

2017 European League Against Rheumatism/American College of Rheumatology
classification criteria for adult and juvenile idiopathic arthritis in
major subgroups

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Citation Report

#	ARTICLE	IF	CITATIONS
1	EULAR/ACR classification criteria for adult and juvenile idiopathic inflammatory myopathies and their major subgroups: a methodology report. <i>RMD Open</i> , 2017, 3, e000507.	1.8	115
2	Consideration of Antisynthetase Syndrome Features in Classifying Patients as Having Idiopathic Inflammatory Myopathy: Comment on the Article by Lundberg et al. <i>Arthritis and Rheumatology</i> , 2018, 70, 975-976.	2.9	6
3	Current diagnosis and treatment of polymyositis and dermatomyositis. <i>Modern Rheumatology</i> , 2018, 28, 913-921.	0.9	82
4	Gastrointestinal Tract Vasculopathy. <i>American Journal of Surgical Pathology</i> , 2018, 42, 866-876.	2.1	46
5	Dermatomyositis Clinical and Pathological Phenotypes Associated with Myositis-Specific Autoantibodies. <i>Current Rheumatology Reports</i> , 2018, 20, 28.	2.1	76
6	Classification of myositis. <i>Nature Reviews Rheumatology</i> , 2018, 14, 269-278.	3.5	210
7	Applicability of EULAR/ACR classification criteria for dermatomyositis to amyopathic disease. <i>Journal of the American Academy of Dermatology</i> , 2018, 79, 77-83.e1.	0.6	42
8	Response to: "Performance of the 2017 European League Against Rheumatism/American College of Rheumatology classification criteria for adult and juvenile idiopathic inflammatory myopathies in clinical practice" by Hoëvar et al. <i>Annals of the Rheumatic Diseases</i> , 2018, 77, e91-e91.	0.5	43
9	Bench to bedside review of myositis autoantibodies. <i>Clinical and Molecular Allergy</i> , 2018, 16, 5.	0.8	59
10	New Myositis Classification Criteria" What We Have Learned Since Bohan and Peter. <i>Current Rheumatology Reports</i> , 2018, 20, 18.	2.1	65
11	2017 EULAR/ACR classification criteria for adult and juvenile idiopathic inflammatory myopathies and their major subgroups: little emphasis on autoantibodies, why?. <i>Annals of the Rheumatic Diseases</i> , 2018, 77, e77-e77.	0.5	23
12	Response to: '2017 EULAR/ACR classification criteria for adult and juvenile idiopathic inflammatory myopathies and their major subgroups: little emphasis on autoantibodies, why?' by Malaviya. <i>Annals of the Rheumatic Diseases</i> , 2018, 77, e78-e78.	0.5	19
13	Performance of the 2017 European League Against Rheumatism/American College of Rheumatology classification criteria for adult and juvenile idiopathic inflammatory myopathies in clinical practice. <i>Annals of the Rheumatic Diseases</i> , 2018, 77, e90-e90.	0.5	14
14	Myositis in clinical practice"relevance of new antibodies. <i>Best Practice and Research in Clinical Rheumatology</i> , 2018, 32, 887-901.	1.4	3
15	OP0147"Aberrant activation of type I interferon system in anti-mda5 dermatomyositis patients. , 2018, , .		0
16	Myositis an evolving spectrum of disease. <i>Immunological Medicine</i> , 2018, 41, 46-54.	1.4	14
18	Increasing incidence of adult idiopathic inflammatory myopathies in the City of Salford, UK: a 10-year epidemiological study. <i>Rheumatology Advances in Practice</i> , 2018, 2, rky035.	0.3	15
19	Current Treatment for Myositis. <i>Current Treatment Options in Rheumatology</i> , 2018, 4, 299-315.	0.6	36

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20	Systemic and Tissue Inflammation in Juvenile Dermatomyositis: From Pathogenesis to the Quest for Monitoring Tools. <i>Frontiers in Immunology</i> , 2018, 9, 2951.	2.2	50
21	Utility of [18F] Fluorodeoxyglucose Positron Emission Tomography/Computed Tomography Scan in Inflammatory Myopathies: Case Report and Literature Review. <i>Case Reports in Rheumatology</i> , 2018, 2018, 1-5.	0.2	1
22	Patients with anti-Jo1 antibodies display a characteristic IgG Fc-glycan profile which is further enhanced in anti-Jo1 autoantibodies. <i>Scientific Reports</i> , 2018, 8, 17958.	1.6	12
23	Miopatías inflamatorias. <i>Revista Médica Clínica Las Condes</i> , 2018, 29, 611-621.	0.2	2
25	Proximal muscle weakness and skin rash. <i>BMJ: British Medical Journal</i> , 2018, 363, k3614.	2.4	0
26	New insights in myositis-specific autoantibodies. <i>Current Opinion in Rheumatology</i> , 2018, 30, 614-622.	2.0	37
28	Autoimmune Myopathies: Updates on Evaluation and Treatment. <i>Neurotherapeutics</i> , 2018, 15, 976-994.	2.1	55
29	The diagnostic work-up of cancer-associated myositis. <i>Current Opinion in Rheumatology</i> , 2018, 30, 630-636.	2.0	37
30	T-cell transcriptomics from peripheral blood highlights differences between polymyositis and dermatomyositis patients. <i>Arthritis Research and Therapy</i> , 2018, 20, 188.	1.6	21
31	Recent developments in classification criteria and diagnosis guidelines for idiopathic inflammatory myopathies. <i>Current Opinion in Rheumatology</i> , 2018, 30, 606-613.	2.0	17
32	Prediction of autoimmune connective tissue disease in an at-risk cohort: prognostic value of a novel two-score system for interferon status. <i>Annals of the Rheumatic Diseases</i> , 2018, 77, 1432-1439.	0.5	79
33	Current Classification and Management of Inflammatory Myopathies. <i>Journal of Neuromuscular Diseases</i> , 2018, 5, 109-129.	1.1	190
34	Interstitial Pneumonia With Autoimmune Features. <i>Arthritis and Rheumatology</i> , 2018, 70, 1901-1913.	2.9	38
35	Long-term follow-up of nailfold videocapillaroscopic changes in dermatomyositis versus systemic sclerosis patients. <i>Clinical Rheumatology</i> , 2018, 37, 2723-2729.	1.0	14
36	Targeted lipidomics analysis identified altered serum lipid profiles in patients with polymyositis and dermatomyositis. <i>Arthritis Research and Therapy</i> , 2018, 20, 83.	1.6	22
37	Classification and management of adult inflammatory myopathies. <i>Lancet Neurology</i> , The, 2018, 17, 816-828.	4.9	267
38	Expression of interleukin-18 in muscle tissue of patients with polymyositis or dermatomyositis and effects of conventional immunosuppressive treatment. <i>Rheumatology</i> , 2018, 57, 2149-2157.	0.9	13
39	Muscle pain syndromes and fibromyalgia: the role of muscle biopsy. <i>Current Opinion in Supportive and Palliative Care</i> , 2018, 12, 382-387.	0.5	17

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40	The incidence of idiopathic inflammatory myopathies in the adult Slovenian population. <i>Clinical Rheumatology</i> , 2019, 38, 279-283.	1.0	3
41	Myositis autoantibody profiles and their clinical associations in Greek patients with inflammatory myopathies. <i>Clinical Rheumatology</i> , 2019, 38, 125-132.	1.0	35
42	Response to: "Time to personalise the treatment of anti-MDA-5 associated lung disease" by Lake et al. <i>Annals of the Rheumatic Diseases</i> , 2019, 78, e53-e53.	0.5	1
44	Including myositis-specific autoantibodies improves performance of the idiopathic inflammatory myopathies classification criteria. <i>Rheumatology</i> , 2019, 58, 2331-2333.	0.9	4
45	Imaging of Inflammatory Myopathies. , 2019, , 2185-2197.		0
46	Effectiveness of Tai Chi on fibromyalgia patients: A meta-analysis of randomized controlled trials. <i>Complementary Therapies in Medicine</i> , 2019, 46, 1-8.	1.3	35
47	Dysphagia as Isolated Manifestation of Jo-1 Associated Myositis?. <i>Frontiers in Neurology</i> , 2019, 10, 739.	1.1	11
48	Undifferentiated connective tissue disease: state of the art on clinical practice guidelines. <i>RMD Open</i> , 2019, 4, e000786.	1.8	28
49	Comment on: Muscle fluorodeoxyglucose uptake assessed by positron emission tomography-computed tomography as a biomarker of inflammatory myopathies disease activity: reply. <i>Rheumatology</i> , 2019, 58, 2345-2346.	0.9	3
50	Idiopathic and immune-related pulmonary fibrosis: diagnostic and therapeutic challenges. <i>Clinical and Translational Immunology</i> , 2019, 8, e1086.	1.7	22
51	Demographic and clinical predictors of progression and mortality in connective tissue disease-associated interstitial lung disease: a retrospective cohort study. <i>BMC Pulmonary Medicine</i> , 2019, 19, 192.	0.8	37
52	The Role of the Multidisciplinary Evaluation of Interstitial Lung Diseases: Systematic Literature Review of the Current Evidence and Future Perspectives. <i>Frontiers in Medicine</i> , 2019, 6, 246.	1.2	59
53	Idiopathic Inflammatory Myopathies. <i>Rheumatic Disease Clinics of North America</i> , 2019, 45, 569-581.	0.8	41
54	Comment on: Muscle fluorodeoxyglucose uptake assessed by positron emission tomography-computed tomography as a biomarker of inflammatory myopathies disease activity. <i>Rheumatology</i> , 2019, 58, 2344-2345.	0.9	0
55	Evaluation of a novel particle-based assay for detection of autoantibodies in idiopathic inflammatory myopathies. <i>Journal of Immunological Methods</i> , 2019, 474, 112661.	0.6	18
56	Clinical Applications of Synovial Biopsy. <i>Frontiers in Medicine</i> , 2019, 6, 102.	1.2	5
57	Tertiary lymphoid organs in the inflammatory myopathy associated with PD-1 inhibitors. , 2019, 7, 256.		13
58	Development and validation of a composite disease activity score for measurement of muscle and skin involvement in juvenile dermatomyositis. <i>Rheumatology</i> , 2019, 58, 1196-1205.	0.9	10

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59	Efficacy and safety of rituximab in anti-synthetase antibody positive and negative subjects with idiopathic inflammatory myopathy: a registry-based study. <i>Rheumatology</i> , 2019, 58, 1214-1220.	0.9	22
60	Imaging in the diagnosis of idiopathic inflammatory myopathies; indications and utility. <i>Expert Review of Neurotherapeutics</i> , 2019, 19, 173-184.	1.4	7
61	Anti-Ku syndrome with elevated CK and anti-Ku syndrome with anti-dsDNA are two distinct entities with different outcomes. <i>Annals of the Rheumatic Diseases</i> , 2019, 78, 1101-1106.	0.5	54
62	Biomarkers in Inflammatory Myopathies—An Expanded Definition. <i>Frontiers in Neurology</i> , 2019, 10, 554.	1.1	48
63	Validation of 2017 classification criteria for adult and juvenile idiopathic inflammatory myopathies proposed by EULAR/ACR in Chinese patients. <i>International Journal of Rheumatic Diseases</i> , 2019, 22, 1278-1282.	0.9	15
64	An Anti-PM/Sci-75 Antibody-positive Japanese Woman Who Developed Inflammatory Myopathy. <i>Internal Medicine</i> , 2019, 58, 2689-2693.	0.3	4
66	Idiopathic inflammatory myopathy comorbid with Sturge-Weber syndrome: a case report. <i>BMC Neurology</i> , 2019, 19, 87.	0.8	1
67	Comparison of Three Immunoassays for the Detection of Myositis Specific Antibodies. <i>Frontiers in Immunology</i> , 2019, 10, 848.	2.2	54
68	Comparison of the 2017 EULAR/ACR criteria with Bohan and Peter criteria for the classification of idiopathic inflammatory myopathies. <i>Clinical Rheumatology</i> , 2019, 38, 1931-1934.	1.0	18
69	Successful treatment of extensive calcifications and acute pulmonary involvement in dermatomyositis with the Janus-Kinase inhibitor tofacitinib — A report of two cases. <i>Journal of Autoimmunity</i> , 2019, 100, 131-136.	3.0	78
70	The IgG2 Isotype of Anti-Transcription Intermediary Factor 1 β Autoantibodies Is a Biomarker of Cancer and Mortality in Adult Dermatomyositis. <i>Arthritis and Rheumatology</i> , 2019, 71, 1360-1370.	2.9	33
71	Strategy for suspected myositis. <i>Joint Bone Spine</i> , 2019, 86, 568-575.	0.8	3
72	Autoantibody testing in idiopathic inflammatory myopathies. <i>Practical Neurology</i> , 2019, 19, 284-294.	0.5	16
73	Imaging of Inflammatory Myopathies. , 2019, , 1-13.		0
74	Idiopathic inflammatory myopathies and antisynthetase syndrome: contribution of antisynthetase antibodies to improve current classification criteria. <i>Annals of the Rheumatic Diseases</i> , 2019, 78, 1291-1292.	0.5	16
75	Muscle fluorodeoxyglucose uptake assessed by positron emission tomography—computed tomography as a biomarker of inflammatory myopathies disease activity. <i>Rheumatology</i> , 2019, 58, 1459-1464.	0.9	20
76	Clinical relevance of HEp-2 indirect immunofluorescent patterns: the International Consensus on ANA patterns (ICAP) perspective. <i>Annals of the Rheumatic Diseases</i> , 2019, 78, 879-889.	0.5	217
77	Acute Coronary Syndrome in Idiopathic Inflammatory Myopathies: A Population-based Study. <i>Journal of Rheumatology</i> , 2019, 46, 1509-1514.	1.0	13

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78	MRI of skeletal muscles in patients with idiopathic inflammatory myopathies: characteristic findings and diagnostic performance in dermatomyositis. RMD Open, 2019, 5, e000850.	1.8	37
79	Idiopathic inflammatory myopathies: state of the art on clinical practice guidelines. RMD Open, 2019, 4, e000784.	1.8	19
80	Autoantibodies at the Center of (sub)Classificationâ€”Issues of Detection. JAMA Neurology, 2019, 76, 867.	4.5	6
81	Novel susceptibility alleles in HLA region for myositis and myositis specific autoantibodies in Korean patients. Seminars in Arthritis and Rheumatism, 2019, 49, 283-287.	1.6	28
82	Autoantibodies to Mi-2 alpha and Mi-2 beta in patients with idiopathic inflammatory myopathy. Rheumatology, 2019, 58, 1655-1661.	0.9	20
83	Intravenous human immunoglobulin and/or methylprednisolone pulse therapies as a possible treat-to-target strategy in immune-mediated necrotizing myopathies. Rheumatology International, 2019, 39, 1201-1212.	1.5	17
84	FRI0308â€…HIGH LEVEL OF SERUM NEOPTERIN IS ASSOCIATED WITH RAPIDLY PROGRESSIVE INTERSTITIAL LUNG DISEASE AND REDUCED SURVIVAL IN DERMATOMYOSITIS. , 2019, , .		0
85	FRI0307â€…LENABASUM, A CANNABINOID TYPE 2 RECEPTOR AGONIST, REDUCES CD4 CELL POPULATIONS AND DOWNREGULATES TYPE 1 AND 2 INTERFERON ACTIVITIES IN LESIONAL DERMATOMYOSITIS SKIN. , 2019, , .		5
86	FRI0327â€…EVALUATION OF AMERICAN COLLEGE OF RHEUMATOLOGY PROVISIONAL COMPOSITE RESPONSE INDEX IN SYSTEMIC SCLEROSIS IN A PHASE II TRIAL OF ABATACEPT VS. PLACEBO. , 2019, , .		0
87	AB0697Bâ€…VALIDATION OF 2017 CLASSIFICATION CRITERIA FOR ADULT AND JUVENILE IDIOPATHIC INFLAMMATORY MYOPATHIES PROPOSED BY EULAR/ACR IN CHINESE PATIENTS. , 2019, , .		0
88	FRI0329â€…ANALYSIS OF 11 CASES OF ANTI-PL-7 ANTIBODY POSITIVE PATIENTS WITH IDIOPATHIC INFLAMMATORY MYOPATHIES. MALIGNANCY MAY NOT BE UNCOMMON COMPLICATION IN ANTI-PL-7 ANTIBODY POSITIVE MYOSITIS PATIENTS. , 2019, , .		0
89	AB0696â€…DETECTION OF COEXISTING MYOSITIS-SPECIFIC AUTOANTIBODIES WITH LINE AND DOT IMMUNOASSAYS IN PATIENTS WITH IDIOPATHIC INFLAMMATORY MYOPATHIES. , 2019, , .		0
90	FRI0328â€…BRANCHED CHAIN AMINO ACIDSIN THE TREATMENT OF POLYMYOSITIS AND DERMATOMYOSITIS: RESULTS FROM THE BTOUGH STUDY. , 2019, , .		0
91	FRI0306â€…IDIOPATHIC INFLAMMATORY MYOPATHIESAND ANTISYNTHEASE SYNDROME: CONTRIBUTION OF ANTISYNTHEASE ANTIBODIES TO IMPROVE CURRENT CLASSIFICATION CRITERIA. , 2019, , .		0
92	AB0697Câ€…THE ASSOCIATION OF ANTI-MELANOMA DIFFERENTIATION-ASSOCIATED PROTEIN 5 AND SEASONAL PATTERNS IN ONSET OF IDIOPATHIC INFLAMMATORY MYOPATHIES IN KOREA. , 2019, , .		0
93	Correlation of PMN elastase and PMN elastase-to-neutrophil ratio with disease activity in patients with myositis. Journal of Translational Medicine, 2019, 17, 420.	1.8	8
94	Advances Toward Precision Medicine in Juvenile Dermatomyositis. Current Rheumatology Reports, 2019, 21, 73.	2.1	1
95	Classification of idiopathic inflammatory myopathies: pathology perspectives. Current Opinion in Neurology, 2019, 32, 704-714.	1.8	61

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96	Pulmonary involvement in antisynthetase syndrome. <i>Current Opinion in Rheumatology</i> , 2019, 31, 603-610.	2.0	29
97	Juvenile Dermatomyositisâ€”Clinical Phenotypes. <i>Current Rheumatology Reports</i> , 2019, 21, 74.	2.1	38
98	Treatment consensus for management of polymyositis and dermatomyositis among rheumatologists, neurologists and dermatologists. <i>Modern Rheumatology</i> , 2019, 29, 1-19.	0.9	28
99	Treatment consensus for management of polymyositis and dermatomyositis among rheumatologists, neurologists and dermatologists. <i>Journal of Dermatology</i> , 2019, 46, e1-e18.	0.6	13
100	Significance of nailfold videocapillaroscopy in patients with idiopathic inflammatory myopathies. <i>Rheumatology</i> , 2019, 58, 120-130.	0.9	19
101	Is it time to depart from dichotomization in ALS diagnosis?. <i>Clinical Neurophysiology</i> , 2019, 130, 303-304.	0.7	0
102	Raynaudâ€™s phenomenon and antiâ€”nuclear antibody are associated with pulmonary function decline in patients with dermatomyositis and polymyositis. <i>International Journal of Rheumatic Diseases</i> , 2019, 22, 507-515.	0.9	3
103	A pattern-based approach to the interpretation of skeletal muscle biopsies. <i>Modern Pathology</i> , 2019, 32, 462-483.	2.9	14
104	Treatment consensus for management of polymyositis and dermatomyositis among rheumatologists, neurologists and dermatologists. <i>Neurology and Clinical Neuroscience</i> , 2019, 7, 3-21.	0.2	6
105	Myositis-specific autoantibodies in dermatomyositis/polymyositis with interstitial lung disease. <i>Journal of the Neurological Sciences</i> , 2019, 397, 123-128.	0.3	28
106	Mechanic hands: clinical and capillaroscopy manifestations of patients with connective tissue diseases presented with and without mechanic hands. <i>Clinical Rheumatology</i> , 2019, 38, 2309-2318.	1.0	9
107	Autoantibodies in idiopathic inflammatory myopathies: Clinical associations and laboratory evaluation by mono- and multispecific immunoassays. <i>Autoimmunity Reviews</i> , 2019, 18, 293-305.	2.5	100
108	Safety of statin drugs in patients with dyslipidemia and stable systemic autoimmune myopathies. <i>Rheumatology International</i> , 2019, 39, 311-316.	1.5	9
109	Cardiorespiratory fitness in long-term juvenile dermatomyositis: a controlled, cross-sectional study of active/inactive disease. <i>Rheumatology</i> , 2019, 58, 492-501.	0.9	14
110	The performance of the European League Against Rheumatism/American College of Rheumatology idiopathic inflammatory myopathies classification criteria in an expert-defined 10 year incident cohort. <i>Rheumatology</i> , 2019, 58, 468-475.	0.9	22
111	<i>In vivo</i> pathogenicity of IgG from patients with anti-SRP or anti-HMGCR autoantibodies in immune-mediated necrotising myopathy. <i>Annals of the Rheumatic Diseases</i> , 2019, 78, 131-139.	0.5	97
112	Diagnostic potential of sarcoplasmic myxovirus resistance protein A expression in subsets of dermatomyositis. <i>Neuropathology and Applied Neurobiology</i> , 2019, 45, 513-522.	1.8	56
113	Detection of myositis-specific antibodies: additional notes. <i>Annals of the Rheumatic Diseases</i> , 2019, 78, e29-e29.	0.5	4

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114	Detection of myositis-specific antibodies. <i>Annals of the Rheumatic Diseases</i> , 2019, 78, e7-e7.	0.5	48
115	Detection of myositis-specific antibodies: additional notes. <i>Annals of the Rheumatic Diseases</i> , 2019, 78, e45-e45.	0.5	24
116	Response to: "Detection of myositis-specific antibodies" by Vulsteke et al. <i>Annals of the Rheumatic Diseases</i> , 2019, 78, e8-e8.	0.5	3
117	Time to personalize the treatment of anti-MDA-5 associated lung disease. <i>Annals of the Rheumatic Diseases</i> , 2019, 78, e52-e52.	0.5	5
118	Response to: "Detection of myositis-specific antibodies: additional notes" by Infantino et al. <i>Annals of the Rheumatic Diseases</i> , 2019, 78, e30-e30.	0.5	3
119	Corneal Involvement of Patients with Polymyositis and Dermatomyositis. <i>Ocular Immunology and Inflammation</i> , 2020, 28, 58-66.	1.0	5
120	EBV-Associated Non-keratinizing Nasopharyngeal Carcinoma with Prominent Spindled Cell and Whorling Patterns: A Previously Unreported Histological Variant in a Patient Presenting with Dermatomyositis. <i>Head and Neck Pathology</i> , 2020, 14, 203-207.	1.3	2
121	Response to: "Comment on: "Idiopathic inflammatory myopathies and antisynthetase syndrome: contribution of antisynthetase antibodies to improve current classification criteria" by Greco et al" by Knitza et al. <i>Annals of the Rheumatic Diseases</i> , 2020, 79, e86-e86.	0.5	4
122	Comment on: "Idiopathic inflammatory myopathies and antisynthetase syndrome: contribution of antisynthetase antibodies to improve current classification criteria" by Greco et al. <i>Annals of the Rheumatic Diseases</i> , 2020, 79, e85-e85.	0.5	7
123	Risk of progression of interstitial pneumonia with autoimmune features to a systemic autoimmune rheumatic disease. <i>Rheumatology</i> , 2020, 59, 1233-1240.	0.9	17
124	Dermatomyositis: Diagnosis and treatment. <i>Journal of the American Academy of Dermatology</i> , 2020, 82, 283-296.	0.6	75
125	239th ENMC International Workshop: Classification of dermatomyositis, Amsterdam, the Netherlands, 14-16 December 2018. <i>Neuromuscular Disorders</i> , 2020, 30, 70-92.	0.3	148
126	The myositis clinical phenotype associated with anti-Zo autoantibodies: a case series of nine UK patients. <i>Rheumatology</i> , 2020, 59, 1626-1631.	0.9	10
127	Retrospective Analysis of Cancer-Associated Myositis Patients over the Past 3 Decades in a Hungarian Myositis Cohort. <i>Pathology and Oncology Research</i> , 2020, 26, 1749-1755.	0.9	30
128	Anti-RNP antibodies delineate a subgroup of myositis: A systematic retrospective study on 46 patients. <i>Autoimmunity Reviews</i> , 2020, 19, 102465.	2.5	14
129	Performance of the new EULAR/ACR classification criteria for idiopathic inflammatory myopathies (IIM) in a large monocentric IIM cohort. <i>Seminars in Arthritis and Rheumatism</i> , 2020, 50, 492-497.	1.6	18
130	Anti-TIF1- β autoantibodies: warning lights of a tumour autoantigen. <i>Rheumatology</i> , 2020, 59, 469-477.	0.9	43
131	Autoantibodies in connective tissue disease. <i>Best Practice and Research in Clinical Rheumatology</i> , 2020, 34, 101462.	1.4	17

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132	Interstitial pneumonia with autoimmune features: A single center prospective follow-up study. <i>Autoimmunity Reviews</i> , 2020, 19, 102451.	2.5	34
133	Needle electromyography, muscle MRI, and muscle pathology: Correlations in idiopathic inflammatory myopathies. <i>Neurology and Clinical Neuroscience</i> , 2020, 8, 28-35.	0.2	1
134	Current understanding and recent advances in myositis-specific and -associated autoantibodies detected in patients with dermatomyositis. <i>Expert Review of Clinical Immunology</i> , 2020, 16, 79-89.	1.3	14
135	Drop Head Syndrome as a Rare Complication in Mixed Connective Tissue Disease. <i>Internal Medicine</i> , 2020, 59, 729-732.	0.3	0
137	Patient insights on living with idiopathic inflammatory myopathy and the limitations of disease activity measurement methods – a qualitative study. <i>BMC Rheumatology</i> , 2020, 4, 47.	0.6	9
138	Functional Progression in Patients with Interstitial Lung Disease Resulted Positive to Antisynthetase Antibodies: A Multicenter, Retrospective Analysis. <i>Journal of Clinical Medicine</i> , 2020, 9, 3033.	1.0	4
139	Relationship between change in physical activity and in clinical status in patients with idiopathic inflammatory myopathy: A prospective cohort study. <i>Seminars in Arthritis and Rheumatism</i> , 2020, 50, 1140-1149.	1.6	14
140	Update on dermatomyositis in adults. <i>JDDG - Journal of the German Society of Dermatology</i> , 2020, 18, 995-1013.	0.4	8
141	A novel autoantibody targeting calreticulin is associated with cancer in patients with idiopathic inflammatory myopathies. <i>Clinical and Translational Immunology</i> , 2020, 9, e1195.	1.7	5
142	MiR-193b modulates osteoarthritis progression through targeting ST3GAL4 via sialylation of CD44 and NF- κ B pathway. <i>Cellular Signalling</i> , 2020, 76, 109814.	1.7	10
143	Myositis-specific antibodies identify a distinct interstitial pneumonia with autoimmune features phenotype. <i>European Respiratory Journal</i> , 2020, 56, 2001205.	3.1	24
144	Severe muscle damage with myofiber necrosis and macrophage infiltrates characterize anti-Mi2 positive dermatomyositis. <i>Rheumatology</i> , 2021, 60, 2916-2926.	0.9	13
145	Increased Levels of Soluble CD206 Associated with Rapidly Progressive Interstitial Lung Disease in Patients with Dermatomyositis. <i>Mediators of Inflammation</i> , 2020, 2020, 1-11.	1.4	8
146	Distinct tissue injury patterns in juvenile dermatomyositis auto-antibody subgroups. <i>Acta Neuropathologica Communications</i> , 2020, 8, 125.	2.4	16
147	Predictors and Mortality of Rapidly Progressive Interstitial Lung Disease in Patients With Idiopathic Inflammatory Myopathy: A Series of 474 Patients. <i>Frontiers in Medicine</i> , 2020, 7, 363.	1.2	47
148	LEF1 mediates osteoarthritis progression through circRNF121/miR-665/MYD88 axis via NF- κ B signaling pathway. <i>Cell Death and Disease</i> , 2020, 11, 598.	2.7	33
149	Inflammatory Myopathies. <i>Neurologic Clinics</i> , 2020, 38, 661-678.	0.8	0
150	Line blot immunoassays in idiopathic inflammatory myopathies: retrospective review of diagnostic accuracy and factors predicting true positive results. <i>BMC Rheumatology</i> , 2020, 4, 28.	0.6	16

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151	Soluble IL-2 Receptor in Dermatomyositis: Its Associations with Skin Ulcers and Disease Activity. Mediators of Inflammation, 2020, 2020, 1-8.	1.4	4
152	Factors Associated With Treatment Response in Patients With Idiopathic Inflammatory Myopathies: A Registry-Based Study. Arthritis Care and Research, 2022, 74, 468-477.	1.5	3
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154	Predicting Survival Across Acute Exacerbation of Interstitial Lung Disease in Patients with Idiopathic Inflammatory Myositis: The GAP-ILD Model. Rheumatology and Therapy, 2020, 7, 967-978.	1.1	7
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488	Profile of specific and associated autoantibodies in patients with idiopathic inflammatory myopathies in a Colombian population. <i>Frontiers in Medicine</i> , 0, 9, .	1.2	1
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490	Concurrence of leukocyte chemotactic factor 2-associated amyloidosis and autoimmune diseases: A case report. <i>Frontiers in Immunology</i> , 0, 13, .	2.2	0
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497	Clinical features of anti-mitochondrial M2 antibody-positive myositis: case series of 17 patients. <i>Journal of the Neurological Sciences</i> , 2022, 442, 120391.	0.3	3
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512	V. Recent Advances in Understanding Inflammatory Myopathy. <i>The Journal of the Japanese Society of Internal Medicine</i> , 2021, 110, 2189-2195.	0.0	0
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535	Efficacy and safety of rituximab treatment in patients with idiopathic inflammatory myopathies: A systematic review and meta-analysis. <i>Frontiers in Immunology</i> , 0, 13, .	2.2	8
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548	Clinical features and outcomes of patients with myositis associated-interstitial lung disease. <i>Frontiers in Medicine</i> , 0, 9, .	1.2	6
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