

# David T Eton

## List of Publications by Citations

**Source:** <https://exaly.com/author-pdf/1780107/david-t-eton-publications-by-citations.pdf>

**Version:** 2024-04-25

This document has been generated based on the publications and citations recorded by exaly.com. For the latest version of this publication list, visit the link given above.

The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

41  
papers

4,114  
citations

19  
h-index

42  
g-index

42  
ext. papers

4,712  
ext. citations

3.4  
avg, IF

4.98  
L-index

#	Paper	IF	Citations
41	Guideline for the management of clinically localized prostate cancer: 2007 update. <i>Journal of Urology</i> , <b>2007</b> , 177, 2106-31	2.5	862
40	Combining anchor and distribution-based methods to derive minimal clinically important differences on the Functional Assessment of Cancer Therapy (FACT) anemia and fatigue scales. <i>Journal of Pain and Symptom Management</i> , <b>2002</b> , 24, 547-61	4.8	574
39	Minimally important differences were estimated for six Patient-Reported Outcomes Measurement Information System-Cancer scales in advanced-stage cancer patients. <i>Journal of Clinical Epidemiology</i> , <b>2011</b> , 64, 507-16	5.7	384
38	What is a clinically meaningful change on the Functional Assessment of Cancer Therapy-Lung (FACT-L) Questionnaire? Results from Eastern Cooperative Oncology Group (ECOG) Study 5592. <i>Journal of Clinical Epidemiology</i> , <b>2002</b> , 55, 285-95	5.7	299
37	Rethinking the patient: using Burden of Treatment Theory to understand the changing dynamics of illness. <i>BMC Health Services Research</i> , <b>2014</b> , 14, 281	2.9	297
36	A combination of distribution- and anchor-based approaches determined minimally important differences (MIDs) for four endpoints in a breast cancer scale. <i>Journal of Clinical Epidemiology</i> , <b>2004</b> , 57, 898-910	5.7	270
35	Combining distribution- and anchor-based approaches to determine minimally important differences: the FACIT experience. <i>Evaluation and the Health Professions</i> , <b>2005</b> , 28, 172-91	2.5	269
34	Improving quality of life in men with prostate cancer: a randomized controlled trial of group education interventions. <i>Health Psychology</i> , <b>2003</b> , 22, 443-52	5	214
33	Building a measurement framework of burden of treatment in complex patients with chronic conditions: a qualitative study. <i>Patient Related Outcome Measures</i> , <b>2012</b> , 3, 39-49	2.9	165
32	Development and description of measurement properties of an instrument to assess treatment burden among patients with multiple chronic conditions. <i>BMC Medicine</i> , <b>2012</b> , 10, 68	11.4	143
31	Development and validation of the Patient Experience with Treatment and Self-management (PETS): a patient-reported measure of treatment burden. <i>Quality of Life Research</i> , <b>2017</b> , 26, 489-503	3.7	88
30	A systematic review of patient-reported measures of burden of treatment in three chronic diseases. <i>Patient Related Outcome Measures</i> , <b>2013</b> , 4, 7-20	2.9	81
29	Factors that lessen the burden of treatment in complex patients with chronic conditions: a qualitative study. <i>Patient Preference and Adherence</i> , <b>2014</b> , 8, 339-51	2.4	80
28	Finalizing a measurement framework for the burden of treatment in complex patients with chronic conditions. <i>Patient Related Outcome Measures</i> , <b>2015</b> , 6, 117-26	2.9	74
27	Responsiveness of 8 Patient-Reported Outcomes Measurement Information System (PROMIS) measures in a large, community-based cancer study cohort. <i>Cancer</i> , <b>2017</b> , 123, 327-335	6.4	43
26	Harmonizing and consolidating the measurement of patient-reported information at health care institutions: a position statement of the Mayo Clinic. <i>Patient Related Outcome Measures</i> , <b>2014</b> , 5, 7-15	2.9	33
25	Impact of childhood psoriasis on parents of affected children. <i>Journal of the American Academy of Dermatology</i> , <b>2017</b> , 76, 286-289.e5	4.5	29

24	Healthcare provider relational quality is associated with better self-management and less treatment burden in people with multiple chronic conditions. <i>Patient Preference and Adherence</i> , <b>2017</b> , 11, 1635-1646	2.4	28
23	Constructing a conceptual framework of patient-reported outcomes for metastatic hormone-refractory prostate cancer. <i>Value in Health</i> , <b>2010</b> , 13, 613-23	3.3	24
22	Comparing Measures to Assess Health-Related Quality of Life in Heart Failure With Preserved Ejection Fraction. <i>JACC: Heart Failure</i> , <b>2018</b> , 6, 552-560	7.9	19
21	Validating the Patient Experience with Treatment and Self-Management (PETS), a patient-reported measure of treatment burden, in people with diabetes. <i>Patient Related Outcome Measures</i> , <b>2017</b> , 8, 143-156	2.0	19
20	Patient-reported outcomes in studies of complementary and alternative medicine: problems, solutions, and future directions. <i>Explore: the Journal of Science and Healing</i> , <b>2011</b> , 7, 314-9	1.4	19
19	Coping with colorectal cancer: a qualitative exploration with patients and their family members. <i>Family Practice</i> , <b>2014</b> , 31, 598-606	1.9	18
18	A method to create a standardized generic and condition-specific patient-reported outcome measure for patient care and healthcare improvement. <i>Quality of Life Research</i> , <b>2018</b> , 27, 367-378	3.7	13
17	Risk factors for poor health-related quality of life in cancer survivors with multiple chronic conditions: exploring the role of treatment burden as a mediator. <i>Patient Related Outcome Measures</i> , <b>2019</b> , 10, 89-99	2.9	12
16	Why we need response shift: an appeal to functionalism. <i>Quality of Life Research</i> , <b>2010</b> , 19, 929-30	3.7	10
15	Reserve-building activities attenuate treatment burden in chronic illness: The mediating role of appraisal and social support. <i>Health Psychology Open</i> , <b>2018</b> , 5, 2055102918773440	1.9	8
14	Patient experience after kidney transplant: a conceptual framework of treatment burden. <i>Journal of Patient-Reported Outcomes</i> , <b>2019</b> , 3, 8	2.6	7
13	Deriving and validating a brief measure of treatment burden to assess person-centered healthcare quality in primary care: a multi-method study. <i>BMC Family Practice</i> , <b>2020</b> , 21, 221	2.6	6
12	Known-groups validity and responsiveness to change of the Patient Experience with Treatment and Self-management (PETS vs. 2.0): a patient-reported measure of treatment burden. <i>Quality of Life Research</i> , <b>2020</b> , 29, 3143-3154	3.7	5
11	Systematic review of patient-reported measures of treatment burden in stroke. <i>BMJ Open</i> , <b>2019</b> , 9, e029258	3.258	5
10	Do erythropoietic-stimulating agents relieve fatigue? A review of reviews. <i>Cancer Treatment and Research</i> , <b>2011</b> , 157, 181-94	3.5	4
9	Data pooling and analysis to build a preliminary item bank: an example using bowel function in prostate cancer. <i>Evaluation and the Health Professions</i> , <b>2005</b> , 28, 142-59	2.5	3
8	Functional Outcomes and Health-Related Quality of Life Before and After Primary Total Knee Replacement for Patients From Diverse Geographic Regions. <i>Journal of Arthroplasty</i> , <b>2021</b> , 36, 1958-1964	4.4	2
7	Confirmatory Factor Analyses and Differential Item Functioning of the Patient Experience with Treatment and Self-Management (PETS vs. 2.0): A Measure of Treatment Burden. <i>Patient Related Outcome Measures</i> , <b>2020</b> , 11, 249-263	2.9	2

6	Ensuring comprehensive assessment of urinary problems in prostate cancer through patient-physician concordance. <i>Urologic Oncology: Seminars and Original Investigations</i> , <b>2014</b> , 32, 26.e25-31	2.8	1
5	Effects of patient-reported outcome assessment order.. <i>Clinical Trials</i> , <b>2022</b> , 17407745211073788	2.2	1
4	Score equivalence of paper-, tablet-, and interactive voice response system-based versions of PROMIS, PRO-CTCAE, and numerical rating scales among cancer patients. <i>Journal of Patient-Reported Outcomes</i> , <b>2021</b> , 5, 95	2.6	1
3	Longitudinal trajectories of treatment burden: A prospective survey study of adults living with multiple chronic conditions in the midwestern United States.. <i>Journal of Multimorbidity and Comorbidity</i> , <b>2022</b> , 12, 26335565221081291		1
2	Lessons Learned: Recruiting Research Participants from an Underrepresented Patient Population at a Safety Net Hospital.. <i>Journal of General Internal Medicine</i> , <b>2022</b> , 37, 922	4	0
1	Treatment Burden in People with Hypertension is Correlated with Patient Experience with Self-Management. <i>Journal of the American Board of Family Medicine</i> , <b>2021</b> , 34, 1243-1245	1.6	0