Veronica Lambert

List of Publications by Citations

Source: https://exaly.com/author-pdf/9286565/veronica-lambert-publications-by-citations.pdf

Version: 2024-04-10

This document has been generated based on the publications and citations recorded by exaly.com. For the latest version of this publication list, visit the link given above.

The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

40 648 15 24 g-index

46 798 2.7 4.27 ext. papers ext. citations avg, IF L-index

#	Paper	IF	Citations
40	Paediatric early warning systems for detecting and responding to clinical deterioration in children: a systematic review. <i>BMJ Open</i> , 2017 , 7, e014497	3	85
39	Engaging with children in research: Theoretical and practical implications of negotiating informed consent/assent. <i>Nursing Ethics</i> , 2011 , 18, 781-801	3.5	58
38	Striving to live a normal life: a review of children and young people's experience of feeling different when living with a long term condition. <i>Journal of Pediatric Nursing</i> , 2015 , 30, 63-77	2.2	55
37	Social spaces for young children in hospital. <i>Child: Care, Health and Development</i> , 2014 , 40, 195-204	2.8	42
36	Stisible-nessS the nature of communication for children admitted to a specialist childrenS hospital in the Republic of Ireland. <i>Journal of Clinical Nursing</i> , 2008 , 17, 3092-102	3.2	42
35	To tell or not to tell: A systematic review of the disclosure practices of children living with epilepsy and their parents. <i>Epilepsy and Behavior</i> , 2015 , 51, 73-95	3.2	30
34	Communication between children and health professionals in a child hospital setting: a Child Transitional Communication Model. <i>Journal of Advanced Nursing</i> , 2011 , 67, 569-82	3.1	30
33	The stigma experiences and perceptions of families living with epilepsy: Implications for epilepsy-related communication within and external to the family unit. <i>Patient Education and Counseling</i> , 2016 , 99, 1473-81	3.1	29
32	Family communication in the context of pediatric epilepsy: A systematic review. <i>Epilepsy and Behavior</i> , 2015 , 51, 225-39	3.2	26
31	"I don's want them to look at me and think of my illness, I just want them to look at me and see me": Child perspectives on the challenges associated with disclosing an epilepsy diagnosis to others. <i>Epilepsy and Behavior</i> , 2015 , 53, 83-91	3.2	25
30	Clarifying the mechanisms and resources that enable the reciprocal involvement of seldom heard groups in health and social care research: A collaborative rapid realist review process. <i>Health Expectations</i> , 2019 , 22, 298-306	3.7	23
29	Meeting the information needs of children in hospital. Journal of Child Health Care, 2013, 17, 338-53	2	21
28	Using a range of methods to access children's voices. <i>Journal of Research in Nursing</i> , 2013 , 18, 601-616	0.9	20
27	Employing an ethnographic approach: key characteristics. Nurse Researcher, 2011, 19, 17-24	1	20
26	The development of an epilepsy electronic patient portal: Facilitating both patient empowerment and remote clinician-patient interaction in a post-COVID-19 world. <i>Epilepsia</i> , 2020 , 61, 1894-1905	6.4	16
25	Parent perspectives of the challenging aspects of disclosing a childs epilepsy diagnosis to others: Why dons they tell?. <i>Chronic Illness</i> , 2017 , 13, 28-48	1.4	12
24	Talking about epilepsy: Challenges parents face when communicating with their child about epilepsy and epilepsy-related issues. <i>Epilepsy and Behavior</i> , 2016 , 57, 9-15	3.2	11

(2011-2014)

23	Health literacy and its importance for effective communication. Part 2. <i>Nursing Children and Young People</i> , 2014 , 26, 32-6; quiz 38	0.3	11
22	The rhetoric and reality of integrated patient-centered care for healthcare providers: An ethnographic exploration of epilepsy care in Ireland. <i>Epilepsy and Behavior</i> , 2019 , 94, 87-92	3.2	10
21	Virtual reality distraction for acute pain in children. The Cochrane Library, 2020, 10, CD010686	5.2	10
20	Stigmatising feelings and disclosure apprehension among children with epilepsy. <i>Nursing Children and Young People</i> , 2014 , 26, 22-6	0.3	9
19	"The Child's World": a creative and visual trigger to stimulate student enquiry in a problem based learning module. <i>Nurse Education Today</i> , 2008 , 28, 962-9	3.7	9
18	School lunches in the Republic of Ireland: a comparison of the nutritional quality of adolescentsS lunches sourced from home or purchased at school or SoutSat local food outlets. <i>Public Health Nutrition</i> , 2017 , 20, 504-514	3.3	7
17	"I don I like talking about it because that I not who I am": Challenges children face during epilepsy-related family communication. <i>Chronic Illness</i> , 2016 , 12, 216-26	1.4	6
16	Exploring the relationship between parent-child communication about epilepsy and psychosocial well-being. <i>Journal of Health Psychology</i> , 2021 , 26, 1207-1221	3.1	5
15	Quality care metrics (QC-M) in nursing and midwifery care processes: a rapid realist review (RRR) protocol. <i>HRB Open Research</i> , 2020 , 3, 85	1.2	5
14	Children and young peoples experiences and perceptions of self-management of type 1 diabetes: A qualitative meta-synthesis. <i>Health Psychology Open</i> , 2019 , 6, 2055102919877105	1.9	4
13	Design of paediatric hospitals. Nursing Children and Young People, 2016, 28, 55	0.3	3
12	Virtual reality simulation for reducing pain in children. The Cochrane Library, 2013,	5.2	3
11	Response to Carter, B (2008) Commentary on Shields L, Pratt J, Hunter J (2006) Family centred care: a review of qualitative studies Journal of Clinical Nursing 15, 1317-1323 in Journal of Clinical Nursing 17, 2091-2093. <i>Journal of Clinical Nursing</i> , 2009 , 18, 623-4	3.2	3
10	Quality care metrics (QC-M) in nursing and midwifery care processes: a rapid realist review (RRR) protocol. <i>HRB Open Research</i> , 2020 , 3, 85	1.2	3
9	Are patients ready for integrated person-centered care? A qualitative study of people with epilepsy in Ireland. <i>Epilepsy and Behavior</i> , 2020 , 102, 106668	3.2	3
8	Coproducing health and well-being in partnership with patients, families, and healthcare providers: A qualitative study exploring the role of an epilepsy patient portal. <i>Epilepsy and Behavior</i> , 2021 , 115, 107664	3.2	3
7	Promoting healthy diet and physical activity in Irish secondary schools: the views of principals and teachers. <i>Proceedings of the Nutrition Society</i> , 2015 , 74,	2.9	2
6	Commentary on Shields L (2010) Questioning family-centred care. Journal of Clinical Nursing 19, 2629-2638. <i>Journal of Clinical Nursing</i> , 2011 , 20, 1791-3	3.2	2

5	Parent and adolescent communication with healthcare professionals about Type 1 diabetes management at adolescentsSoutpatient clinic appointments. <i>Diabetic Medicine</i> , 2020 , 37, 785-796	3.5	2
4	Development and psychometric evaluation of the youth and parent versions of the Epilepsy Disclosure Scale (EDS). <i>Epilepsy and Behavior</i> , 2018 , 82, 111-118	3.2	1
3	Igniting intersectoral collaboration in chronic disease management: a participatory action research study on epilepsy care in Ireland. <i>Journal of Interprofessional Care</i> , 2020 , 34, 500-508	2.7	1
2	Democratizing epilepsy care: Utility and usability of an electronic patient portal. <i>Epilepsy and Behavior</i> , 2021 , 122, 108197	3.2	0
1	Self-management in children and young people with epilepsy: A systematic review and qualitative meta-synthesis. <i>Journal of Health Psychology</i> , 2021 , 26, 126-139	3.1	