Lotte Haverman

List of Publications by Year in descending order

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279487 276539 2,065 76 23 41 h-index citations g-index papers 81 81 81 2328 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Training clinicians in how to use patient-reported outcome measures in routine clinical practice. Quality of Life Research, 2015, 24, 1707-1718.	1.5	161
2	Using an implementation science approach to implement and evaluate patient-reported outcome measures (PROM) initiatives in routine care settings. Quality of Life Research, 2021, 30, 3015-3033.	1.5	127
3	The impact of lockdown during the COVID-19 pandemic on mental and social health of children and adolescents. Quality of Life Research, 2021, 30, 2795-2804.	1.5	124
4	Development and Validation of the Distress Thermometer for Parents of a Chronically Ill Child. Journal of Pediatrics, 2013, 163, 1140-1146.e2.	0.9	107
5	Innovations in e-health. Quality of Life Research, 2014, 23, 195-203.	1.5	105
6	Monitoring health-related quality of life in paediatric practice: development of an innovative web-based application. BMC Pediatrics, 2011, 11, 3.	0.7	102
7	Effectiveness of a Web-Based Application to Monitor Health-Related Quality of Life. Pediatrics, 2013, 131, e533-e543.	1.0	90
8	Implementation of electronic patient reported outcomes in pediatric daily clinical practice: The KLIK experience Clinical Practice in Pediatric Psychology, 2014, 2, 50-67.	0.2	69
9	Real-world implementation of electronic patient-reported outcomes in outpatient pediatric cancer care. Psycho-Oncology, 2017, 26, 951-959.	1.0	61
10	Patient-Reported Outcome Measures in Routine Pediatric Clinical Care: A Systematic Review. Frontiers in Pediatrics, 2020, 8, 364.	0.9	55
11	Health related quality of life in Dutch young adults: psychometric properties of the PedsQL generic core scales young adult version. Health and Quality of Life Outcomes, 2014, 12, 9.	1.0	52
12	Development and implementation of a patient reported outcome intervention (QLIC-ON PROfile) in clinical paediatric oncology practice. Patient Education and Counseling, 2010, 81, 235-244.	1.0	51
13	Healthcare Professionals' Preferences and Perceived Barriers for Routine Assessment of Patientâ€Reported Outcomes in Pediatric Oncology Practice: Moving Toward International Processes of Change. Pediatric Blood and Cancer, 2016, 63, 2181-2188.	0.8	47
14	Children on dialysis as well as renal transplanted children report severely impaired health-related quality of life. Quality of Life Research, 2018, 27, 1445-1454.	1.5	46
15	Parental distress 6 months after a pediatric cancer diagnosis in relation to family psychosocial risk at diagnosis. Cancer, 2018, 124, 381-390.	2.0	45
16	Dutch normative data and psychometric properties for the Distress Thermometer for Parents. Quality of Life Research, 2017, 26, 177-182.	1.5	44
17	Health-related quality of life in children with newly diagnosed immune thrombocytopenia. Haematologica, 2014, 99, 1525-1531.	1.7	34
18	Parental Distress and Quality of Life in Pediatric Inflammatory Bowel Disease. Journal of Pediatric Gastroenterology and Nutrition, 2018, 66, 630-636.	0.9	34

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19	Health related quality of life of infants and children with avoidant restrictive food intake disorder. International Journal of Eating Disorders, 2019, 52, 410-418.	2.1	31
20	Mental and Social Health of Children and Adolescents With Pre-existing Mental or Somatic Problems During the COVID-19 Pandemic Lockdown. Frontiers in Psychiatry, 2021, 12, 692853.	1.3	29
21	Dutch–Flemish translation of nine pediatric item banks from the Patient-Reported Outcomes Measurement Information System (PROMIS)®. Quality of Life Research, 2016, 25, 761-765.	1.5	28
22	Attitudes of the general population towards preconception expanded carrier screening for autosomal recessive disorders including inborn errors of metabolism. Molecular Genetics and Metabolism, 2019, 126, 14-22.	0.5	28
23	Options for the Interpretation of and Recommendations for Acting on Different PROMs in Daily Clinical Practice Using KLIK. Medical Care, 2019, 57, S52-S58.	1.1	26
24	Health-related quality of life, anxiety, depression and distress of mothers and fathers of children on Home parenteral nutrition. Clinical Nutrition, 2019, 38, 1905-1912.	2.3	25
25	Psychosocial Functioning in Parents of MPS III Patients. JIMD Reports, 2018, 44, 33-41.	0.7	23
26	Health related quality of life and parental perceptions of child vulnerability among parents of a child with juvenile idiopathic arthritis: results from a web-based survey. Pediatric Rheumatology, 2014, 12, 34.	0.9	22
27	A retrospective assessment of the KLIK PROM portal implementation using the Consolidated Framework for Implementation Research (CFIR). Quality of Life Research, 2021, 30, 3049-3061.	1.5	21
28	Common patient-reported outcomes across ICHOM Standard Sets: the potential contribution of PROMIS®. BMC Medical Informatics and Decision Making, 2021, 21, 259.	1.5	21
29	COVID-19 lockdown impacts the wellbeing of parents with infants on a Dutch neonatal intensive care unit. Journal of Pediatric Nursing, 2022, 62, 106-112.	0.7	20
30	Paediatric patients report lower healthâ€related quality of life in daily clinical practice compared to new normative PedsQL TM data. Acta Paediatrica, International Journal of Paediatrics, 2021, 110, 2267-2279.	0.7	18
31	Quality of Life and Behavioral Functioning in Dutch Children with a History of Kawasaki Disease. Journal of Pediatrics, 2012, 161, 314-319.e1.	0.9	17
32	Psychometric properties of the Patient-Reported Outcomes Measurement Information System (PROMIS®) pediatric item bank peer relationships in the Dutch general population. Quality of Life Research, 2021, 30, 2061-2070.	1.5	17
33	Health-related quality of life in perinatally HIV-infected children in the Netherlands. AIDS Care - Psychological and Socio-Medical Aspects of AIDS/HIV, 2015, 27, 1279-1288.	0.6	16
34	Prevention of relapses with levamisole as adjuvant therapy in children with a first episode of idiopathic nephrotic syndrome: study protocol for a double blind, randomised placebo-controlled trial (the LEARNS study). BMJ Open, 2019, 9, e027011.	0.8	16
35	Psychometric Properties of the Pediatric Patientâ€Reported Outcomes Measurement Information System Item Banks in a Dutch Clinical Sample of Children With Juvenile Idiopathic Arthritis. Arthritis Care and Research, 2020, 72, 1780-1789.	1.5	16
36	Clinicians' perspective on the implemented KLIK PROM portal in clinical practice. Quality of Life Research, 2021, 30, 3267-3277.	1.5	16

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37	Validation of PROMIS Profileâ€29 in adults with hemophilia in the Netherlands. Journal of Thrombosis and Haemostasis, 2021, 19, 2687-2701.	1.9	16
38	Health-related quality of life and psychosocial developmental trajectory in young female beneficiaries with JIA. Rheumatology, 2012, 51, 368-374.	0.9	15
39	Children of non-Western origin with end-stage renal disease in the Netherlands, Belgium and a part of Germany have impaired health-related quality of life compared with Western children. Nephrology Dialysis Transplantation, 2014, 29, 448-457.	0.4	15
40	Distress and everyday problems in Dutch mothers and fathers of young adolescents with Down syndrome. Research in Developmental Disabilities, 2017, 67, 19-27.	1.2	15
41	Use of Patient-Reported Experience Measures in Pediatric Care: A Systematic Review. Frontiers in Pediatrics, 2021, 9, 753536.	0.9	15
42	Psychometric properties and Dutch norm data of the PedsQL Multidimensional Fatigue Scale for Young Adults. Quality of Life Research, 2014, 23, 2841-2847.	1.5	14
43	Healthâ€related quality of life in children with Robin sequence. American Journal of Medical Genetics, Part A, 2017, 173, 54-61.	0.7	14
44	Internalizing problems before and during the COVID-19 pandemic in independent samples of Dutch children and adolescents with and without pre-existing mental health problems. European Child and Adolescent Psychiatry, 2023, 32, 1873-1883.	2.8	13
45	Health related quality of life and perceptions of child vulnerability among parents of children with a history of Kawasaki disease. Acta Paediatrica, International Journal of Paediatrics, 2014, 103, 671-677.	0.7	12
46	From statistics to clinics: the visual feedback of PROMIS® CATs. Journal of Patient-Reported Outcomes, 2021, 5, 55.	0.9	12
47	Impact of using patient-reported outcome measures in routine clinical care of paediatric patients with chronic conditions: a systematic review protocol. BMJ Open, 2019, 9, e027354.	0.8	11
48	Preconception expanded carrier screening: Impact of information presented by text or video on genetic knowledge and attitudes. Journal of Genetic Counseling, 2021, 30, 457-469.	0.9	11
49	Managing womenâ€specific bleeding in inherited bleeding disorders: A multidisciplinary approach. Haemophilia, 2021, 27, 463-469.	1.0	10
50	Patients' and parents' perspective on the implementation of Patient Reported Outcome Measures in pediatric clinical practice using the KLIK PROM portal. Quality of Life Research, 2022, 31, 241-254.	1.5	10
51	Psychometric properties and CAT performance of the PROMIS pediatric sleep disturbance, sleep-related impairment, and fatigue item banks in Dutch children and adolescents Psychological Assessment, 2022, 34, 860-869.	1.2	9
52	Parenting a child with Marfan syndrome: Distress and everyday problems. American Journal of Medical Genetics, Part A, 2021, 185, 50-59.	0.7	8
53	Generic PROMIS item banks in adults with hemophilia for patientâ€reported outcome assessment: Feasibility, measurement properties, and relevance. Research and Practice in Thrombosis and Haemostasis, 2021, 5, e12621.	1.0	8
54	Matching Psychosocial Support Needs of Parents of a Child with a Chronic Illness to a Feasible Intervention. Maternal and Child Health Journal, 2020, 24, 1238-1247.	0.7	7

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55	Patient-reported outcome measures in pediatric solid organ transplantation: Exploring stakeholder perspectives on clinical implementation through qualitative description. Quality of Life Research, 2021, 30, 1355-1364.	1.5	7
56	Advances in measuring pediatric overall health: the PROMIS® Pediatric Global Health scale (PGH-7). European Journal of Pediatrics, 2022, 181, 2117-2125.	1.3	7
57	Views of patients and parents of children with genetic disorders on populationâ€based expanded carrier screening. Prenatal Diagnosis, 0, , .	1.1	7
58	Attitudes of relatives of mucopolysaccharidosis type III patients toward preconception expanded carrier screening. European Journal of Human Genetics, 2020, 28, 1331-1340.	1.4	6
59	Preconception expanded carrier screening: a focus group study with relatives of mucopolysaccharidosis type III patients and the general population. Journal of Community Genetics, 2021, 12, 311-323.	0.5	6
60	Psychometric properties of the Patient-Reported Outcomes Measurement Information System (PROMIS®) Pediatric Anger Scale in the Dutch general population Psychological Assessment, 2021, 33, 1261-1266.	1.2	6
61	Prevalence and determinants of persistent symptoms after infection with SARS-CoV-2: protocol for an observational cohort study (LongCOVID-study). BMJ Open, 2022, 12, e062439.	0.8	6
62	Pediatric Perceived Cognitive Functioning: Psychometric Properties and Normative Data of the Dutch Item Bank and Short Form. Journal of the International Neuropsychological Society, 2019, 25, 845-856.	1.2	5
63	Healthâ€Related Quality of Life and Distress of Parents of Children With Avoidant Restrictive Food Intake Disorder. Journal of Pediatric Gastroenterology and Nutrition, 2021, 73, 115-124.	0.9	5
64	Longitudinal Development of Healthâ€related Quality of Life and Fatigue in Children on Home Parenteral Nutrition. Journal of Pediatric Gastroenterology and Nutrition, 2022, 74, 116-122.	0.9	5
65	Professional functioning of young adults with congenital coagulation disorders in the Netherlands. Haemophilia, 2019, 25, e138-e145.	1.0	4
66	Reducing posttraumatic stress in parents of patients with a rare inherited metabolic disorder using eye movement desensitization and reprocessing therapy: a case study. Orphanet Journal of Rare Diseases, 2021, 16, 126.	1.2	4
67	A qualitative study on the experiences of haemophilia carriers before, during and after pregnancy. Haemophilia, 2021, 27, e675-e682.	1.0	4
68	The Use of Patient-Reported Outcome Measures in Daily Clinical Practice of a Pediatric Nephrology Department. International Journal of Environmental Research and Public Health, 2022, 19, 5338.	1.2	4
69	Including the voice of paediatric patients: Cocreation of an engagement game. Health Expectations, 2022, 25, 1861-1871.	1.1	4
70	Illness cognitions associated with healthâ€related quality of life in young adult men with haemophilia. Haemophilia, 2020, 26, 793-799.	1.0	2
71	How to use the patient-reported outcomes measurement information system (PROMIS) in international orthopaedic trauma research: a practical approach. Injury, 2021, 52, 3537-3539.	0.7	1
72	The sensory-reactivity PROM set: identification of a parent reported outcome measure set for autism spectrum disorder. Journal of Patient-Reported Outcomes, 2021, 5, 123.	0.9	1

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73	Linking the RCADS-25 to the PROMIS® pediatric item banks Anxiety and Depressive Symptoms in a general Dutch population sample. Quality of Life Research, 2022, 31, 1587-1595.	1.5	1
74	Perceived stress, family impact, and changes in physical and social daily life activities of children with chronic somatic conditions during the COVID-19 pandemic. BMC Public Health, 2022, 22, .	1.2	1
75	Stap 9: hoe ga je de meetresultaten gebruiken?. , 2020, , 129-147.		O
76	One-Stop Surgery: An Innovation to Limit Hospital Visits in Children. European Journal of Pediatric Surgery, 2021, , .	0.7	0