

Linda Kwakkenbos

List of Publications by Year in descending order

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Version: 2024-02-01

87
papers

1,628
citations

448610

19
h-index

406436

35
g-index

89
all docs

89
docs citations

89
times ranked

2219
citing authors

#	ARTICLE	IF	CITATIONS
1	Addressing overestimation of the prevalence of depression based on self-report screening questionnaires. <i>Cmaj</i> , 2018, 190, E44-E49.	0.9	139
2	The Scleroderma Patient-centered Intervention Network (SPIN) Cohort: protocol for a cohort multiple randomised controlled trial (cmRCT) design to support trials of psychosocial and rehabilitation interventions in a rare disease context. <i>BMJ Open</i> , 2013, 3, e003563.	0.8	104
3	Systematic review of body image measures. <i>Body Image</i> , 2019, 30, 170-211.	1.9	97
4	Reporting of Positive Results in Randomized Controlled Trials of Mindfulness-Based Mental Health Interventions. <i>PLoS ONE</i> , 2016, 11, e0153220.	1.1	80
5	Psychological health and well-being in systemic sclerosis: State of the science and consensus research agenda. <i>Arthritis Care and Research</i> , 2010, 62, 1181-1189.	1.5	79
6	CONSORT extension for the reporting of randomised controlled trials conducted using cohorts and routinely collected data (CONSORT-ROUTINE): checklist with explanation and elaboration. <i>BMJ</i> , The, 2021, 373, n857.	3.0	65
7	Structural validation of the Self-Compassion Scale with a German general population sample. <i>PLoS ONE</i> , 2018, 13, e0190771.	1.1	59
8	Psychosocial Aspects of Scleroderma. <i>Rheumatic Disease Clinics of North America</i> , 2015, 41, 519-528.	0.8	58
9	Disease-related and psychosocial factors associated with depressive symptoms in patients with systemic sclerosis, including fear of progression and appearance self-esteem. <i>Journal of Psychosomatic Research</i> , 2012, 72, 199-204.	1.2	54
10	Effectiveness of Nonpharmacologic Interventions in Systemic Sclerosis: A Systematic Review. <i>Arthritis Care and Research</i> , 2015, 67, 1426-1439.	1.5	53
11	The Scleroderma Patient-Centered Intervention Network Cohort: baseline clinical features and comparison with other large scleroderma cohorts. <i>Rheumatology</i> , 2018, 57, 1623-1631.	0.9	53
12	Performance of the Patient-Reported Outcomes Measurement Information System-29 in scleroderma: a Scleroderma Patient-centered Intervention Network Cohort Study. <i>Rheumatology</i> , 2017, 56, 1302-1311.	0.9	51
13	Re-validation and screening capacity of the 6-item version of the Cancer Worry Scale. <i>Psycho-Oncology</i> , 2018, 27, 2609-2615.	1.0	43
14	Validity of the Fear of Progression Questionnaire-Short Form in patients with systemic sclerosis. <i>Arthritis Care and Research</i> , 2012, 64, 930-934.	1.5	34
15	Validation of the Self-Efficacy for Managing Chronic Disease Scale: A Scleroderma Patient-Centered Intervention Network Cohort Study. <i>Arthritis Care and Research</i> , 2016, 68, 1195-1200.	1.5	31
16	Protocol for the development of a CONSORT extension for RCTs using cohorts and routinely collected health data. <i>Research Integrity and Peer Review</i> , 2018, 3, 9.	2.2	28
17	Major Depression Diagnoses Among Patients With Systemic Sclerosis: Baseline and One-Month Followup. <i>Arthritis Care and Research</i> , 2015, 67, 411-416.	1.5	26
18	Changes in mental health symptoms from pre-COVID-19 to COVID-19 among participants with systemic sclerosis from four countries: A Scleroderma Patient-centered Intervention Network (SPIN) Cohort study. <i>Journal of Psychosomatic Research</i> , 2020, 139, 110262.	1.2	25

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19	Effects of a multi-faceted education and support programme on anxiety symptoms among people with systemic sclerosis and anxiety during COVID-19 (SPIN-CHAT): a two-arm parallel, partially nested, randomised, controlled trial. <i>Lancet Rheumatology</i> , The, 2021, 3, e427-e437.	2.2	24
20	Using Optimal Test Assembly Methods for Shortening Patient-Reported Outcome Measures: Development and Validation of the Cochin Hand Function Scale: A Scleroderma Patient-Centered Intervention Network Cohort Study. <i>Arthritis Care and Research</i> , 2016, 68, 1704-1713.	1.5	23
21	Exploring Sources of Emotional Distress among People Living with Scleroderma: A Focus Group Study. <i>PLoS ONE</i> , 2016, 11, e0152419.	1.1	22
22	Protocol for a partially nested randomised controlled trial to evaluate the effectiveness of the scleroderma patient-centered intervention network COVID-19 home-isolation activities together (SPIN-CHAT) program to reduce anxiety among at-risk scleroderma patients. <i>Journal of Psychosomatic Research</i> , 2020, 135, 110132.	1.2	21
23	Validation of the COVID-19 Fears Questionnaires for Chronic Medical Conditions: A Scleroderma Patient-centered Intervention Network COVID-19 Cohort study. <i>Journal of Psychosomatic Research</i> , 2020, 139, 110271.	1.2	20
24	Validation of the Social Appearance Anxiety Scale in Patients With Systemic Sclerosis: A Scleroderma Patient-Centered Intervention Network Cohort Study. <i>Arthritis Care and Research</i> , 2018, 70, 1557-1562.	1.5	17
25	Exercise habits and factors associated with exercise in systemic sclerosis: a Scleroderma Patient-centered Intervention Network (SPIN) cohort study. <i>Disability and Rehabilitation</i> , 2018, 40, 1997-2003.	0.9	17
26	Not as Stable as We Think: A Descriptive Study of 12 Monthly Assessments of Fear of Cancer Recurrence Among Curatively-Treated Breast Cancer Survivors 5 Years After Surgery. <i>Frontiers in Psychology</i> , 2020, 11, 580979.	1.1	16
27	Prevalence of posttraumatic stress disorder (PTSD) in women with breast cancer. <i>Journal of Psychosomatic Research</i> , 2014, 76, 485-486.	1.2	15
28	Cross-Language Measurement Equivalence of the Center for Epidemiologic Studies Depression (CES-D) Scale in Systemic Sclerosis: A Comparison of Canadian and Dutch Patients. <i>PLoS ONE</i> , 2013, 8, e53923.	1.1	14
29	Prospective longitudinal study on fear of cancer recurrence in patients newly diagnosed with head and neck cancer: Course, trajectories, and associated factors. <i>Head and Neck</i> , 2022, 44, 914-925.	0.9	14
30	The Comparability of English, French and Dutch Scores on the Functional Assessment of Chronic Illness Therapy-Fatigue (FACIT-F): An Assessment of Differential Item Functioning in Patients with Systemic Sclerosis. <i>PLoS ONE</i> , 2014, 9, e91979.	1.1	13
31	Reasons for Not Participating in Scleroderma Patient Support Groups: A Cross-Sectional Study. <i>Arthritis Care and Research</i> , 2018, 70, 275-283.	1.5	13
32	Shortening patient-reported outcome measures through optimal test assembly: application to the Social Appearance Anxiety Scale in the Scleroderma Patient-centered Intervention Network Cohort. <i>BMJ Open</i> , 2019, 9, e024010.	0.8	13
33	Reasons for attending support groups and organizational preferences: the European scleroderma support group members survey. <i>Disability and Rehabilitation</i> , 2019, 41, 974-982.	0.9	13
34	Development and Validation of the Body Concealment Scale for Scleroderma. <i>Arthritis Care and Research</i> , 2016, 68, 1158-1165.	1.5	11
35	Physical or Occupational Therapy Use in Systemic Sclerosis: A Scleroderma Patient-centered Intervention Network Cohort Study. <i>Journal of Rheumatology</i> , 2019, 46, 1605-1613.	1.0	11
36	Using Marital Status and Continuous Marital Satisfaction Ratings to Predict Depressive Symptoms in Married and Unmarried Women With Systemic Sclerosis: A Canadian Scleroderma Research Group Study. <i>Arthritis Care and Research</i> , 2016, 68, 1143-1149.	1.5	10

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37	Protocol for a scoping review to support development of a CONSORT extension for randomised controlled trials using cohorts and routinely collected health data. <i>BMJ Open</i> , 2018, 8, e025266.	0.8	10
38	The association of sociodemographic and objectively-assessed disease variables with fatigue in systemic sclerosis: an analysis of 785 Canadian Scleroderma Research Group Registry patients. <i>Clinical Rheumatology</i> , 2017, 36, 373-379.	1.0	9
39	Factors associated with fears due to COVID-19: A Scleroderma Patient-centered Intervention Network (SPIN) COVID-19 cohort study. <i>Journal of Psychosomatic Research</i> , 2021, 140, 110314.	1.2	9
40	Methods and results used in the development of a consensus-driven extension to the Consolidated Standards of Reporting Trials (CONSORT) statement for trials conducted using cohorts and routinely collected data (CONSORT-ROUTINE). <i>BMJ Open</i> , 2021, 11, e049093.	0.8	9
41	An Assessment of the Measurement Equivalence of English and French Versions of the Center for Epidemiologic Studies Depression (CES-D) Scale in Systemic Sclerosis. <i>PLoS ONE</i> , 2014, 9, e102897.	1.1	9
42	Pain levels and associated factors in the Scleroderma Patient-centered Intervention Network (SPIN) cohort: a multicentre cross-sectional study. <i>Lancet Rheumatology, The</i> , 2021, 3, e844-e854.	2.2	9
43	Mental health before and during the pandemic in people with systemic sclerosis. <i>Lancet Rheumatology, The</i> , 2022, 4, e82-e85.	2.2	9
44	Systemic sclerosis and COVID-19 vaccines: a SPIN Cohort study. <i>Lancet Rheumatology, The</i> , 2022, 4, e243-e246.	2.2	9
45	Health care use and patients' perceptions on quality of care in systemic sclerosis. <i>Clinical and Experimental Rheumatology</i> , 2013, 31, 64-70.	0.4	9
46	Comparison of Self-Efficacy for Managing Chronic Disease between patients with systemic sclerosis and other chronic conditions: a systematic review. <i>Rheumatology International</i> , 2017, 37, 281-292.	1.5	8
47	Reliability and Validity of Three Versions of the Brief Fear of Negative Evaluation Scale in Patients With Systemic Sclerosis: A Scleroderma Patient-centered Intervention Network Cohort Study. <i>Arthritis Care and Research</i> , 2018, 70, 1646-1652.	1.5	8
48	Understanding coping strategies among people living with scleroderma: a focus group study. <i>Disability and Rehabilitation</i> , 2018, 40, 3012-3021.	0.9	8
49	Protocol for a partially nested randomized controlled trial to evaluate the effectiveness of the Scleroderma Patient-centered Intervention Network Support Group Leader Education (SPIN-SSLED) Program. <i>Trials</i> , 2019, 20, 717.	0.7	8
50	Reporting transparency and completeness in Trials: Paper 2 - reporting of randomised trials using registries was often inadequate and hindered the interpretation of results. <i>Journal of Clinical Epidemiology</i> , 2022, 141, 175-186.	2.4	8
51	The patient engagement evaluation tool was valid for clinical practice guideline development. <i>Journal of Clinical Epidemiology</i> , 2022, 143, 61-72.	2.4	8
52	Randomized feasibility trial of the Scleroderma Patient-centered Intervention Network hand exercise program (SPIN-HAND): Study protocol. <i>Journal of Scleroderma and Related Disorders</i> , 2018, 3, 91-97.	1.0	7
53	Reasons for attending support groups and organizational preferences: A replication study using the North American Scleroderma Support Group Survey. <i>Journal of Scleroderma and Related Disorders</i> , 2019, 4, 173-186.	1.0	7
54	Reporting transparency and completeness in trials: Paper 4 - reporting of randomised controlled trials conducted using routinely collected electronic records "room for improvement. <i>Journal of Clinical Epidemiology</i> , 2022, 141, 198-209.	2.4	7

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55	The Scleroderma Patient-Centered Intervention Network Self-Management Program: Protocol for a Randomized Feasibility Trial. <i>JMIR Research Protocols</i> , 2020, 9, e16799.	0.5	7
56	Reporting transparency and completeness in trials: Paper 3 “ trials conducted using administrative databases do not adequately report elements related to use of databases. <i>Journal of Clinical Epidemiology</i> , 2022, 141, 187-197.	2.4	7
57	Reasons for not participating in scleroderma patient support groups: a comparison of results from the North American and European scleroderma support group surveys. <i>Disability and Rehabilitation</i> , 2021, 43, 1279-1286.	0.9	6
58	Addressing patient health care demands in systemic sclerosis: pre-post assessment of a psycho-educational group programme. <i>Clinical and Experimental Rheumatology</i> , 2011, 29, S60-5.	0.4	6
59	Can the Cancer-related Fatigue Case-definition Criteria Be Applied to Chronic Medical Illness? A Comparison between Breast Cancer and Systemic Sclerosis. <i>Journal of Rheumatology</i> , 2015, 42, 1156-1162.	1.0	5
60	Assessment of English-French differential item functioning of the Satisfaction with Appearance Scale (SWAP) in systemic sclerosis. <i>Body Image</i> , 2017, 22, 97-102.	1.9	5
61	Factors associated with patient-reported likelihood of using online self-care interventions: a Scleroderma Patient-centered Intervention Network (SPIN) cohort study. <i>BMJ Open</i> , 2019, 9, e029542.	0.8	5
62	Cognitive-Behavioural Therapy Targeting Fear of Progression in an Interdisciplinary Care Program: A Case Study in Systemic Sclerosis. <i>Journal of Clinical Psychology in Medical Settings</i> , 2014, 21, 297-312.	0.8	4
63	Factor structure and convergent validity of the Derriford Appearance Scale-24 using standard scoring versus treating “not applicable” responses as missing data: a Scleroderma Patient-centered Intervention Network (SPIN) cohort study. <i>BMJ Open</i> , 2018, 8, e018641.	0.8	4
64	Barriers and Facilitators to Physical Activity for People With Scleroderma: A Scleroderma Patient-Centered Intervention Network Cohort Study. <i>Arthritis Care and Research</i> , 2022, 74, 1300-1310.	1.5	4
65	A study protocol for the development of a SPIRIT extension for trials conducted using cohorts and routinely collected data (SPIRIT-ROUTINE). <i>HRB Open Research</i> , 2021, 4, 82.	0.3	4
66	The Scleroderma Patient-centered Intervention Network Self-Management (SPIN-SELF) Program: protocol for a two-arm parallel partially nested randomized controlled feasibility trial with progression to full-scale trial. <i>Trials</i> , 2021, 22, 856.	0.7	4
67	Validation of the Body Concealment Scale for Scleroderma (BCSS): Replication in the Scleroderma Patient-centered Intervention Network (SPIN) Cohort. <i>Body Image</i> , 2017, 20, 99-106.	1.9	3
68	Validation of the Social Interaction Anxiety Scale in scleroderma: a Scleroderma Patient-centered Intervention Network Cohort study. <i>Journal of Scleroderma and Related Disorders</i> , 2018, 3, 98-105.	1.0	3
69	Effects of non-pharmacological and non-surgical interventions on health outcomes in systemic sclerosis: protocol for a living systematic review. <i>BMJ Open</i> , 2021, 11, e047428.	0.8	3
70	Pain and Self-Efficacy Among Patients With Systemic Sclerosis. <i>Nursing Research</i> , 2021, 70, 334-343.	0.8	3
71	Psychological aspects in patients with advanced cancer receiving lifelong systemic treatment: protocol for a scoping review. <i>BMJ Open</i> , 2021, 11, e042404.	0.8	3
72	Assessing differential item functioning for the Social Appearance Anxiety Scale: a Scleroderma patient-centred Intervention Network (SPIN) Cohort Study. <i>BMJ Open</i> , 2020, 10, e037639.	0.8	3

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73	Examination of the association of sex and race/ethnicity with appearance concerns: a Scleroderma Patient-centered Intervention Network (SPIN) Cohort study. <i>Clinical and Experimental Rheumatology</i> , 2016, 34 Suppl 100, 92-99.	0.4	3
74	The association of sociodemographic and disease variables with hand function: a Scleroderma Patient-centered Intervention Network cohort study. <i>Clinical and Experimental Rheumatology</i> , 2018, 36 Suppl 113, 88-94.	0.4	3
75	Randomized feasibility trial of the Scleroderma Patient-centered Intervention Network Self-Management (SPIN-SELF) Program. <i>Pilot and Feasibility Studies</i> , 2022, 8, 45.	0.5	3
76	The Comparability of Functional Assessment of Chronic Illness Therapy - Fatigue Scores between Cancer and Systemic Sclerosis. <i>Journal of Scleroderma and Related Disorders</i> , 2017, 2, 57-63.	1.0	2
77	Development and preliminary validation of the Scleroderma Support Group Leader Self-efficacy Scale. <i>Journal of Scleroderma and Related Disorders</i> , 2018, 3, 106-111.	1.0	2
78	The Scleroderma Research Topics Survey for patients and health care professionals: A Scleroderma Patient-centered Intervention Network Project. <i>Journal of Scleroderma and Related Disorders</i> , 2019, 4, 165-172.	1.0	2
79	Trajectories of adjustment disorder symptoms in post-treatment breast cancer survivors. <i>Supportive Care in Cancer</i> , 2022, 30, 3521-3530.	1.0	2
80	Three-year trajectories of disability and fatigue in systemic sclerosis: a cohort study. <i>Clinical and Experimental Rheumatology</i> , 2017, 35 Suppl 106, 48-55.	0.4	2
81	Reassessing the clinical utility of the Patient Health Questionnaire (PHQ)-9 for depression screening in prenatal women: a commentary on Sidebottom et al.. <i>Archives of Women's Mental Health</i> , 2013, 16, 253-254.	1.2	1
82	Don't forget about the "in cmRCT: reply to Groenwold and van Smeden. <i>Journal of Clinical Epidemiology</i> , 2019, 106, 143-144.	2.4	1
83	Effects of cosmetic and other camouflage interventions on appearance-related and psychological outcomes among adults with visible differences in appearance: a systematic review. <i>BMJ Open</i> , 2021, 11, e046634.	0.8	1
84	Assessing differential item functioning for the Social Appearance Anxiety Scale: a Scleroderma patient-centred Intervention Network (SPIN) Cohort Study. <i>BMJ Open</i> , 2020, 10, e037639.	0.8	1
85	Performance of the Patient-Reported Outcomes Measurement Information System-29 in scleroderma: a Scleroderma Patient-centered Intervention Network Cohort Study. <i>Rheumatology</i> , 2019, , .	0.9	0
86	Validation of the "knowledge about melanoma early detection scale" in a sample of melanoma survivors. <i>Current Psychology</i> , 0, , 1.	1.7	0
87	Mental Health Care Use and Associated Factors in Systemic Sclerosis: A Scleroderma Patient-centered Intervention Network Cohort Study. <i>ACR Open Rheumatology</i> , 0, , .	0.9	0