

Linda Kwakkenbos

List of Publications by Year in descending order

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Version: 2024-02-01

87
papers

1,628
citations

393982

19
h-index

360668

35
g-index

89
all docs

89
docs citations

89
times ranked

2106
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 |

| # | ARTICLE | IF | CITATIONS |
|----|--|-----|-----------|
| 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-centered 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 |