Marianna BoreckÃ¡

List of Publications by Year
in descending order

13	221	1163117 8	1125743 13
papers	citations	h-index	g-index

14	14	14	551
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Identification of deleterious germline CHEK2 mutations and their association with breast and ovarian cancer. International Journal of Cancer, 2019, 145, 1782-1797.	5.1	62
2	Validation of CZECANCA (CZEch CAncer paNel for Clinical Application) for targeted NGS-based analysis of hereditary cancer syndromes. PLoS ONE, 2018, 13, e0195761.	2.5	31
3	Mutation analysis of the PALB2 gene in unselected pancreatic cancer patients in the Czech Republic. Cancer Genetics, 2016, 209, 199-204.	0.4	27
4	Estrogen Receptor Status Oppositely Modifies Breast Cancer Prognosis in BRCA1/BRCA2 Mutation Carriers Versus Non-Carriers. Cancers, 2019, 11, 738.	3.7	22
5	Multigene Panel Germline Testing of 1333 Czech Patients with Ovarian Cancer. Cancers, 2020, 12, 956.	3.7	19
6	BRCA1 and BRCA2 5â€² noncoding region variants identified in breast cancer patients alter promoter activity and protein binding. Human Mutation, 2018, 39, 2025-2039.	2.5	15
7	The c.657del5 variant in the NBN gene predisposes to pancreatic cancer. Gene, 2016, 587, 169-172.	2.2	13
8	Truncated PPM1D impairs stem cell response to genotoxic stress and promotes growth of APC-deficient tumors in the mouse colon. Cell Death and Disease, 2019, 10, 818.	6.3	12
9	Prevalence of Germline Pathogenic Variants in Cancer Predisposing Genes in Czech and Belgian Pancreatic Cancer Patients. Cancers, 2021, 13, 4430.	3.7	8
10	Genetic analysis of subsequent second primary malignant neoplasms in long-term pancreatic cancer survivors suggests new potential hereditary genetic alterations. Cancer Management and Research, 2019, Volume 11, 599-609.	1.9	4
11	Germline CHEK2 Gene Mutations in Hereditary Breast Cancer Predisposition â€“ Mutation Types and their Biological and Clinical Relevance. Klinicka Onkologie, 2019, 32, 36-50.	0.3	2
12	Novel presentation of the c.1856Aâ€‰>â€‰G (p.Asp619Gly) TSHR gene-activating variant: relapsing hyperthyroidism in three subsequent generations manifesting in early childhood and an in vitro functional study. Hormones, 2021, 20, 803-812.	1.9	1
13	Contribution of Massive Parallel Sequencing to Diagnosis of Hereditary Ovarian Cancer in the Czech Republic. Klinicka Onkologie, 2019, 32, 72-78.	0.3	1