

W Benjamin Nowell

List of Publications by Year in descending order

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43
papers

683
citations

623734

14
h-index

642732

23
g-index

50
all docs

50
docs citations

50
times ranked

991
citing authors

#	ARTICLE	IF	CITATIONS
1	Patient Perceptions and Preferences Regarding Telemedicine for Autoimmune Rheumatic Diseases Care During the COVID-19 Pandemic. Arthritis Care and Research, 2022, 74, 1049-1057.	3.4	10
2	Employment, Work Productivity, and Biologic Treatments in Self-Reported Axial Spondyloarthritis: a Cross-Sectional Study in a Female Predominant Population from the ArthritisPower Registry. Rheumatology and Therapy, 2022, 9, 663-677.	2.3	7
3	Clinical integration of patient-reported outcome measures to enhance the care of patients with SLE: a multi-centre prospective cohort study. Rheumatology, 2022, 61, 4763-4774.	1.9	3
4	Patient Perspectives on Biologics for Axial Spondyloarthritis in a Cross-sectional Study in a Predominantly Female Population: Treatment Satisfaction, Wear-off Between Doses, and Use of Supplemental Medication. Rheumatology and Therapy, 2022, 9, 509-520.	2.3	4
5	Evaluation of an Intervention to Support Patient-Rheumatologist Conversations About Escalating Treatment in Patients with Rheumatoid Arthritis: A Proof-of-Principle Study. ACR Open Rheumatology, 2022, 4, 279-287.	2.1	1
6	Protocol for the pilot randomized trial of the Cardiovascular Risk assessment for Rheumatoid Arthritis (CARE RA) intervention: a peer coach behavioral intervention. Pilot and Feasibility Studies, 2022, 8, 84.	1.2	3
7	Cannabis for Rheumatic Disease Pain: a Review of Current Literature. Current Rheumatology Reports, 2022, 24, 119-131.	4.7	6
8	Rheumatologist and Patient Mental Models for Treatment of Rheumatoid Arthritis Help Explain Low Treatment-Target Rates. ACR Open Rheumatology, 2022, 4, 700-710.	2.1	3
9	Concerns, Healthcare Use, and Treatment Interruptions in Patients With Common Autoimmune Rheumatic Diseases During the COVID-19 Pandemic. Journal of Rheumatology, 2021, 48, 603-607.	2.0	56
10	Effects of the COVID-19 Pandemic on Patients Living With Vasculitis. ACR Open Rheumatology, 2021, 3, 17-24.	2.1	10
11	Which patient-reported outcomes do rheumatology patients find important to track digitally? A real-world longitudinal study in ArthritisPower. Arthritis Research and Therapy, 2021, 23, 53.	3.5	35
12	Data Sharing Goals for Nonprofit Funders of Clinical Trials. Journal of Participatory Medicine, 2021, 13, e23011.	1.3	2
13	Patient-Powered Research Networks of the Autoimmune Research Collaborative: Rationale, Capacity, and Future Directions. Patient, 2021, 14, 699-710.	2.7	13
14	Identifying and prioritizing benefits and risks of using privacy-enhancing software through participatory design: a nominal group technique study with patients living with chronic conditions. Journal of the American Medical Informatics Association: JAMIA, 2021, 28, 1746-1755.	4.4	5
15	Social Distancing, Health Care Disruptions, Telemedicine Use, and Treatment Interruption During the COVID-19 Pandemic in Patients With or Without Autoimmune Rheumatic Disease. ACR Open Rheumatology, 2021, 3, 381-389.	2.1	19
16	Peer Review of "Machine Learning for Risk Group Identification and User Data Collection in a Herpes Simplex Virus Patient Registry: Algorithm Development and Validation Study". Jmirx Med, 2021, 2, e28919.	0.4	2
17	Tough Choices: Exploring Medication Decision-Making During Pregnancy and Lactation Among Women With Inflammatory Arthritis. ACR Open Rheumatology, 2021, 3, 475-483.	2.1	11
18	Disruptions in Rheumatology Care and the Rise of Telehealth in Response to the COVID-19 Pandemic in a Community Practice-Based Network. Arthritis Care and Research, 2021, 73, 1153-1161.	3.4	32

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19	OPEX: Development of a novel overall patient experience measure to facilitate interpretation of comparison effectiveness studies. PLoS ONE, 2021, 16, e0245598.	2.5	6
20	Treatment Satisfaction and Decision-Making from the Patient Perspective in Axial Spondyloarthritis: Real-World Data from a Descriptive Cross-Sectional Survey Study from the ArthritisPower Registry. ACR Open Rheumatology, 2021, , .	2.1	2
21	Assessing Rheumatoid Arthritis Disease Activity With Patient-Reported Outcomes Measurement Information System Measures Using Digital Technology. Arthritis Care and Research, 2020, 72, 553-560.	3.4	13
22	Identifying patient decisions and related information needs during decision making related to total knee arthroplasty. Journal of Comparative Effectiveness Research, 2020, 9, 1153-1166.	1.4	6
23	Harnessing health plan enrollee data to boost membership in patient-powered research networks. BMC Health Services Research, 2020, 20, 462.	2.2	2
24	Barriers to treatment optimization and achievement of patients' goals: perspectives from people living with rheumatoid arthritis enrolled in the ArthritisPower registry. Arthritis Research and Therapy, 2020, 22, 4.	3.5	13
25	Communicating With Patients About Software for Enhancing Privacy in Secondary Database Research Involving Record Linkage: Delphi Study. Journal of Medical Internet Research, 2020, 22, e20783.	4.3	9
26	Impact of Information Presentation Format on Preference for Total Knee Replacement Surgery. Arthritis Care and Research, 2019, 71, 379-384.	3.4	4
27	Preference Phenotypes in Support of Shared Decision-Making at Point-of-Care for Patients With Rheumatoid Arthritis: A Proof-of-Concept Study. Arthritis Care and Research, 2019, 71, 629-637.	3.4	12
28	Patient Perceptions of Unmet Medical Need in Rheumatoid Arthritis: A Cross-Sectional Survey in the USA. Rheumatology and Therapy, 2019, 6, 461-471.	2.3	52
29	PROMIS Fatigue short forms are reliable and valid in adults with rheumatoid arthritis. Journal of Patient-Reported Outcomes, 2019, 3, 14.	1.9	33
30	Information Patients Can Provide Will Strengthen the Real-World Evidence That Matters to Them. Clinical Pharmacology and Therapeutics, 2019, 106, 49-51.	4.7	10
31	Pregnancy, Periods, and "The Pill": Exploring the Reproductive Experiences of Women with Inflammatory Arthritis. ACR Open Rheumatology, 2019, 1, 125-132.	2.1	10
32	Data linkages between patient-powered research networks and health plans: a foundation for collaborative research. Journal of the American Medical Informatics Association: JAMIA, 2019, 26, 594-602.	4.4	14
33	Real-World Patient Experience on the Path to Diagnosis of Ankylosing Spondylitis. Rheumatology and Therapy, 2019, 6, 255-267.	2.3	32
34	Digital Interventions to Build a Patient Registry for Rheumatology Research. Rheumatic Disease Clinics of North America, 2019, 45, 173-186.	1.9	33
35	Contraception methods used by women with rheumatoid arthritis and psoriatic arthritis. Clinical Rheumatology, 2019, 38, 1207-1212.	2.2	9
36	Validity of Privacy-Protecting Analytical Methods That Use Only Aggregate-Level Information to Conduct Multivariable-Adjusted Analysis in Distributed Data Networks. American Journal of Epidemiology, 2019, 188, 709-723.	3.4	25

#	ARTICLE	IF	CITATIONS
37	Digital Tracking of Rheumatoid Arthritis Longitudinally (DIGITAL) Using Biosensor and Patient-Reported Outcome Data: Protocol for a Real-World Study. JMIR Research Protocols, 2019, 8, e14665.	1.0	15
38	Preference phenotypes to facilitate shared decision-making in rheumatoid arthritis. Annals of the Rheumatic Diseases, 2018, 77, 678-683.	0.9	41
39	Patient Governance in a Patient-Powered Research Network for Adult Rheumatologic Conditions. Medical Care, 2018, 56, S16-S21.	2.4	31
40	Stakeholdersâ€™ views on data sharing in multicenter studies. Journal of Comparative Effectiveness Research, 2017, 6, 537-547.	1.4	36
41	Social media for arthritis-related comparative effectiveness and safety research and the impact of direct-to-consumer advertising. Arthritis Research and Therapy, 2017, 19, 48.	3.5	20
42	When Patients Write the Guidelines: Patient Panel Recommendations for the Treatment of Rheumatoid Arthritis. Arthritis Care and Research, 2016, 68, 26-35.	3.4	33
43	Taking the Long View: Patients Perceive Benefits and Risks of Treatment as Multidimensional. Journal of Rheumatology, 0, , jrheum.220637.	2.0	0