Rachel M Taylor,, Dipres, Rscn, Rgn

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/2803279/publications.pdf

Version: 2024-02-01

66 papers

1,845 citations

23 h-index 40 g-index

69 all docs 69 docs citations

69 times ranked

2212 citing authors

#	Article	IF	Citations
1	Impact of leadership on the nursing workforce during the COVID-19 pandemic. BMJ Leader, 2023, 7, 21-27.	0.8	4
2	Virtual care and the impact of COVIDâ€19 on nursing: A single centre evaluation. Journal of Advanced Nursing, 2022, 78, 498-509.	1.5	22
3	The impact of COVID-19 on the wellbeing of the UK nursing and midwifery workforce during the first pandemic wave: A longitudinal survey study. International Journal of Nursing Studies, 2022, 127, 104155.	2.5	60
4	Associations between diagnostic time intervals and health-related quality of life, clinical anxiety and depression in adolescents and young adults with cancer: cross-sectional analysis of the BRIGHTLIGHT cohort. British Journal of Cancer, 2022, 126, 1725-1734.	2.9	10
5	Developing, Nurturing, and Sustaining an Adolescent and Young Adult-Centered Culture of Care. Qualitative Health Research, 2022, 32, 956-969.	1.0	2
6	Care coordination, consistency and continuity: the case of the key worker role in children's cancer care. International Journal of Qualitative Studies on Health and Well-being, 2022, 17, .	0.6	1
7	The BRIGHTLIGHT National Survey of the Impact of Specialist Teenage and Young Adult Cancer Care on Caregivers' Information and Support Needs. Cancer Nursing, 2021, 44, 235-243.	0.7	9
8	"Holistic Competence― How Is it Developed, Shared, and Shaped by Health Care Professionals Caring for Adolescents and Young Adults with Cancer?. Journal of Adolescent and Young Adult Oncology, 2021, 10, 503-511.	0.7	5
9	Processes of care and survival associated with treatment in specialist teenage and young adult cancer centres: results from the BRIGHTLIGHT cohort study. BMJ Open, 2021, 11, e044854.	0.8	5
10	International perspectives on suboptimal patientâ€reported outcome trial design and reporting in cancer clinical trials: A qualitative study. Cancer Medicine, 2021, 10, 5475-5487.	1.3	11
11	Exploring Young People's Experience of Ending Active Cancer Treatment. Cancer Nursing, 2021, 44, 288-294.	0.7	9
12	Specialist cancer services for teenagers and young adults in England: BRIGHTLIGHT research programme. Programme Grants for Applied Research, 2021, 9, 1-82.	0.4	3
13	The support and information needs of adolescents and young adults with cancer when active treatment ends. BMC Cancer, 2020, 20, 697.	1.1	21
14	BRIGHTLIGHT researchers as †dramaturgs': creating There is a Light from complex research data. Research Involvement and Engagement, 2020, 6, 48.	1.1	3
15	Association of Self-reported Presenting Symptoms With Timeliness of Help-Seeking Among Adolescents and Young Adults With Cancer in the BRIGHTLIGHT Study. JAMA Network Open, 2020, 3, e2015437.	2.8	10
16	Development of a patient-reported experience questionnaire for patients with sarcoma: the Sarcoma Assessment Measure (SAM). Quality of Life Research, 2020, 29, 2287-2297.	1.5	19
17	Longitudinal cohort study of the impact of specialist cancer services for teenagers and young adults on quality of life: outcomes from the BRIGHTLIGHT study. BMJ Open, 2020, 10, e038471.	0.8	17
18	Psychological adaptation and recovery in youth with sarcoma: a qualitative study with practical implications for clinical care and research. BMJ Open, 2020, 10, e038799.	0.8	0

#	Article	IF	Citations
19	Psychological adaptation and recovery in youth with sarcoma: a qualitative study with practical implications for clinical care and research. BMJ Open, 2020, 10, e038799.	0.8	9
20	Understanding care when cure is not likely for young adults who face cancer: a realist analysis of data from patients, families and healthcare professionals. BMJ Open, 2019, 9, e024397.	0.8	10
21	Qualitative study exploring patients experiences of being diagnosed and living with primary bone cancer in the UK. BMJ Open, 2019, 9, e028693.	0.8	17
22	Health care professional perceptions of online information and support for young people with cancer in the United Kingdom. Adolescent Health, Medicine and Therapeutics, 2019, Volume 10, 103-116.	0.7	2
23	Barriers to advance care planning with patients as perceived by nurses and other healthcare professionals: A systematic review. Journal of Clinical Nursing, 2019, 28, 4276-4297.	1.4	70
24	The impact of spine sarcoma on mental health and well-being. Spine Journal, 2019, 19, 1603-1605.	0.6	0
25	The culture of young people's cancer care: A narrative review and synthesis of the UK literature. European Journal of Cancer Care, 2019, 28, e13099.	0.7	6
26	Description of the BRIGHTLIGHT cohort: the evaluation of teenage and young adult cancer services in England. BMJ Open, 2019, 9, e027797.	0.8	17
27	Systematic Evaluation of Patient-Reported Outcome Protocol Content and Reporting in Cancer Trials. Journal of the National Cancer Institute, 2019, 111, 1170-1178.	3.0	76
28	A Critical Review of the Impact of Sarcoma on Psychosocial Wellbeing. Sarcoma, 2019, 2019, 1-18.	0.7	34
29	Patient perspectives on a national multidisciplinary team meeting for a rare cancer. European Journal of Cancer Care, 2019, 28, e12971.	0.7	13
30	Diagnostic timeliness in adolescents and young adults with cancer: a cross-sectional analysis of the BRIGHTLIGHT cohort. The Lancet Child and Adolescent Health, 2018, 2, 180-190.	2.7	42
31	Evaluation of patient-reported outcome protocol content and reporting in UK cancer clinical trials: the EPiC study qualitative protocol. BMJ Open, 2018, 8, e017282.	0.8	6
32	Involving young people in BRIGHTLIGHT from study inception to secondary data analysis: insights from 10 years of user involvement. Research Involvement and Engagement, 2018, 4, 50.	1.1	17
33	Transitioning adolescent and young adult cancer care research out of its adolescence. European Journal of Cancer Care, 2018, 27, e12962.	0.7	3
34	Issues experienced and support provided to adolescents and young adults at the end of active treatment for cancer: A rapid review of the literature. European Journal of Cancer Care, 2018, 27, e12972.	0.7	16
35	Online information and support needs of young people with cancer: a participatory action research study. Adolescent Health, Medicine and Therapeutics, 2018, Volume 9, 121-135.	0.7	28
36	Conceptualizing age-appropriate care for teenagers and young adults with cancer: a qualitative mixed-methods study. Adolescent Health, Medicine and Therapeutics, 2018, Volume 9, 149-166.	0.7	25

#	Article	IF	Citations
37	Sex, Body Image, and Relationships: A BRIGHTLIGHT Workshop on Information and Support Needs of Adolescents and Young Adults. Journal of Adolescent and Young Adult Oncology, 2018, 7, 572-578.	0.7	16
38	Enhancing accrual to clinical trials of adolescents and young adults with cancer. Pediatric Blood and Cancer, 2018, 65, e27233.	0.8	17
39	Experiences and Preferences for End-of-Life Care for Young Adults with Cancer and Their Informal Carers: A Narrative Synthesis. Journal of Adolescent and Young Adult Oncology, 2017, 6, 200-212.	0.7	28
40	Exploring the digital technology preferences of teenagers and young adults (TYA) with cancer and survivors: a cross-sectional service evaluation questionnaire. Journal of Cancer Survivorship, 2017, 11, 670-682.	1.5	53
41	Optimizing a Retention Strategy with Young People for BRIGHTLIGHT, a Longitudinal Cohort Study Examining the Value of Specialist Cancer Care for Young People. Journal of Adolescent and Young Adult Oncology, 2017, 6, 459-469.	0.7	15
42	Qualitative study to understand the barriers to recruiting young people with cancer to BRIGHTLIGHT: a national cohort study in England. BMJ Open, 2017, 7, e018291.	0.8	13
43	Mapping Adolescent Cancer Services. Cancer Nursing, 2016, 39, 358-366.	0.7	23
44	Systematic evaluation of patient-reported outcome (PRO) protocol content and reporting in UK cancer clinical trials: the EPiC study protocol. BMJ Open, 2016, 6, e012863.	0.8	15
45	Psychosocial Interventions for Adolescents and Young Adults Diagnosed with Cancer During Adolescence: A Critical Review. Journal of Adolescent and Young Adult Oncology, 2016, 5, 310-321.	0.7	39
46	Modified international e-Delphi survey to define healthcare professional competencies for working with teenagers and young adults with cancer. BMJ Open, 2016, 6, e011361.	0.8	57
47	Direct access to potential research participants for a cohort study using a confidentiality waiver included in UK National Health Service legal statutes. BMJ Open, 2016, 6, e011847.	0.8	6
48	A participatory study of teenagers and young adults views on access and participation in cancer research. European Journal of Oncology Nursing, 2016, 20, 156-164.	0.9	31
49	A Critical Review of the Use of Technology to Provide Psychosocial Support for Children and Young People with Long-Term Conditions. Journal of Pediatric Nursing, 2015, 30, 87-101.	0.7	20
50	Novel participatory methods of involving patients in research: naming and branding a longitudinal cohort study, BRIGHTLIGHT. BMC Medical Research Methodology, 2015, 15, 20.	1.4	32
51	Development and validation of the BRIGHTLIGHT Survey, a patient-reported experience measure for young people with cancer. Health and Quality of Life Outcomes, 2015, 13, 107.	1.0	50
52	Identifying the supportive care needs of adolescent and young adult survivors of cancer: a qualitative analysis and systematic literature review. Supportive Care in Cancer, 2014, 22, 947-959.	1.0	77
53	Development and Validation of the Pediatric Liver Transplantation Quality of Life: A Disease-Specific Quality of Life Measure for Pediatric Liver Transplant Recipients. Journal of Pediatrics, 2014, 165, 547-555.e7.	0.9	37
54	Developing a conceptual model of teenage and young adult experiences of cancer through meta-synthesis. International Journal of Nursing Studies, 2013, 50, 832-846.	2.5	65

#	Article	IF	CITATIONS
55	The Art of Age-Appropriate Care. Cancer Nursing, 2013, 36, E27-E38.	0.7	88
56	"Your Place or Mine?―Priorities for a Specialist Teenage and Young Adult (TYA) Cancer Unit: Disparity Between TYA and Professional Perceptions. Journal of Adolescent and Young Adult Oncology, 2011, 1, 145-151.	0.7	33
57	Measuring quality of life in children and young people after transplantation: Methodological considerations. Pediatric Transplantation, 2010, 14, 445-458.	0.5	17
58	The Stories of Young People Living With a Liver Transplant. Qualitative Health Research, 2010, 20, 1076-1090.	1.0	56
59	A concept analysis of healthâ€related quality of life in young people with chronic illness. Journal of Clinical Nursing, 2008, 17, 1823-1833.	1.4	125
60	The experience of living with a chronic illness during adolescence: a critical review of the literature. Journal of Clinical Nursing, 2008, 17, 3083-3091.	1.4	147
61	A critical review of the health-related quality of life of children and adolescents after liver transplantation. Liver Transplantation, 2005, 11, 51-60.	1.3	83
62	Liver transplantation in children: part 2 - long-term issues. Journal of Child Health Care, 2005, 9, 274-287.	0.7	4
63	Liver transplantation in children: part 1 - peri-operative issues. Journal of Child Health Care, 2005, 9, 256-273.	0.7	6
64	Immunosuppression for solid organ transplantation in children. Paediatric Nursing, 2004, 16, 39-43.	0.1	1
65	Humoral immunity in children with biliary atresia splenic malformation syndrome. European Journal of Pediatrics, 2003, 162, 539-540.	1.3	9
66	Can energy expenditure be predicted in critically ill children?*. Pediatric Critical Care Medicine, 2003, 4, 176-180.	0.2	77