

Rebekah Clare Laidsaar-Powell

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/4053498/publications.pdf>

Version: 2024-02-01

21
papers

620
citations

759233

12
h-index

713466

21
g-index

21
all docs

21
docs citations

21
times ranked

778
citing authors

#	ARTICLE	IF	CITATIONS
1	Workforce participation in relation to cancer diagnosis, type and stage: Australian population-based study of 163,556 middle-aged people. <i>Journal of Cancer Survivorship</i> , 2022, 16, 461-473.	2.9	7
2	Measuring female breast cancer survivors' concerns about late effects of treatment: initial development of the Concerns about Late Effects in Oncology Questionnaire (CLEO). <i>Supportive Care in Cancer</i> , 2022, 30, 3563-3572.	2.2	2
3	Colorectal cancer survivorship: A systematic review and thematic synthesis of qualitative research. <i>European Journal of Cancer Care</i> , 2021, 30, e13421.	1.5	43
4	Improving breast cancer nurses' management of challenging situations involving family carers: Pilot evaluation of a brief targeted online education module (TRIO-Conflict). <i>Patient Education and Counseling</i> , 2021, 104, 3023-3031.	2.2	6
5	Avoiding the "survivorship abyss": Qualitative insights from 15-year prostate cancer survivors. <i>Psycho-Oncology</i> , 2021, 30, 1745-1755.	2.3	4
6	Considering the type and timing of breast reconstruction after mastectomy: Qualitative insights into women's decision-making. <i>European Journal of Oncology Nursing</i> , 2021, 54, 102024.	2.1	7
7	Return to work after a cancer diagnosis: a meta-review of reviews and a meta-synthesis of recent qualitative studies. <i>Journal of Cancer Survivorship</i> , 2020, 14, 114-134.	2.9	84
8	Disability, psychological distress and quality of life in relation to cancer diagnosis and cancer type: population-based Australian study of 22,505 cancer survivors and 244,000 people without cancer. <i>BMC Medicine</i> , 2020, 18, 372.	5.5	53
9	Online resources for family caregivers of cognitively competent patients: A review of user-driven reputable health website content on caregiver communication with health professionals. <i>Patient Education and Counseling</i> , 2020, 103, 2408-2419.	2.2	5
10	Recognising and managing the psychosocial needs of family carers: It's time for change. <i>Patient Education and Counseling</i> , 2019, 102, 401-403.	2.2	5
11	A meta-review of qualitative research on adult cancer survivors: current strengths and evidence gaps. <i>Journal of Cancer Survivorship</i> , 2019, 13, 852-889.	2.9	35
12	Facilitating collaborative and effective family involvement in the cancer setting: Guidelines for clinicians (TRIO Guidelines-1). <i>Patient Education and Counseling</i> , 2018, 101, 970-982.	2.2	31
13	Health care providers' perceptions of family caregivers' involvement in consultations within a geriatric hospital setting. <i>Geriatric Nursing</i> , 2018, 39, 419-427.	1.9	11
14	Managing challenging interactions with family caregivers in the cancer setting: Guidelines for clinicians (TRIO Guidelines-2). <i>Patient Education and Counseling</i> , 2018, 101, 983-994.	2.2	40
15	A qualitative exploration of patient and family views and experiences of treatment decision-making in bipolar II disorder. <i>Journal of Mental Health</i> , 2018, 27, 66-79.	1.9	25
16	An exploration of Australian psychologists' role in assessing women considering risk-reducing or contralateral prophylactic mastectomy. <i>Breast</i> , 2017, 32, 105-111.	2.2	9
17	The TRIO Framework: Conceptual insights into family caregiver involvement and influence throughout cancer treatment decision-making. <i>Patient Education and Counseling</i> , 2017, 100, 2035-2046.	2.2	57
18	A Qualitative Exploration of Clinician Views and Experiences of Treatment Decision-Making in Bipolar II Disorder. <i>Community Mental Health Journal</i> , 2017, 53, 958-971.	2.0	15

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19	Exploring the communication of oncologists, patients and family members in cancer consultations: development and application of a coding system capturing family-relevant behaviours (KINcode). <i>Psycho-Oncology</i> , 2016, 25, 787-794.	2.3	24
20	Attitudes and experiences of family involvement in cancer consultations: a qualitative exploration of patient and family member perspectives. <i>Supportive Care in Cancer</i> , 2016, 24, 4131-4140.	2.2	46
21	Family involvement in cancer treatment decision-making: A qualitative study of patient, family, and clinician attitudes and experiences. <i>Patient Education and Counseling</i> , 2016, 99, 1146-1155.	2.2	111