JosÃ© CÃ©sar Milisenda

List of Publications by Year
in descending order

61 papers	1,451 citations	430874 18 h-index	361022 35 g-index
70 all docs	70 docs citations	70 times ranked	1912 citing authors

#	Article	IF	Citations
1	Clinicoâ€“pathological phenotypes of systemic sclerosisâ€“associated myopathy: analysis of a large multicentre cohort. Rheumatology, 2023, 62, SI82-SI90.	1.9	8
2	Esophageal diffuse spasms in patient with Pompe disease. Medicina ClÃ¬nica, 2022, , .	0.6	0
3	Gastrointestinal Involvement in Dermatomyositis. Diagnostics, 2022, 12, 1200.	2.6	6
4	A multidisciplinary registry of patients with autoimmune and immune-mediated diseases with symptomatic COVID-19 from a single center. Journal of Autoimmunity, 2021, 117, 102580.	6.5	23
5	Association of Initial Maximal Motor Ability With Long-term Functional Outcome in Patients With COL6-Related Dystrophies. Neurology, 2021, 96, e1413-e1424.	1.1	10
6	Response to: â€˜Correspondence on â€˜Machine learning algorithms reveal unique gene expression profiles in muscle biopsies from patients with different types of myositisâ€™â€™ by Takanashi et al. Annals of the Rheumatic Diseases, 2021, , annrheumdis-2020-219767.	0.9	1
7	Clinical characteristics and outcome of patients aged over 80 years with covid-19. Medicine (United) Tj ETQq1 1 0.784314 rgBT /Over	1.0	5
8	Differential diagnosis of necrotizing myopathy. Current Opinion in Rheumatology, 2021, 33, 544-553.	4.3	11
9	Accumulation of autophagosome cargo protein p62 is common in idiopathic inflammatory myopathies. Clinical and Experimental Rheumatology, 2021, 39, 351-356.	0.8	2
10	Accumulation of autophagosome cargo protein p62 is common in idiopathic inflammatory myopathies. Clinical and Experimental Rheumatology, 2021, 39, 351-356.	0.8	8
11	ANCA-associated vasculitic neuropathy during treatment with ipilimumab. Rheumatology, 2020, 59, 251-252.	1.9	7
12	Emerging PD-1 and PD-1L inhibitors-associated myopathy with a characteristic histopathological pattern. Autoimmunity Reviews, 2020, 19, 102455.	5.8	51
13	Statin use and myopathy. Not always guilty. Rheumatology, 2020, 59, 3853-3857.	1.9	1
14	Ultra-efficient sequencing of T Cell receptor repertoires reveals shared responses in muscle from patients with Myositis. EBioMedicine, 2020, 59, 102972.	6.1	11
15	A disabling case of chronic external ophthalmoplegia cleverly overcomed. Medicina ClÃ¬nica (English) Tj ETQq1 1 0.784314 rgBT /Over	0.2	0
16	Mitochondrial Dysfunction: A Common Hallmark Underlying Comorbidity between sIBM and Other Degenerative and Age-Related Diseases. Journal of Clinical Medicine, 2020, 9, 1446.	2.4	4
17	Targeted Next-Generation Sequencing in a Large Cohort of Genetically Undiagnosed Patients with Neuromuscular Disorders in Spain. Genes, 2020, 11, 539.	2.4	25
18	Anoctamin 5 (ANO5) muscular dystrophyâ€”three different phenotypes and a new histological pattern. Neurological Sciences, 2020, 41, 2967-2971.	1.9	6

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19	Machine learning algorithms reveal unique gene expression profiles in muscle biopsies from patients with different types of myositis. Annals of the Rheumatic Diseases, 2020, 79, 1234-1242.	0.9	80
20	Massive deep venous thrombosis in a patient addicted to cocaine. Medicina ClÃnica (English Edition), 2020, 154, 110-111.	0.2	0
21	Neutrophil dysregulation is pathogenic in idiopathic inflammatory myopathies. JCI Insight, 2020, 5, .	5.0	65
22	A disabling case of chronic external ophthalmoplegia cleverly overcomed. Medicina ClÃnica, 2020, 155, 570.	0.6	0
23	Massive deep venous thrombosis in a patient addicted to cocaine. Medicina ClÃnica, 2020, 154, 110-111.	0.6	0
24	Deep morphological analysis of muscle biopsies from type III glycogenesis (GSDIII), debranching enzyme deficiency, revealed stereotyped vacuolar myopathy and autophagy impairment. Acta Neuropathologica Communications, 2019, 7, 167.	5.2	17
25	Identification of distinctive interferon gene signatures in different types of myositis. Neurology, 2019, 93, e1193-e1204.	1.1	115
26	Diffusionâ€weighted magnetic resonance imaging is useful for assessing inflammatory myopathies. Muscle and Nerve, 2019, 59, 555-560.	2.2	10
27	Expanding the importance of HMERF titinopathy: new mutations and clinical aspects. Journal of Neurology, 2019, 266, 680-690.	3.6	31
28	Myositis Autoantigen Expression Correlates With Muscle Regeneration but Not Autoantibody Specificity. Arthritis and Rheumatology, 2019, 71, 1371-1376.	5.6	29
29	Mitochondrial implications in human pregnancies with intrauterine growth restriction and associated cardiac remodelling. Journal of Cellular and Molecular Medicine, 2019, 23, 3962-3973.	3.6	19
30	Sporadic inclusion body myositis: Diagnostic value of p62 immunostaining. Medicina ClÃnica (English) Tj ETQq0 0 0 rgBT /Overlock 10 T	0.2	0
31	Whole-body MRI and pathological findings in adult patients with myopathies. Skeletal Radiology, 2019, 48, 653-676.	2.0	21
32	Sporadic inclusion body myositis: Diagnostic value of p62 immunostaining. Medicina ClÃnica, 2019, 153, 437-440.	0.6	0
33	Correlation between quantitative and semiquantitative magnetic resonance imaging and histopathology findings in dermatomyositis. Clinical and Experimental Rheumatology, 2019, 37, 633-640.	0.8	5
34	Statin-induced myalgia and myositis: an update on pathogenesis and clinical recommendations. Expert Review of Clinical Immunology, 2018, 14, 215-224.	3.0	112
35	Transcriptional alterations in skin fibroblasts from Parkinson's disease patients with parkin mutations. Neurobiology of Aging, 2018, 65, 206-216.	3.1	13
36	Tumour TIF1 mutations and loss of heterozygosity related to cancer-associated myositis. Rheumatology, 2018, 57, 388-396.	1.9	81

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37	Cardiac and placental mitochondrial characterization in a rabbit model of intrauterine growth restriction. Biochimica Et Biophysica Acta - General Subjects, 2018, 1862, 1157-1167.	2.4	14
38	Amyloid polyneuropathy in 2 patients after liver transplantation. Neurology, 2018, 90, 38-38.	1.1	1
39	Exhaustion of mitochondrial and autophagic reserve may contribute to the development of LRRK2 G2019S -Parkinsonâ€™s disease. Journal of Translational Medicine, 2018, 16, 160.	4.4	22
40	Muscular and extramuscular clinical features of patients with anti-PM/Scl autoantibodies. Neurology, 2018, 90, e2068-e2076.	1.1	76
41	Classification and management of adult inflammatory myopathies. Lancet Neurology, The, 2018, 17, 816-828.	10.2	267
42	Clinical characteristics of adult patients with inborn errors of metabolism in Spain: A review of 500 cases from university hospitals. Molecular Genetics and Metabolism Reports, 2017, 10, 92-95.	1.1	10
43	â€œExtendedâ€•Beevor's sing as a new clinical sign in sporadic inclusion body myositis. Medicina ClÃnica (English Edition), 2017, 148, e43.	0.2	3
44	Placental Mitochondrial Toxicity, Oxidative Stress, Apoptosis, and Adverse Perinatal Outcomes in HIV Pregnancies Under Antiretroviral Treatment Containing Zidovudine. Journal of Acquired Immune Deficiency Syndromes (1999), 2017, 75, e113-e119.	2.1	28
45	Miositis focal secundaria a radiculopatÃa. Medicina ClÃnica, 2017, 149, 371-372.	0.6	2
46	Imbalance in mitochondrial dynamics and apoptosis in pregnancies among HIV-infected women on HAART with obstetric complications. Journal of Antimicrobial Chemotherapy, 2017, 72, 2578-2586.	3.0	11
47	Influence of Mitochondrial Genetics on the Mitochondrial Toxicity of Linezolid in Blood Cells and Skin Nerve Fibers. Antimicrobial Agents and Chemotherapy, 2017, 61, .	3.2	33
48	HIV â€‹promonocytic and lymphoid cell lines: an in vitro model of in vivo mitochondrial and apoptotic lesion. Journal of Cellular and Molecular Medicine, 2017, 21, 402-409.	3.6	11
49	Signo de Beevor Â«extendidoÂ». Un nuevo signo clÃnico en miositis con cuerpos de inclusiÃ³n. Medicina ClÃnica, 2017, 148, e43.	0.6	3
50	Mitochondrial toxicity and caspase activation in HIV pregnant women. Journal of Cellular and Molecular Medicine, 2017, 21, 26-34.	3.6	5
51	Focal myositis associated to radiculopathy. Medicina ClÃnica (English Edition), 2017, 149, 371-372.	0.2	0
52	Colonic Oxidative and Mitochondrial Function in Parkinsonâ€™s Disease and Idiopathic REM Sleep Behavior Disorder. Parkinson's Disease, 2017, 2017, 1-7.	1.1	3
53	Mitochondrial DNA disturbances and deregulated expression of oxidative phosphorylation and mitochondrial fusion proteins in sporadic inclusion body myositis. Clinical Science, 2016, 130, 1741-1751.	4.3	33
54	Not only bright tongue sign in Pompe disease. Neurology, 2016, 87, 1629-1630.	1.1	5

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55	Enteropathy associated with chronic use of olmesartan. Medicina ClÃnica (English Edition), 2015, 144, 139-140.	0.2	0
56	Polymyositis, a very uncommon isolated disease: clinical and histological re-evaluation after long-term follow-up. Rheumatology International, 2015, 35, 915-920.	3.0	32
57	Infected False Aneurysm of the Aortic Arch After Endoscopic Transurethral Instillation of Bacillus Calmette-GuÃ©rin. Annals of Thoracic Surgery, 2015, 100, 717-720.	1.3	10
58	Dermatomyositis presenting with severe subcutaneous edema: Five additional cases and review of the literature. Seminars in Arthritis and Rheumatism, 2014, 44, 228-233.	3.4	50
59	The diagnosis and classification of polymyositis. Journal of Autoimmunity, 2014, 48-49, 118-121.	6.5	36
60	A case of Japanese encephalitis in a 20 year-old Spanish sportsman, February 2013. Eurosurveillance, 2013, 18, 20573.	7.0	10
61	Miositis con cuerpos de inclusiÃ³n (forma esporÃ¡dica). Seminarios De La Fundaciâˆšâ‰¥n EspaâˆšÂ±ola De Reumatologâˆšâ‰a, 2012, 13, 23-30.	0.1	3