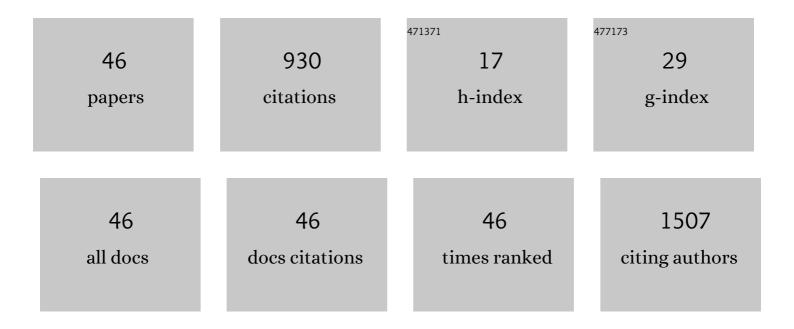
## Laura Elenor Forrest

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/2934397/publications.pdf Version: 2024-02-01



#	Article	IF	CITATIONS
1	Communicating genetic information in families – a review of guidelines and position papers. European Journal of Human Genetics, 2007, 15, 612-618.	1.4	118
2	Increased genetic counseling support improves communication of genetic information in families. Genetics in Medicine, 2008, 10, 167-172.	1.1	94
3	Nurse delivered lifestyle interventions in primary health care to treat chronic disease risk factors associated with obesity: a systematic review. Obesity Reviews, 2012, 13, 1148-1171.	3.1	80
4	Population-based carrier screening for cystic fibrosis: a systematic review of 23 years of research. Genetics in Medicine, 2014, 16, 207-216.	1.1	72
5	How acceptable are primary health care nurse practitioners to Australian consumers?. Collegian, 2013, 20, 35-41.	0.6	70
6	Health first, genetics second: exploring families' experiences of communicating genetic information. European Journal of Human Genetics, 2008, 16, 1329-1335.	1.4	47
7	Genetic health professionals and the communication of genetic information in families: Practice during and after a genetic consultation. American Journal of Medical Genetics, Part A, 2010, 152A, 1458-1466.	0.7	41
8	Breast Cancer Polygenic Risk Score and Contralateral Breast Cancer Risk. American Journal of Human Genetics, 2020, 107, 837-848.	2.6	39
9	Making Sense of SNPs: Women's Understanding and Experiences of Receiving a Personalized Profile of Their Breast Cancer Risks. Journal of Genetic Counseling, 2018, 27, 702-708.	0.9	30
10	A national survey of general practitioners' experiences of patientâ€initiated aggression in Australia. Medical Journal of Australia, 2011, 194, 605-608.	0.8	29
11	"A Natural Progression― Australian Women's Attitudes About an Individualized Breast Screening Model. Cancer Prevention Research, 2019, 12, 383-390.	0.7	26
12	Nurseâ€led primary healthcare walkâ€in centres: an integrative literature review. Journal of Advanced Nursing, 2012, 68, 248-263.	1.5	23
13	Evaluation of telephone genetic counselling to facilitate germline BRCA1/2 testing in women with high-grade serous ovarian cancer. European Journal of Human Genetics, 2019, 27, 1186-1196.	1.4	21
14	The work, education and career pathways of nurses in Australian general practice. Australian Journal of Primary Health, 2011, 17, 227.	0.4	20
15	A Systematic Review of How Young People Live with Inherited Disease: What Can We Learn for Li-Fraumeni Syndrome?. Journal of Adolescent and Young Adult Oncology, 2018, 7, 525-545.	0.7	20
16	An audit of clinical service examining the uptake of genetic testing by at-risk family members. Genetics in Medicine, 2012, 14, 122-128.	1.1	18
17	Clinically Significant Germline Mutations in Cancer-Causing Genes Identified Through Research Studies Should Be Offered to Research Participants by Genetic Counselors. Journal of Clinical Oncology, 2016, 34, 898-901.	0.8	17
18	Patient initiated aggression and violence in Australian general practice. Australian Family Physician, 2010, 39, 323-6.	0.5	17

#	Article	IF	CITATIONS
19	High-risk women's risk perception after receiving personalized polygenic breast cancer risk information. Journal of Community Genetics, 2019, 10, 197-206.	0.5	16
20	Connecting patients, researchers and clinical genetics services: the experiences of participants in the Australian Ovarian Cancer Study (AOCS). European Journal of Human Genetics, 2015, 23, 152-158.	1.4	14
21	"l need to know if l'm going to die youngâ€ŧ Adolescent and young adult experiences of genetic testing for Li–Fraumeni syndrome. Journal of Psychosocial Oncology, 2021, 39, 54-73.	0.6	14
22	Stakeholder perceptions of a nurse led walk-in centre. BMC Health Services Research, 2012, 12, 382.	0.9	13
23	What primary healthâ€eare services are <scp>A</scp> ustralian consumers willing to accept from nurse practitioners? A <scp>N</scp> ational <scp>S</scp> urvey. Health Expectations, 2014, 17, 733-740.	1.1	10
24	Are "part-time―general practitioners workforce idlers or committed professionals?. BMC Family Practice, 2014, 15, 154.	2.9	9
25	Development and implementation of a nurse-led walk-in centre: evidence lost in translation?. Journal of Health Services Research and Policy, 2013, 18, 174-178.	0.8	6
26	Consumer attitudes towards the establishment of a national Australian familial cancer research database by the Inherited Cancer Connect (ICCon) Partnership. Journal of Community Genetics, 2018, 9, 57-64.	0.5	6
27	Social media usage in family communication about genetic information: †I no longer speak with my sister but she needed to know'. Journal of Genetic Counseling, 2021, 30, 180-190.	0.9	6
28	Patient initiated aggression - prevalence and impact for general practice staff. Australian Family Physician, 2011, 40, 415-8.	0.5	6
29	Feasibility and acceptability of nurse-led youth clinics in Australian general practice. Australian Journal of Primary Health, 2013, 19, 159.	0.4	5
30	Young people's experiences of a CDH1 pathogenic variant: Decisionâ€making about gastric cancer risk management. Journal of Genetic Counseling, 2021, , .	0.9	5
31	The impact of participation in genetic research for families with cleft lip with and without cleft palate: a qualitative study. Journal of Community Genetics, 2014, 5, 249-256.	0.5	4
32	Health professionals' practice for young people with, or at risk of, Li–Fraumeni syndrome: An Australasian survey. Journal of Genetic Counseling, 2020, 29, 737-747.	0.9	4
33	Parental experiences and genetic counsellor roles in Pierre Robin sequence. Journal of Community Genetics, 2020, 11, 475-484.	0.5	4
34	Men's experiences of recontact about a potential increased risk of prostate cancer due to Lynch Syndrome: "Just another straw on the stack― Journal of Genetic Counseling, 2019, 28, 750-759.	0.9	3
35	Finding the fiveâ€year window: A qualitative study examining young women's decisionâ€making and experience of using tamoxifen to reduce BRCA1/2 breast cancer risk. Psycho-Oncology, 2021, 30, 159-166.	1.0	3
36	Benefits and burdens of risk management for young people with inherited cancer: A focus on Li-Fraumeni syndrome. Australian Journal of General Practice, 2021, 50, 538-544.	0.3	3

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#	Article	IF	CITATIONS
37	Reproduction and Genetic Responsibility: An Interpretive Description of Reproductive Decision-Making for Young People With Li-Fraumeni Syndrome. Qualitative Health Research, 2022, 32, 168-181.	1.0	3
38	TRACEBACK: Testing of Historical Tubo-Ovarian Cancer Patients for Hereditary Risk Genes as a Cancer Prevention Strategy in Family Members. Journal of Clinical Oncology, 2022, , JCO2102108.	0.8	3
39	Experience of Asian males communicating cardiac genetic risk within the family. Journal of Community Genetics, 2018, 9, 293-303.	0.5	2
40	Mainstreaming genomics: training experience of hospital medical officers at the Royal Melbourne Hospital. Internal Medicine Journal, 2021, 51, 268-271.	0.5	2
41	Assessing the acceptability, feasibility, and usefulness of a psychosocial screening tool to patients and clinicians in a clinical genetics service in Australia. Journal of Genetic Counseling, 2022, 31, 653-662.	0.9	2
42	Development and pilot testing of an online decision aid for women considering risk-stratified breast screening. Journal of Community Genetics, 2022, 13, 137-141.	0.5	2
43	The Clinical and Psychosocial Outcomes for Women Who Received Unexpected Clinically Actionable Germline Information Identified through Research: An Exploratory Sequential Mixed-Methods Comparative Study. Journal of Personalized Medicine, 2022, 12, 1112.	1.1	2
44	Unselected Women's Experiences of Receiving Genetic Research Results for Hereditary Breast and Ovarian Cancer: A Qualitative Study. Genetic Testing and Molecular Biomarkers, 2021, 25, 741-748.	0.3	1
45	A comparison of Australian and French families affected by sarcoma: perceptions of genetics and incidental findings. Personalized Medicine, 2018, 15, 13-24.	0.8	0
46	Oncologists' perspectives of telephone genetic counseling to facilitate germline BRCA1/2 testing for their patients with high-grade serous ovarian cancer. Journal of Community Genetics, 2021, 12, 449-457.	0.5	0