

# Lori Frank

## List of Publications by Year in descending order

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Version: 2024-02-01

47  
papers

3,582  
citations

236925

25  
h-index

243625

44  
g-index

47  
all docs

47  
docs citations

47  
times ranked

5755  
citing authors

#	ARTICLE	IF	CITATIONS
1	Guidelines for Inclusion of Patient-Reported Outcomes in Clinical Trial Protocols. JAMA - Journal of the American Medical Association, 2018, 319, 483.	7.4	507
2	The Patient-Centered Outcomes Research Institute (PCORI) National Priorities for Research and Initial Research Agenda. JAMA - Journal of the American Medical Association, 2012, 307, 1583.	7.4	497
3	The PCORI Perspective on Patient-Centered Outcomes Research. JAMA - Journal of the American Medical Association, 2014, 312, 1513.	7.4	424
4	Patient and Stakeholder Engagement in the PCORI Pilot Projects: Description and Lessons Learned. Journal of General Internal Medicine, 2016, 31, 13-21.	2.6	225
5	Health-related quality of life associated with irritable bowel syndrome: Comparison with other chronic diseases. Clinical Therapeutics, 2002, 24, 675-689.	2.5	215
6	Conceptual and practical foundations of patient engagement in research at the patient-centered outcomes research institute. Quality of Life Research, 2015, 24, 1033-1041.	3.1	202
7	Quality of Life and Disability in Patients with Treatment-Failure Gout. Journal of Rheumatology, 2009, 36, 1041-1048.	2.0	154
8	Upper gastrointestinal symptoms in North America: prevalence and relationship to healthcare utilization and quality of life. Digestive Diseases and Sciences, 2000, 45, 809-818.	2.3	144
9	Emerging Guidelines for Patient Engagement in Research. Value in Health, 2017, 20, 481-486.	0.3	144
10	Methods and impact of engagement in research, from theory to practice and back again: early findings from the Patient-Centered Outcomes Research Institute. Quality of Life Research, 2018, 27, 17-31.	3.1	128
11	Impact of cognitive impairment on mild dementia patients and mild cognitive impairment patients and their informants. International Psychogeriatrics, 2006, 18, 151-162.	1.0	104
12	SPIRIT-PRO Extension explanation and elaboration: guidelines for inclusion of patient-reported outcomes in protocols of clinical trials. BMJ Open, 2021, 11, e045105.	1.9	65
13	Engaging Patients and Stakeholders in Research Proposal Review: The Patient-Centered Outcomes Research Institute. Annals of Internal Medicine, 2014, 161, 122.	3.9	63
14	Researchers, patients, and other stakeholders's perspectives on challenges to and strategies for engagement. Research Involvement and Engagement, 2020, 6, 60.	2.9	51
15	Primary care providers' experiences with trauma patients: A qualitative study.. Psychological Trauma: Theory, Research, Practice, and Policy, 2011, 3, 37-41.	2.1	48
16	Patient-centered priorities for improving medication management and adherence. Patient Education and Counseling, 2015, 98, 102-110.	2.2	44
17	Findings from the National Memory Screening Day Program. Journal of the American Geriatrics Society, 2015, 63, 309-314.	2.6	41
18	Use of a Self-Report Constipation Questionnaire With Older Adults in Long-Term Care. Gerontologist, The, 2001, 41, 778-786.	3.9	40

#	ARTICLE	IF	CITATIONS
19	Development and psychometric performance of the schizophrenia objective functioning instrument: An interviewer administered measure of function. <i>Schizophrenia Research</i> , 2009, 107, 275-285.	2.0	40
20	Patient self-report for evaluating mild cognitive impairment and prodromal Alzheimer's disease. <i>Alzheimer's Research and Therapy</i> , 2011, 3, 35.	6.2	40
21	Development and Evaluation of a Measure to Assess Restorative Sleep. <i>Journal of Clinical Sleep Medicine</i> , 2014, 10, 733-741.	2.6	40
22	The Effect of Risperidone on Nursing Burden Associated with Caring for Patients with Dementia. <i>Journal of the American Geriatrics Society</i> , 2004, 52, 1449-1455.	2.6	38
23	Implications of response shift for micro-, meso-, and macro-level healthcare decision-making using results of patient-reported outcome measures. <i>Quality of Life Research</i> , 2021, 30, 3343-3357.	3.1	38
24	The Economics of Selective Serotonin Reuptake Inhibitors In Depression. <i>CNS Drugs</i> , 2001, 15, 59-83.	5.9	36
25	Patient and clinician views on comparative effectiveness research and engagement in research. <i>Journal of Comparative Effectiveness Research</i> , 2015, 4, 11-25.	1.4	29
26	Validation of a new symptom impact questionnaire for mild to moderate cognitive impairment. <i>International Psychogeriatrics</i> , 2006, 18, 135-149.	1.0	27
27	Measuring change in quality of life in response to <i>Helicobacter pylori</i> eradication in peptic ulcer disease: the QOLRAD. <i>Digestive Diseases and Sciences</i> , 2001, 46, 571-580.	2.3	21
28	Development of a patient-reported outcome instrument to assess complex activities of daily living and interpersonal functioning in persons with mild cognitive impairment: The qualitative research phase. <i>Alzheimer's and Dementia</i> , 2016, 12, 75-84.	0.8	19
29	Contributions of Persons Living With Dementia to Scientific Research Meetings. Results From the National Research Summit on Care, Services, and Supports for Persons With Dementia and Their Caregivers. <i>American Journal of Geriatric Psychiatry</i> , 2020, 28, 421-430.	1.2	19
30	All together now: findings from a PCORI workshop to align patient-reported outcomes in the electronic health record. <i>Journal of Comparative Effectiveness Research</i> , 2016, 5, 561-567.	1.4	18
31	Development and Validation of the Congestion Quantifier Seven-Item Test (CQ7): A Screening Tool for Nasal Congestion. <i>Value in Health</i> , 2007, 10, 457-465.	0.3	17
32	Improving the quality of person-centred healthcare from the patient perspective: development of person-centred quality indicators. <i>BMJ Open</i> , 2020, 10, e037323.	1.9	17
33	A Qualitative Evaluation of Barriers to Care for Trauma-Related Mental Health Problems Among Low-Income Minorities in Primary Care. <i>Journal of Nervous and Mental Disease</i> , 2012, 200, 438-443.	1.0	15
34	Engaging Patients and Other Non-Researchers in Health Research: Defining Research Engagement. <i>Journal of General Internal Medicine</i> , 2020, 35, 307-314.	2.6	15
35	Time and economic cost of constipation care in nursing homes. <i>Journal of the American Medical Directors Association</i> , 2002, 3, 215-23.	2.5	14
36	The Experiences of Persons Living with Dementia Planning for a Dementia Research Meeting. Lessons Learned From the National Research Summit on Care, Services, and Supports for Persons With Dementia and Their Caregivers. <i>American Journal of Geriatric Psychiatry</i> , 2020, 28, 434-442.	1.2	12

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37	Participation of persons with dementia and their caregivers in research. Journal of the American Geriatrics Society, 2021, 69, 1784-1792.	2.6	10
38	Engaging Stakeholders in the Design and Conduct of Embedded Pragmatic Clinical Trials for Alzheimer's Disease and Alzheimer's Disease-Related Dementias. Journal of the American Geriatrics Society, 2020, 68, S62-S67.	2.6	7
39	The Patient Experience of Depression and Remission. Journal of Nervous and Mental Disease, 2007, 195, 647-654.	1.0	5
40	Genetic Risk of Alzheimer's Disease: Three Wishes Now That the Genie is Out of the Bottle. Journal of Alzheimer's Disease, 2018, 66, 421-423.	2.6	3
41	Development and Analysis of Item Response Theory-based Short-form Depression Severity Scales Based on the HDRS and MADRS. Health Outcomes Research in Medicine, 2010, 1, e111-e122.	0.6	2
42	Author Response: Patient Engagement. Value in Health, 2017, 20, 1433.	0.3	2
43	RESPONSE LETTER TO DR. ASHGAR ALI. Journal of the American Geriatrics Society, 2005, 53, 1262-1263.	2.6	1
44	Meta-issues: On Writing Scientific Manuscripts With a Stakeholder Group of Persons Living With Dementia. American Journal of Geriatric Psychiatry, 2020, 28, 445-447.	1.2	1
45	MEDICAL DECISION-MAKING IN OLDER HOSPITALIZED PATIENTS. Journal of the American Geriatrics Society, 1996, 44, 336-337.	2.6	0
46	Constipation Medications. Journal of the American Medical Directors Association, 2003, 4, 56.	2.5	0
47	Raising the Bar for Self-reported Health Status. Medical Care, 2014, 52, 1007-1009.	2.4	0