

# Caroline Sanders

## List of Publications by Year in descending order

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115  
papers

5,103  
citations

101543  
36  
h-index

102487  
66  
g-index

121  
all docs

121  
docs citations

121  
times ranked

7014  
citing authors

#	ARTICLE	IF	CITATIONS
1	Primary Care Providers Involvement in Caring for Young Adults with Complex Chronic Conditions Exiting Pediatric Care: An Integrative Literature Review. Comprehensive Child and Adolescent Nursing, 2023, 46, 201-222.	0.9	4
2	The perceptions of general practice among Central and Eastern Europeans in the United Kingdom: A systematic scoping review. Health Expectations, 2022, , .	2.6	4
3	Remote primary care during the COVID-19 pandemic for people experiencing homelessness: a qualitative study. British Journal of General Practice, 2022, 72, e492-e500.	1.4	16
4	Ethnic inequalities in COVID-19 vaccine uptake and comparison to seasonal influenza vaccine uptake in Greater Manchester, UK: A cohort study. PLoS Medicine, 2022, 19, e1003932.	8.4	30
5	Lifespan healthcare transitions among individuals with intersex traits in Canada: a mixed-methods and qualitative study. BMJ Open, 2022, 12, e055759.	1.9	4
6	Pediatric practicums in undergraduate nursing programs: an integrative review. International Journal of Nursing Education Scholarship, 2022, 19, .	1.0	1
7	Exploring engagement with digital screens for collecting patient feedback in clinical waiting rooms: The role of touch and place. Health (United Kingdom), 2021, 25, 454-474.	1.5	5
8	Exploring the experiences of changes to support access to primary health care services and the impact on the quality and safety of care for homeless people during the COVID-19 pandemic: a study protocol for a qualitative mixed methods approach. International Journal for Equity in Health, 2021, 20, 29.	3.5	17
9	Protocol for a non-randomised feasibility study evaluating a codesigned patient safety guide in primary care. BMJ Open, 2021, 11, e039752.	1.9	1
10	<i>On the borderline of diabetes</i>: understanding how individuals resist and reframe diabetes risk. Health, Risk and Society, 2021, 23, 34-51.	1.7	16
11	Public Perspectives of Using Social Media Data to Improve Adverse Drug Reaction Reporting: A Mixed-Methods Study. Drug Safety, 2021, 44, 553-564.	3.2	8
12	Childrenâ€™s Independent Mobility and Physical Activity during the COVID-19 Pandemic: A Qualitative Study with Families. International Journal of Environmental Research and Public Health, 2021, 18, 4481.	2.6	36
13	Co-designing an Adaption of a Mobile App to Enhance Communication, Safety, and Well-being Among People Living at Home With Early-Stage Dementia: Protocol for an Exploratory Multiple Case Study. JMIR Research Protocols, 2021, 10, e19543.	1.0	3
14	Co-designing new tools for collecting, analysing and presenting patient experience data in NHS services: working in partnership with patients and carers. Research Involvement and Engagement, 2021, 7, 85.	2.9	5
15	Understanding the implementation, impact and sustainable use of an electronic pharmacy referral service at hospital discharge: A qualitative evaluation from a sociotechnical perspective.. PLoS ONE, 2021, 16, e0261153.	2.5	3
16	Providing â€˜the bigger pictureâ€™: benefits and feasibility of integrating remote monitoring from smartphones into the electronic health record. Rheumatology, 2020, 59, 367-378.	1.9	75
17	Implementing a digital patient feedback system: an analysis using normalisation process theory. BMC Health Services Research, 2020, 20, 387.	2.2	11
18	Patient safety in marginalised groups: a narrative scoping review. International Journal for Equity in Health, 2020, 19, 26.	3.5	38

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19	Engagement and Participant Experiences With Consumer Smartwatches for Health Research: Longitudinal, Observational Feasibility Study. JMIR MHealth and UHealth, 2020, 8, e14368.	3.7	43
20	Smartphone-Enhanced Symptom Management In Psychosis: Open, Randomized Controlled Trial. Journal of Medical Internet Research, 2020, 22, e17019.	4.3	25
21	Digital methods to enhance the usefulness of patient experience data in services for long-term conditions: the DEPEND mixed-methods study. Health Services and Delivery Research, 2020, 8, 1-128.	1.4	3
22	Dementia and mild cognitive impairment in prisoners aged over 50 years in England and Wales: a mixed-methods study. Health Services and Delivery Research, 2020, 8, 1-116.	1.4	14
23	Evaluating a telehealth intervention for urinalysis monitoring in children with neurogenic bladder. Journal of Child Health Care, 2019, 23, 45-62.	1.4	9
24	Understanding the diagnosis of pre-diabetes in patients aged over 85 in English primary care: a qualitative study. BMC Family Practice, 2019, 20, 90.	2.9	12
25	How the weather affects the pain of citizen scientists using a smartphone app. Npj Digital Medicine, 2019, 2, 105.	10.9	49
26	Canadian Pediatric Populations and Specific Challenges. Comprehensive Child and Adolescent Nursing, 2019, 42, 167-171.	0.9	0
27	Collecting Symptoms and Sensor Data With Consumer Smartwatches (the Knee OsteoArthritis, Linking) Tj ETQq1 1 0.784314 rgBT /Ov Protocols, 2019, 8, e10238.	1.0	18
28	Training to enhance user and carer involvement in mental health-care planning: the EQUIP research programme including a cluster RCT. Programme Grants for Applied Research, 2019, 7, 1-140.	1.0	8
29	Empowering people to help speak up about safety in primary care: Using codesign to involve patients and professionals in developing new interventions for patients with multimorbidity. Health Expectations, 2018, 21, 539-548.	2.6	38
30	Is it time to abandon care planning in mental health services? A qualitative study exploring the views of professionals, service users and carers. Health Expectations, 2018, 21, 597-605.	2.6	28
31	A core outcome set for clinical trials of chemoradiotherapy interventions for anal cancer (CORMAC): a patient and health-care professional consensus. The Lancet Gastroenterology and Hepatology, 2018, 3, 865-873.	8.1	51
32	Exploring implementation of an electronic referral management system and enhanced primary care service for oral surgery: perspectives of patients, providers and practitioners. BMC Health Services Research, 2018, 18, 646.	2.2	5
33	Implementing and evaluating a primary care service for oral surgery: a case study. BMC Health Services Research, 2018, 18, 636.	2.2	7
34	Embedding shared decision-making in the care of patients with severe and enduring mental health problems: The EQUIP pragmatic cluster randomised trial. PLoS ONE, 2018, 13, e0201533.	2.5	33
35	Critical moments in long-term condition management: A longitudinal qualitative social network study. Chronic Illness, 2018, 14, 119-134.	1.5	5
36	Enacting personâ€centredness in integrated care: A qualitative study of practice and perspectives within multidisciplinary groups in the care of older people. Health Expectations, 2018, 21, 1066-1074.	2.6	10

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37	An evaluation of a referral management and triage system for oral surgery referrals from primary care dentists: a mixed-methods study. <i>Health Services and Delivery Research</i> , 2018, 6, 1-126.	1.4	11
38	Improving care for older people with long-term conditions and social care needs in Salford: the CLASSIC mixed-methods study, including RCT. <i>Health Services and Delivery Research</i> , 2018, 6, 1-188.	1.4	9
39	Quantifying beliefs regarding telehealth: Development of the Whole Systems Demonstrator Service User Technology Acceptability Questionnaire. <i>Journal of Telemedicine and Telecare</i> , 2017, 23, 460-469.	2.7	91
40	Parent-to-parent peer support for parents of children with a disability: A mixed method study. <i>Patient Education and Counseling</i> , 2017, 100, 1537-1543.	2.2	71
41	An evaluation study examining penile bacterial flora in boys undergoing hypospadias surgery with foreskin reconstruction. <i>International Journal of Urological Nursing</i> , 2017, 11, 67-72.	0.2	1
42	Accessing support and empowerment online: The experiences of individuals with diabetes. <i>Health Expectations</i> , 2017, 20, 1088-1095.	2.6	32
43	Core outcome research measures in anal cancer (CORMAC): protocol for systematic review, qualitative interviews and Delphi survey to develop a core outcome set in anal cancer. <i>BMJ Open</i> , 2017, 7, e018726.	1.9	24
44	A narrative systematic review of factors affecting diabetes prevention in primary care settings. <i>PLoS ONE</i> , 2017, 12, e0177699.	2.5	39
45	Threats to patient safety in primary care reported by older people with multimorbidity: baseline findings from a longitudinal qualitative study and implications for intervention. <i>BMC Health Services Research</i> , 2017, 17, 754.	2.2	24
46	Tea, talk and technology: patient and public involvement to improve connected health “wearables” research in dementia. <i>Research Involvement and Engagement</i> , 2017, 3, 12.	2.9	38
47	Seeking to understand lived experiences of personal recovery in personality disorder in community and forensic settings “ a qualitative methods investigation. <i>BMC Psychiatry</i> , 2017, 17, 282.	2.6	11
48	Cloudy with a Chance of Pain: Engagement and Subsequent Attrition of Daily Data Entry in a Smartphone Pilot Study Tracking Weather, Disease Severity, and Physical Activity in Patients With Rheumatoid Arthritis. <i>JMIR MHealth and UHealth</i> , 2017, 5, e37.	3.7	60
49	Recruitment and Ongoing Engagement in a UK Smartphone Study Examining the Association Between Weather and Pain: Cohort Study. <i>JMIR MHealth and UHealth</i> , 2017, 5, e168.	3.7	41
50	Personal recovery within forensic settings “ Systematic review and meta-synthesis of qualitative methods studies. <i>Criminal Behaviour and Mental Health</i> , 2016, 26, 59-75.	0.8	53
51	Trust, temporality and systems: how do patients understand patient safety in primary care? A qualitative study. <i>Health Expectations</i> , 2016, 19, 253-263.	2.6	53
52	Sensemaking and the co-production of safety: a qualitative study of primary medical care patients. <i>Sociology of Health and Illness</i> , 2016, 38, 270-285.	2.1	35
53	Evolving “self”-management: exploring the role of social network typologies on individual long-term condition management. <i>Health Expectations</i> , 2016, 19, 1044-1061.	2.6	24
54	Hidden caring, hidden carers? Exploring the experience of carers for people with long-term conditions. <i>Health and Social Care in the Community</i> , 2016, 24, 203-213.	1.6	52

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55	A case of tightrope walking. <i>International Journal of Workplace Health Management</i> , 2016, 9, 238-250.	1.9	13
56	“You get to know the people and whether they’re talking sense or not” Negotiating trust on health-related forums. <i>Social Science and Medicine</i> , 2016, 162, 151-157.	3.8	27
57	“Knowing the Places of Care” How Nurses Facilitate Transition of Children with Complex Health Care Needs from Hospital to Home. <i>Comprehensive Child and Adolescent Nursing</i> , 2016, 39, 139-153.	0.9	8
58	Personal recovery in personality disorder: Systematic review and meta-synthesis of qualitative methods studies. <i>International Journal of Social Psychiatry</i> , 2016, 62, 41-50.	3.1	16
59	Patient Perspectives on Sharing Anonymized Personal Health Data Using a Digital System for Dynamic Consent and Research Feedback: A Qualitative Study. <i>Journal of Medical Internet Research</i> , 2016, 18, e66.	4.3	139
60	“I Always Vet Things” Navigating Privacy and the Presentation of Self on Health Discussion Boards Among Individuals with Long-Term Conditions. <i>Journal of Medical Internet Research</i> , 2016, 18, e274.	4.3	25
61	A cluster randomised controlled trial and process evaluation of a training programme for mental health professionals to enhance user involvement in care planning in service users with severe mental health issues (EQUIP): study protocol for a randomised controlled trial. <i>Trials</i> , 2015, 16, 348.	1.6	20
62	Re-inventing care planning in mental health: stakeholder accounts of the imagined implementation of a user/carer involved intervention. <i>BMC Health Services Research</i> , 2015, 15, 490.	2.2	20
63	Blame the Patient, Blame the Doctor or Blame the System? A Meta-Synthesis of Qualitative Studies of Patient Safety in Primary Care. <i>PLoS ONE</i> , 2015, 10, e0128329.	2.5	69
64	The Contradictions of Telehealth User Experience in Chronic Obstructive Pulmonary Disease (COPD): A Qualitative Meta-Synthesis. <i>PLoS ONE</i> , 2015, 10, e0139561.	2.5	82
65	A study of the provision of hospital based dental General Anaesthetic services for children in the North West of England: Part 2 - the views and experience of families and dentists regarding service needs, treatment and prevention. <i>BMC Oral Health</i> , 2015, 15, 47.	2.3	14
66	Perceptions of recovery and prognosis from long-term conditions: The relevance of hope and imagined futures. <i>Chronic Illness</i> , 2015, 11, 3-20.	1.5	28
67	Connecting local support: A qualitative study exploring the role of voluntary organisations in long-term condition management. <i>Chronic Illness</i> , 2015, 11, 140-155.	1.5	5
68	Issues arising following a referral and subsequent wait for extraction under general anaesthetic: impact on children. <i>BMC Oral Health</i> , 2015, 15, 3.	2.3	48
69	Using social media for support and feedback by mental health service users: thematic analysis of a twitter conversation. <i>BMC Psychiatry</i> , 2015, 15, 29.	2.6	92
70	A study of the provision of hospital based dental general anaesthetic services for children in the northwest of England: part 1 - a comparison of service delivery between six hospitals. <i>BMC Oral Health</i> , 2015, 15, 50.	2.3	25
71	“You don't get told anything, they don't do anything and nothing changes”™. Medicine as a resource and constraint in progressive ataxia. <i>Health Expectations</i> , 2015, 18, 177-187.	2.6	13
72	Dynamic Consent: A Possible Solution to Improve Patient Confidence and Trust in How Electronic Patient Records Are Used in Medical Research. <i>JMIR Medical Informatics</i> , 2015, 3, e3.	2.6	95

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73	The effect of telecare on the quality of life and psychological well-being of elderly recipients of social care over a 12-month period: the Whole Systems Demonstrator cluster randomised trial. <i>Age and Ageing</i> , 2014, 43, 334-341.	1.6	58
74	Social status and living with a chronic illness: An exploration of assessment and meaning attributed to work and employment. <i>Chronic Illness</i> , 2014, 10, 273-290.	1.5	26
75	The mediation of social influences on smoking cessation and awareness of the early signs of lung cancer. <i>BMC Public Health</i> , 2014, 14, 1043.	2.9	9
76	How potentially serious symptom changes are talked about and managed in COPD clinical review consultations: A micro-analysis. <i>Social Science and Medicine</i> , 2014, 113, 120-136.	3.8	7
77	Relationship continuity: when and why do primary care patients think it is safer?. <i>British Journal of General Practice</i> , 2014, 64, e758-e764.	1.4	36
78	A dynamic model of patient consent to sharing of medical record data. <i>BMJ, The</i> , 2014, 348, g1294-g1294.	6.0	19
79	A Constellation of Misfortune. <i>SAGE Open</i> , 2014, 4, 215824401455804.	1.7	3
80	Experiences of front-line health professionals in the delivery of telehealth: a qualitative study. <i>British Journal of General Practice</i> , 2014, 64, e401-e407.	1.4	84
81	Qualitative Meta-Synthesis of User Experience of Computerised Therapy for Depression and Anxiety. <i>PLoS ONE</i> , 2014, 9, e84323.	2.5	148
82	Implementation of self management support for long term conditions in routine primary care settings: cluster randomised controlled trial. <i>BMJ, The</i> , 2013, 346, f2882-f2882.	6.0	195
83	Effect of telehealth on quality of life and psychological outcomes over 12 months (Whole Systems) Tj ETQq1 1 0.784314 rgBT /Overlocl pragmatic, cluster randomised controlled trial. <i>BMJ, The</i> , 2013, 346, f653-f653.	6.0	191
84	The influence of social factors on help-seeking for people with lung cancer. <i>European Journal of Cancer Care</i> , 2013, 22, 709-713.	1.5	34
85	What influences withdrawal because of rejection of telehealth â€” the whole systems demonstrator evaluation. <i>Journal of Assistive Technologies</i> , 2013, 7, 219-227.	0.8	20
86	Trouble with ataxia: A longitudinal qualitative study of the diagnosis and medical management of a group of rare, progressive neurological conditions. <i>SAGE Open Medicine</i> , 2013, 1, 205031211350556.	1.8	11
87	Social Networks, the â€”Workâ€”™ and Work Force of Chronic Illness Self-Management: A Survey Analysis of Personal Communities. <i>PLoS ONE</i> , 2013, 8, e59723.	2.5	132
88	Exploring barriers to participation and adoption of telehealth and telecare within the Whole System Demonstrator trial: a qualitative study. <i>BMC Health Services Research</i> , 2012, 12, 220.	2.2	358
89	An organisational analysis of the implementation of telecare and telehealth: the whole systems demonstrator. <i>BMC Health Services Research</i> , 2012, 12, 403.	2.2	104
90	Changes to financial incentives in <sc>English dentistry 2006â€”2009: a qualitative study. <i>Community Dentistry and Oral Epidemiology</i> , 2012, 40, 468-473.	1.9	26

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91	Experience of contractual change in UK general practice: a qualitative study of salaried GPs. British Journal of General Practice, 2012, 62, e282-e287.	1.4	7
92	A cluster randomised controlled trial of the clinical and cost-effectiveness of a 'whole systems' model of self-management support for the management of long-term conditions in primary care: trial protocol. Implementation Science, 2012, 7, 7.	6.9	51
93	The role of information in supporting self-care in vascular conditions: a conceptual and empirical review. Health and Social Care in the Community, 2011, 19, 449-459.	1.6	24
94	A comprehensive evaluation of the impact of telemonitoring in patients with long-term conditions and social care needs: protocol for the whole systems demonstrator cluster randomised trial. BMC Health Services Research, 2011, 11, 184.	2.2	105
95	Social networks, work and network-based resources for the management of long-term conditions: a framework and study protocol for developing self-care support. Implementation Science, 2011, 6, 56.	6.9	106
96	Getting a diagnosis v. learning to live with it? The case of the progressive ataxias. Chronic Illness, 2011, 7, 120-133.	1.5	10
97	Social networks, social capital and chronic illness self-management: a realist review. Chronic Illness, 2011, 7, 60-86.	1.5	167
98	Shifting priorities in multimorbidity: a longitudinal qualitative study of patients' prioritization of multiple conditions. Chronic Illness, 2011, 7, 147-161.	1.5	153
99	Managing 'difficult emotions' and family life: exploring insights and social support within online self-management training. Chronic Illness, 2011, 7, 134-146.	1.5	18
100	Bodies in Context: Potential Avenues of Inquiry for the Sociology of Chronic Illness and Disability Within a New Policy Era. Handbooks of Sociology and Social Research, 2011, , 483-504.	0.1	11
101	Professional status in a changing world: The case of medicines use reviews in English community pharmacy. Social Science and Medicine, 2010, 71, 451-458.	3.8	73
102	Unstated factors in orthopaedic decision-making: a qualitative study. BMC Musculoskeletal Disorders, 2010, 11, 213.	1.9	36
103	Routes to total joint replacement surgery: Patients' and clinicians' perceptions of need. Arthritis Care and Research, 2010, 62, 1252-1257.	3.4	48
104	Evaluating a peer education programme for advance end-of-life care planning for older adults: The peer educators' perspective. International Journal on Disability and Human Development, 2009, 8, .	0.2	10
105	Are some more equal than others? Social comparison in self-management skills training for long-term conditions. Chronic Illness, 2009, 5, 305-317.	1.5	27
106	Creating 'good' self-managers?: Facilitating and governing an online self care skills training course. BMC Health Services Research, 2009, 9, 93.	2.2	7
107	Planning for end of life care within lay-led chronic illness self-management training: The significance of 'death awareness' and biographical context in participant accounts. Social Science and Medicine, 2008, 66, 982-993.	3.8	40
108	Theorising inequalities in the experience and management of chronic illness: Bringing social networks and social capital back in (critically). Research in the Sociology of Health Care, 2007, , 15-42.	0.1	8

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109	Re-thinking the relationship between long-term condition self-management education and the utilisation of health services. <i>Social Science and Medicine</i> , 2007, 65, 934-945.	3.8	102
110	Development of a peer education programme for advance end-of-life care planning. <i>International Journal of Palliative Nursing</i> , 2006, 12, 216-223.	0.5	37
111	Unmet need for joint replacement: a qualitative investigation of barriers to treatment among individuals with severe pain and disability of the hip and knee. <i>British Journal of Rheumatology</i> , 2003, 43, 353-357.	2.3	82
112	The significance and consequences of having painful and disabled joints in older age: co-existing accounts of normal and disrupted biographies. <i>Sociology of Health and Illness</i> , 2002, 24, 227-253.	2.1	209
113	Reporting on quality of life in randomised controlled trials: bibliographic study. <i>BMJ: British Medical Journal</i> , 1998, 317, 1191-1194.	2.3	259
114	A review of menstrual sex education and management in women with congenital adrenal hyperplasia. <i>Sex Education</i> , 0, , 1-17.	2.0	3
115	An integrative literature review of menstruation patterns in people with congenital adrenal hyperplasia. <i>Canadian Journal of Human Sexuality</i> , 0, , .	1.6	0