	4
31	Timed immersion expiration measures in patients with muscular dystrophies. Archives of Physiotherapy, 2020, 10, 4.	1.8	3
32	Ã“rteses de membros superiores para pessoas com lesÃ£o medular a nÃvel cervical: revisÃ£o integrativa. Revista Neurociencias, 0, 29, 1-19.	0.0	2
33	AvaliaÃ§Ã£o das atividades da ABRELA:. Revista Neurociencias, 2006, 14, 191-197.	0.0	2
34	Efeitos da LiberaÃ§Ã£o Miofascial Sobre a Flexibilidade de um Paciente com Distrofia MiotÃ´nica de Steinert. Revista Neurociencias, 2012, 20, 404-409.	0.0	2
35	Motor learning through a non-immersive virtual task in people with limb-girdle muscular dystrophies. Journal of Human Growth and Development, 2020, 30, 461-471.	0.6	2
36	Comparison of motor function in patients with Duchenne muscular dystrophy in physical therapy in and out of water: 2-year follow-up. Acta FisiÃ¡trica, 2015, 22, .	0.1	1

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37	IndependÃªncia funcional de crianÃ§as de um a quatro anos com mielomeningocele. Fisioterapia E Pesquisa, 2018, 25, 196-201.	0.1	1
38	Perfil clÃnico de pacientes com doenÃ§a do neurÃ´nio motor no ambulatÃ³rio da Unifesp. Revista Neurociencias, 2008, 16, 189-193.	0.0	1
39	A ImportÃ¢ncia das Ã“rteses de Membros Inferiores na Distrofia Muscular de Duchenne. Revista Neurociencias, 2012, 20, 584-587.	0.0	1
40	Funcionalidade de membro superior em pacientes deambuladores e nÃ£o deambuladores com distrofia muscular de Duchenne. Fisioterapia E Pesquisa, 2020, 27, 188-193.	0.1	1
41	AnÃ¡lise da imagem e esquema corporal em indivÃduos com diagnÃ³stico de sÃndrome pÃ³s-poliomielite. Revista Neurociencias, 2012, 20, 50-57.	0.0	0
42	OrientaÃ§Ãµes emergenciais para profissionais que assistem pacientes com Esclerose Lateral AmiotrÃ³fica. Revista Neurociencias, 2012, 20, 260-265.	0.0	0
43	Fatores preditivos para a falÃªncia respiratÃ³ria na sÃndrome de Guillain-BarrÃ©. Revista Neurociencias, 2010, 18, 87-94.	0.0	0
44	A Equoterapia na Distrofia Muscular de Duchenne. Revista Neurociencias, 2010, 18, 479-484.	0.0	0
45	CaracterizaÃ§Ã£o de alteraÃ§Ãµes do sono de pacientes com SÃndrome PÃ³s-Poliomielite pela Polissonografia. Revista Neurociencias, 2011, 19, 18-25.	0.0	0
46	A PercepÃ§Ã£o dos Pacientes com Distrofia Muscular de Duchenne em RelaÃ§Ã£o a sua Acessibilidade Domiciliar. Revista Neurociencias, 2013, 21, 94-101.	0.0	0
47	A PercepÃ§Ã£o dos Pacientes com Distrofia Muscular de Duchenne em RelaÃ§Ã£o a sua Acessibilidade Domiciliar. Revista Neurociencias, 0, 21, 94-101.	0.0	0
48	AvaliaÃ§Ã£o espacial e temporal do ciclo da marcha nas distrofias musculares. Revista Neurociencias, 2014, 22, 286-293.	0.0	0
49	Space and temporal assessment of the gait cycle in muscular dystrophies. Revista Neurociencias, 2014, 22, 286-293.	0.0	0
50	Retrospective study on the behavior of muscular strength of Post-Poliomyelitis Syndrome patients. Revista Neurociencias, 2014, 22, 351-358.	0.0	0
51	Epidemiological and clinical factors impact on survival in ALS/ MND: a cohort study. International Archive of Medicine, 0, , .	1.2	0
52	Effects of a single-session massage for sedentary older women with prehypertension: a pilot study. International Archive of Medicine, 0, , .	1.2	0
53	Effects of modified Restricted Environmental Stimulation Therapy on relaxation, heart rate, blood pressure and flexibility. International Archive of Medicine, 0, , .	1.2	0
54	Genetic and functional differences between Bethlem miopathyand Ullrich congenital muscular dystrophy â€“ case studies. Cadernos De PÃ³s-GraduaÃ§Ã£o Em DistÃºrbios Do Desenvolvimento, 2018, 18, .	0.1	0

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55	ExercÃcios aquÃ¡ticos e em solo para pacientes com Esclerose Lateral AmiotrÃ³fica. Revista Neurociencias, 2009, 17, 170-177.	0.0	0
56	Esclerose Lateral AmiotrÃ³fica (ELA). Revista Neurociencias, 0, 17, 1-2.	0.0	0
57	MudanÃ§a na biomecÃ¢nica da postura sentada afeta a funÃ§Ã£o pulmonar. Fisioterapia E Pesquisa, 2019, 26, 265-274.	0.1	0
58	Sobreviventes de Poliomielite em Ã©poca de pandemia por COVID-19 (SARS CoV-2). Research, Society and Development, 2022, 11, e54811730014.	0.1	0