Dmitry Khodyakov

List of Publications by Year
in descending order

65 papers	1,607 citations	304743 22 h-index	330143 37 g-index
68 all docs	68 docs citations	68 times ranked	2257 citing authors

#	Article	IF	Citations
1	Ethical Community-Engaged Research: A Literature Review. American Journal of Public Health, 2013, 103, e7-e14.	2.7	158
2	ExpertLens: A system for eliciting opinions from a large pool of non-collocated experts with diverse knowledge. Technological Forecasting and Social Change, 2011, 78, 1426-1444.	11.6	98
3	pSCANNER: patient-centered Scalable National Network for Effectiveness Research. Journal of the American Medical Informatics Association: JAMIA, 2014, 21, 621-626.	4.4	80
4	Conducting Online Expert panels: a feasibility and experimental replicability study. BMC Medical Research Methodology, 2011, 11, 174.	3.1	77
5	From Subject to Participant: Ethics and the Evolving Role of Community in Health Research. American Journal of Public Health, 2015, 105, 900-908.	2.7	67
6	An Exploration of the Effect of Community Engagement in Research on Perceived Outcomes of Partnered Mental Health Services Projects. Society and Mental Health, 2011, 1, 185-199.	2.2	64
7	How can we recognize continuous quality improvement?. International Journal for Quality in Health Care, 2014, 26, 6-15.	1.8	64
8	Development of System-level Performance Measures for Evaluation of Models of Care for Inflammatory Arthritis in Canada. Journal of Rheumatology, 2016, 43, 530-540.	2.0	63
9	On Measuring Community Participation in Research. Health Education and Behavior, 2013, 40, 346-354.	2.5	62
10	Practical Considerations in Using Online Modified-Delphi Approaches to Engage Patients and Other Stakeholders in Clinical Practice Guideline Development. Patient, 2020, 13, 11-21.	2.7	62
11	Reducing the Burden of Suicide in the U.S.. American Journal of Preventive Medicine, 2014, 47, 309-314.	3.0	52
12	Development of Cardiovascular Quality Indicators for Rheumatoid Arthritis: Results from an International Expert Panel Using a Novel Online Process. Journal of Rheumatology, 2015, 42, 1548-1555.	2.0	42
13	Development of key performance indicators to evaluate centralized intake for patients with osteoarthritis and rheumatoid arthritis. Arthritis Research and Therapy, 2015, 17, 322.	3.5	37
14	Lack of preregistered analysis plans allows unacceptable data mining for and selective reporting of consensus in Delphi studies. Journal of Clinical Epidemiology, 2018, 99, 96-105.	5.0	37
15	On using ethical principles of community-engaged research in translational science. Translational Research, 2016, 171, 52-62.e1.	5.0	36
16	A Novel Stakeholder Engagement Approach for Patient-centered Outcomes Research. Medical Care, 2018, 56, S41-S47.	2.4	34
17	Patient engagement in the process of planning and designing outpatient care improvements at the Veterans Administration Healthâ€care System: findings from an online expert panel. Health Expectations, 2017, 20, 130-145.	2.6	31
18	Comparative analysis of stakeholder experiences with an online approach to prioritizing patient-centered research topics. Journal of the American Medical Informatics Association: JAMIA, 2017, 24, 537-543.	4.4	29

#	Article	IF	Citations
19	Screening Tool of Older Person's Prescriptions/Screening Tools to Alert Doctors to Right Treatment Medication Criteria Modified for U.S. Nursing Home Setting. Journal of the American Geriatrics Society, 2017, 65, 586-591.	2.6	28
20	Standardizing the Clinical Definition of Opioid Withdrawal in the Neonate. Journal of Pediatrics, 2022, 243, 33-39.e1.	1.8	28
21	Acceptability of an online modified Delphi panel approach for developing health services performance measures: results from 3 panels on arthritis research. Journal of Evaluation in Clinical Practice, 2017, 23, 354-360.	1.8	27
22	An Implementation Evaluation of the Community Engagement and Planning Intervention in the CPIC Depression Care Improvement Trial. Community Mental Health Journal, 2014, 50, 312-324.	2.0	24
23	Collaborative learning framework for online stakeholder engagement. Health Expectations, 2016, 19, 868-882.	2.6	23
24	Using an Online, Modified Delphi Approach to Engage Patients and Caregivers in Determining the Patient-Centeredness of Duchenne Muscular Dystrophy Care Considerations. Medical Decision Making, 2019, 39, 1019-1031.	2.4	22
25	A Community-Partnered, Participatory, Cluster-Randomized Study of Depression Care Quality Improvement: Three-Year Outcomes. Psychiatric Services, 2017, 68, 1262-1270.	2.0	21
26	Practical Considerations for Using Online Methods to Engage Patients in Guideline Development. Patient, 2018, 11, 155-166.	2.7	19
27	Importance and Feasibility of Transitional Care for Children With Medical Complexity: Results of a Multistakeholder Delphi Process. Academic Pediatrics, 2018, 18, 94-101.	2.0	18
28	Patient and caregiver perspectives on guideline adherence: the case of endocrine and bone health recommendations for Duchenne muscular dystrophy. Orphanet Journal of Rare Diseases, 2019, 14, 205.	2.7	17
29	Expert Panel Consensus on Management of Advanced Cancer–Related Pain in Individuals With Opioid Use Disorder. JAMA Network Open, 2021, 4, e2139968.	5.9	17
30	Engaging Patients and Caregivers Managing Rare Diseases to Improve the Methods of Clinical Guideline Development: A Research Protocol. JMIR Research Protocols, 2017, 6, e57.	1.0	15
31	Consensus-Based Guidance on Opioid Management in Individuals With Advanced Cancer-Related Pain and Opioid Misuse or Use Disorder. JAMA Oncology, 2022, 8, 1107.	7.1	15
32	Response changes in Delphi processes: why is it important to provide high-quality feedback to Delphi participants?. Journal of Clinical Epidemiology, 2020, 125, 160-161.	5.0	14
33	Partnered evaluation of a community engagement intervention: use of a kickoff conference in a randomized trial for depression care improvement in underserved communities. Ethnicity and Disease, 2011, 21, S1-78-88.	2.3	13
34	ADAPT study: adaptation of evidence-informed complex population health interventions for implementation and/or re-evaluation in new contexts: protocol for a Delphi consensus exercise to develop guidance. BMJ Open, 2020, 10, e038965.	1.9	12
35	Community Partners in Care: Leveraging Community Diversity to Improve Depression Care for Underserved Populations. International Journal of Diversity in Organisations, Communities and Nations, 2009, 9, 167-182.	0.2	12
36	Community Partners in Care: Leveraging Community Diversity to Improve Depression Care for Underserved Populations. International Journal of Diversity in Organisations, Communities and Nations, 2009, 9, 167-182.	0.2	12

#	Article	IF	Citations
37	Community-Partnered Research Conference Model: The Experience of Community Partners in Care Study. Progress in Community Health Partnerships: Research, Education, and Action, 2014, 8, 83-97.	0.3	11
38	Development of a Canadian Core Clinical Dataset to Support High-quality Care for Canadian Patients with Rheumatoid Arthritis. Journal of Rheumatology, 2017, 44, 1813-1822.	2.0	11
39	Return of Value in the New Era of Biomedical Researchâ€"One Size Will Not Fit All. AJOB Empirical Bioethics, 2019, 10, 265-275.	1.6	11
40	Standardizing care to manage bleeding disorders in adolescents with heavy mensesâ€"A joint project from the ISTH pediatric/neonatal and women's health SSCs. Journal of Thrombosis and Haemostasis, 2020, 18, 2759-2774.	3.8	11
41	A qualitative analysis of the impact of healthcare personnel influenza vaccination requirements in California. Vaccine, 2014, 32, 3082-3087.	3.8	10
42	From â€œInformedâ€•to â€œEngagedâ€•Consent: Risks and Obligations in Consent for Participation in a Health Data Repository. Journal of Law, Medicine and Ethics, 2020, 48, 172-182.	0.9	10
43	Best Practices for Participant and Stakeholder Engagement in the All of Us Research Program. , 2018, , .		10
44	Development of a Metric to Detect and Decrease Low-Value Prescribing in Older Adults. JAMA Network Open, 2022, 5, e2148599.	5.9	10
45	Nature and Predictors of Response Changes in Modified-Delphi Panels. Value in Health, 2020, 23, 1630-1638.	0.3	9
46	Comparative Effectiveness of Two Models of Depression Services Quality Improvement in Health and Community Sectors. Psychiatric Services, 2017, 68, 1315-1320.	2.0	8
47	Online Modified-Delphi: a Potential Method for Continuous Patient Engagement Across Stages of Clinical Practice Guideline Development. Journal of General Internal Medicine, 2021, 36, 1746-1750.	2.6	8
48	Trust and the ethical conduct of community-engaged research. European Journal for Person Centered Healthcare, 2017, 5, 522.	0.3	8
49	Rating the seriousness of maternal and child health outcomes linked with pregnancy weight gain. Paediatric and Perinatal Epidemiology, 2021, 35, 459-468.	1.7	7
50	Getting in tune: A qualitative analysis of guest conductorâ€"musicians relationships in symphony orchestras. Poetics, 2014, 44, 64-83.	1.3	6
51	Priority-Setting to Address the Geriatric Pharmacoparadox for Pain Management: A Nursing Home Stakeholder Delphi Study. Drugs and Aging, 2021, 38, 327-340.	2.7	6
52	Evaluation of the Work Loss Data Institute's Official Disability Guidelines. Journal of Occupational and Environmental Medicine, 2018, 60, e146-e151.	1.7	5
53	Engaging Patients and Professionals to Evaluate the Seriousness of Maternal and Child Health Outcomes: Protocol for a Modified Delphi Study. JMIR Research Protocols, 2020, 9, e16478.	1.0	5
54	Ethics and Science in the Participatory Era: A Vignette-Based Delphi Study. Journal of Empirical Research on Human Research Ethics, 2017, 12, 295-309.	1.3	4

#	ARTICLE	IF	CITATIONS
55	Whole Person Care in Underresourced Communities: Stakeholder Priorities at Long-Term Follow-Up in Community Partners in Care. Ethnicity and Disease, 2018, 28, 371-380.	2.3	4
56	Participant experiences with a new online modified-Delphi approach for engaging patients and caregivers in developing clinical guidelines. European Journal for Person Centered Healthcare, 2019, 7, 476-489.	0.3	4
57	Prioritizing Needs and Outcomes for Adolescent Substance Use Treatment Planning: An Online Modified-Delphi Process. Journal of Addiction Medicine, 2020, 14, e83-e88.	2.6	3
58	Consensus-based approach to managing opioids, including opioid misuse and opioid use disorder, in patients with serious illness: protocol for a modified Delphi process. BMJ Open, 2021, 11, e045402.	1.9	3
59	The value of consent for biobanking. Nature Human Behaviour, 2021, 5, 1125-1126.	12.0	3
60	Using Stakeholder Input to Inform an Innovative Research and Policy Initiative to Improve Depression in Safety Net Communities. Progress in Community Health Partnerships: Research, Education, and Action, 2017, 11, 93-98.	0.3	2
61	Informing the Development of a Standardized Clinical Definition of Neonatal Abstinence Syndrome: Protocol for a Modified-Delphi Expert Panel. JMIR Research Protocols, 2021, 10, e25387.	1.0	2
62	The RAND/PPMD Patient-Centeredness Method: a novel online approach to engaging patients and their representatives in guideline development. European Journal for Person Centered Healthcare, 2019, 7, 470-475.	0.3	2
63	Improving Mental Health Guardianship: From Prevention to Treatment. Psychiatric Services, 2022, 73, 642-649.	2.0	2
64	The impact of panel composition and topic on stakeholder perspectives: Generating hypotheses from online maternal and child health modified-Delphi panels. Health Expectations, 2022, 25, 732-743.	2.6	2
65	Insurers' perspectives on MA value-based insurance design model. American Journal of Managed Care, 2019, 25, e198-e203.	1.1	0