

Esi M Morgan

List of Publications by Year in descending order

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times ranked

1383
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#	ARTICLE	IF	CITATIONS
1	Treatment of Sarcoidosis in <scp>US</scp> Rheumatology Practices: Data From the American College of Rheumatology's Rheumatology Informatics System for Effectiveness (<scp>RISE</scp>) Registry. Arthritis Care and Research, 2022, 74, 371-376.	1.5	8
2	Consensus Approach to a Treat-to-target Strategy in Juvenile Idiopathic Arthritis Care: Report From the 2020 PR-COIN Consensus Conference. Journal of Rheumatology, 2022, 49, 497-503.	1.0	4
3	Development and Psychometric Evaluation of the PROMIS Pediatric Pain Intensity Measure in Children and Adolescents with Chronic Pain. Journal of Pain, 2021, 22, 48-56.	0.7	11
4	Engaging patients and parents to improve mental health intervention for youth with rheumatological disease. Pediatric Rheumatology, 2021, 19, 19.	0.9	12
5	Exploring Pediatric Tele-Rheumatology Practices During COVID-19: A Survey of the PROIN Network. Frontiers in Pediatrics, 2021, 9, 642460.	0.9	11
6	Prescribing Patterns and Impact of Factors Associated with Time to Initial Biologic Therapy among Children with Non-systemic Juvenile Idiopathic Arthritis. Paediatric Drugs, 2021, 23, 171-182.	1.3	7
7	Endorsement of the OMERACT core domain set for shared decision making interventions in rheumatology trials: Results from a multi-stepped consensus-building approach. Seminars in Arthritis and Rheumatism, 2021, 51, 593-600.	1.6	13
8	Comparative effectiveness and persistence of TNFi and non-TNFi in juvenile idiopathic arthritis: a large paediatric rheumatology centre in the USA. Rheumatology, 2021, 60, 4063-4073.	0.9	7
9	Quality of Care in Childhood-onset Systemic Lupus Erythematosus: Report of an Intervention to Improve Cardiovascular and Bone Health Screening. Journal of Rheumatology, 2020, 47, 1506-1513.	1.0	7
10	Development and Acceptability of a Patient Decision Aid for Pain Management in Juvenile Idiopathic Arthritis: The JIA Option Map. Patient, 2020, 13, 719-728.	1.1	5
11	Timing matters: real-world effectiveness of early combination of biologic and conventional synthetic disease-modifying antirheumatic drugs for treating newly diagnosed polyarticular course juvenile idiopathic arthritis. RMD Open, 2020, 6, e001091.	1.8	16
12	Measures of Health Status and Quality of Life in Juvenile Idiopathic Arthritis. Arthritis Care and Research, 2020, 72, 565-576.	1.5	3
13	Identifying Research Priorities among Patients and Families of Children with Rheumatic Diseases Living in the United States. Journal of Rheumatology, 2020, 47, 1800-1806.	1.0	9
14	OMERACT Development of a Core Domain Set of Outcomes for Shared Decision-making Interventions. Journal of Rheumatology, 2019, 46, 1409-1414.	1.0	14
15	Establishing an Updated Core Domain Set for Studies in Juvenile Idiopathic Arthritis: A Report from the OMERACT 2018 JIA Workshop. Journal of Rheumatology, 2019, 46, 1006-1013.	1.0	34
16	Utility of the PROMIS Pediatric Pain Interference Scale in Juvenile Fibromyalgia. Journal of Pediatric Psychology, 2019, 44, 436-441.	1.1	10
17	Barriers to Adherence in Juvenile Idiopathic Arthritis: A Multicenter Collaborative Experience and Preliminary Results. Journal of Rheumatology, 2018, 45, 690-696.	1.0	32
18	Treating juvenile idiopathic arthritis to target: recommendations of an international task force. Annals of the Rheumatic Diseases, 2018, 77, annrheumdis-2018-213030.	0.5	183

#	ARTICLE	IF	CITATIONS
19	The Childhood Arthritis and Rheumatology Research Alliance Consensus Treatment Plans. <i>Arthritis and Rheumatology</i> , 2018, 70, 669-678.	2.9	40
20	Treat to Target in Juvenile Idiopathic Arthritis: Challenges and Opportunities. <i>Current Treatment Options in Rheumatology</i> , 2018, 4, 29-43.	0.6	3
21	Lack of Concordance in Interrater Scoring of the Provider's Global Assessment of Children With Juvenile Idiopathic Arthritis With Low Disease Activity. <i>Arthritis Care and Research</i> , 2018, 70, 162-166.	1.5	15
22	Systemwide Implementation of Patient-Reported Outcomes in Routine Clinical Care at a Children's Hospital. <i>Joint Commission Journal on Quality and Patient Safety</i> , 2018, 44, 441-453.	0.4	25
23	Development and validation of the self-reported PROMIS pediatric pain behavior item bank and short form scale. <i>Pain</i> , 2017, 158, 1323-1331.	2.0	55
24	Toward the Development of a Core Set of Outcome Domains to Assess Shared Decision-making Interventions in Rheumatology: Results from an OMERACT Delphi Survey and Consensus Meeting. <i>Journal of Rheumatology</i> , 2017, 44, 1544-1550.	1.0	21
25	Evidence for Updating the Core Domain Set of Outcome Measures for Juvenile Idiopathic Arthritis: Report from a Special Interest Group at OMERACT 2016. <i>Journal of Rheumatology</i> , 2017, 44, 1884-1888.	1.0	11
26	Establishing clinical meaning and defining important differences for Patient-Reported Outcomes Measurement Information System (PROMIS [®]) measures in juvenile idiopathic arthritis using standard setting with patients, parents, and providers. <i>Quality of Life Research</i> , 2017, 26, 565-586.	1.5	60
27	Pilot study comparing the Childhood Arthritis & Rheumatology Research Alliance (CARRA) systemic Juvenile Idiopathic Arthritis Consensus Treatment Plans. <i>Pediatric Rheumatology</i> , 2017, 15, 23.	0.9	41
28	Design and implementation of a decision aid for juvenile idiopathic arthritis medication choices. <i>Pediatric Rheumatology</i> , 2017, 15, 48.	0.9	33
29	Multicenter inception cohort of enthesitis-related arthritis: variation in disease characteristics and treatment approaches. <i>Arthritis Research and Therapy</i> , 2017, 19, 84.	1.6	46
30	Improving care delivery and outcomes in pediatric rheumatic diseases. <i>Current Opinion in Rheumatology</i> , 2016, 28, 110-116.	2.0	31
31	Rheumatology Informatics System for Effectiveness: A National Informatics-Enabled Registry for Quality Improvement. <i>Arthritis Care and Research</i> , 2016, 68, 1866-1873.	1.5	61
32	Information technology in paediatric rheumatology. <i>Clinical and Experimental Rheumatology</i> , 2016, 34, S11-S16.	0.4	4