## Eva Tomiak

## List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/5944731/publications.pdf

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		1163117	1281871	
13	574	8	11	
papers	citations	h-index	g-index	
14	14	14	1332	
all docs	docs citations	times ranked	citing authors	

#	Article	IF	CITATIONS
1	A decade of <i>RAD51C</i> and <i>RAD51D</i> germline variants in cancer. Human Mutation, 2022, 43, 285-298.	2.5	6
2	Meaningful relationships as a driving force in the experience of parents of a child living with polyposis conditions. Psychology, Health and Medicine, $2021$ , , $1-12$ .	2.4	0
3	Canadian Urological Association Best Practice Report on the long-term followup for patients with pheochromocytomas. Canadian Urological Association Journal, 2019, 13, 372-376.	0.6	3
4	Broad spectrum of neuropsychiatric phenotypes associated with white matter disease in <i>PTEN</i> hamartoma tumor syndrome. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2018, 177, 101-109.	1.7	34
5	Evolution of genetic assessment for BRCA-associated gynaecologic malignancies: a Canadian multisociety roadmap. Journal of Medical Genetics, 2018, 55, 571-577.	3.2	33
6	Functionally Null <i>RAD51D</i> Missense Mutation Associates Strongly with Ovarian Carcinoma. Cancer Research, 2017, 77, 4517-4529.	0.9	34
7	Recurrent spontaneous pneumothoraces and bullous emphysema. A novel mutation causing Birt-Hogg-Dube syndrome. Respiratory Medicine Case Reports, 2016, 19, 106-108.	0.4	5
8	Multigene panels in prostate cancer risk assessment: a systematic review. Genetics in Medicine, 2016, 18, 535-544.	2.4	11
9	Genetic testing: When prediction generates stigmatization. Journal of Health Psychology, 2015, 20, 393-400.	2.3	13
10	Evaluation of clinical and histological criteria and outcome in management of hereditary non-polyposis colorectal carcinoma (HNPCC) Journal of Clinical Oncology, 2015, 33, e14552-e14552.	1.6	0
11	<i>DICER1</i> mutations in an adolescent with cervical embryonal rhabdomyosarcoma (cERMS). Pediatric Blood and Cancer, 2014, 61, 568-569.	1.5	33
12	Mutation analysis of PALB2 in BRCA1 and BRCA2-negative breast and/or ovarian cancer families from Eastern Ontario, Canada. Hereditary Cancer in Clinical Practice, 2014, 12, 19.	1.5	19
13	Germline and somatic SMARCA4 mutations characterize small cell carcinoma of the ovary, hypercalcemic type. Nature Genetics, 2014, 46, 438-443.	21.4	383