

Leila Dorling

List of Publications by Citations

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Version: 2024-04-28

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

21
papers

974
citations

13
h-index

24
g-index

24
ext. papers

1,442
ext. citations

10.2
avg, IF

2.87
L-index

#	Paper	IF	Citations
21	Randomized controlled trial of intensity-modulated radiotherapy for early breast cancer: 5-year results confirm superior overall cosmesis. <i>Journal of Clinical Oncology</i> , 2013 , 31, 4488-95	2.2	150
20	Breast Cancer Risk Genes - Association Analysis in More than 113,000 Women. <i>New England Journal of Medicine</i> , 2021 , 384, 428-439	59.2	143
19	Cancer Risks Associated With Germline Pathogenic Variants: An International Study of 524 Families. <i>Journal of Clinical Oncology</i> , 2020 , 38, 674-685	2.2	133
18	A genome wide association study (GWAS) providing evidence of an association between common genetic variants and late radiotherapy toxicity. <i>Radiotherapy and Oncology</i> , 2014 , 111, 178-85	5.3	102
17	A three-stage genome-wide association study identifies a susceptibility locus for late radiotherapy toxicity at 2q24.1. <i>Nature Genetics</i> , 2014 , 46, 891-4	36.3	92
16	Replication of genetic polymorphisms reported to be associated with taxane-related sensory neuropathy in patients with early breast cancer treated with Paclitaxel. <i>Clinical Cancer Research</i> , 2014 , 20, 2466-75	12.9	78
15	Individual patient data meta-analysis shows a significant association between the ATM rs1801516 SNP and toxicity after radiotherapy in 5456 breast and prostate cancer patients. <i>Radiotherapy and Oncology</i> , 2016 , 121, 431-439	5.3	69
14	Meta-analysis of Genome Wide Association Studies Identifies Genetic Markers of Late Toxicity Following Radiotherapy for Prostate Cancer. <i>EBioMedicine</i> , 2016 , 10, 150-63	8.8	50
13	Patient reported outcome measures (PROMs) following forward planned field-in field IMRT: results from the Cambridge Breast IMRT trial. <i>Radiotherapy and Oncology</i> , 2014 , 111, 270-5	5.3	33
12	Inherited mutations in and in an unselected multiethnic cohort of Asian patients with breast cancer and healthy controls from Malaysia. <i>Journal of Medical Genetics</i> , 2018 , 55, 97-103	5.8	24
11	A nested cohort study of 6,248 early breast cancer patients treated in neoadjuvant and adjuvant chemotherapy trials investigating the prognostic value of chemotherapy-related toxicities. <i>BMC Medicine</i> , 2015 , 13, 306	11.4	18
10	Prevalence of BRCA1 and BRCA2 pathogenic variants in a large, unselected breast cancer cohort. <i>International Journal of Cancer</i> , 2019 , 144, 1195-1204	7.5	18
9	Differential Burden of Rare and Common Variants on Tumor Characteristics, Survival, and Mode of Detection in Breast Cancer. <i>Cancer Research</i> , 2018 , 78, 6329-6338	10.1	13
8	Hereditary breast and ovarian cancer: successful systematic implementation of a group approach to genetic counselling. <i>Familial Cancer</i> , 2017 , 16, 51-56	3	12
7	Common genetic variation associated with increased susceptibility to prostate cancer does not increase risk of radiotherapy toxicity. <i>British Journal of Cancer</i> , 2016 , 114, 1165-74	8.7	12
6	Patients with a High Polygenic Risk of Breast Cancer do not have An Increased Risk of Radiotherapy Toxicity. <i>Clinical Cancer Research</i> , 2016 , 22, 1413-20	12.9	11
5	The Relationship between Common Genetic Markers of Breast Cancer Risk and Chemotherapy-Induced Toxicity: A Case-Control Study. <i>PLoS ONE</i> , 2016 , 11, e0158984	3.7	7

4	First international workshop of the ATM and cancer risk group (4-5 December 2019). <i>Familial Cancer</i> , 2021 , 1	3	5
3	Pathology of Tumors Associated With Pathogenic Germline Variants in 9 Breast Cancer Susceptibility Genes.. <i>JAMA Oncology</i> , 2022 ,	13.4	4
2	Rare germline copy number variants (CNVs) and breast cancer risk.. <i>Communications Biology</i> , 2022 , 5, 65	6.7	0
1	Breast cancer risks associated with missense variants in breast cancer susceptibility genes.. <i>Genome Medicine</i> , 2022 , 14, 51	14.4	0