Eva Machackova

List of Publications by Year in descending order

30 papers	2,865 citations	516215 16 h-index	454577 30 g-index
33 all docs	33 docs citations	33 times ranked	4948 citing authors

#	Article	IF	Citations
1	Polygenic risk modeling for prediction of epithelial ovarian cancer risk. European Journal of Human Genetics, 2022, 30, 349-362.	1.4	23
2	Re: ERCC3, a new ovarian cancer susceptibility gene?. European Journal of Cancer, 2021, 150, 278-280.	1.3	1
3	Identification of Germline Mutations in Melanoma Patients with Early Onset, Double Primary Tumors, or Family Cancer History by NGS Analysis of 217 Genes. Biomedicines, 2020, 8, 404.	1.4	10
4	Characterization of the Cancer Spectrum in Men With GermlineBRCA1andBRCA2Pathogenic Variants. JAMA Oncology, 2020, 6, 1218.	3.4	48
5	Multigene Panel Germline Testing of 1333 Czech Patients with Ovarian Cancer. Cancers, 2020, 12, 956.	1.7	19
6	Identification of deleterious germline CHEK2 mutations and their association with breast and ovarian cancer. International Journal of Cancer, 2019, 145, 1782-1797.	2.3	62
7	Functional evaluation of variants of unknown significance in theBRCA2gene identified in genetic testing. Cancer Biology and Therapy, 2019, 20, 633-641.	1.5	1
8	GAPPS â€“ Gastric Adenocarcinoma and Proximal Polyposis of the Stomach Syndrome in 8 Families Tested at Masaryk Memorial Cancer Institute â€“ Prevention and Prophylactic Gastrectomies. Klinicka Onkologie, 2019, 32, 109-117.	0.1	20
9	BAP1 Syndrome â€“ Predisposition to Malignant Mesothelioma, Skin and Uveal Melanoma, Renal and Other Cancers. Klinicka Onkologie, 2019, 32, 118-122.	0.1	7
10	Germline CHEK2 Gene Mutations in Hereditary Breast Cancer Predisposition â€“ Mutation Types and their Biological and Clinical Relevance. Klinicka Onkologie, 2019, 32, 36-50.	0.1	2
11	Recommendations for Preventive Care for Women with Rare Genetic Cause of Breast and Ovarian Cancer. Klinicka Onkologie, 2019, 32, 6-13.	0.1	8
12	Contribution of Massive Parallel Sequencing to Diagnosis of Hereditary Ovarian Cancer in the Czech Republic. Klinicka Onkologie, 2019, 32, 72-78.	0.1	1
13	Twenty Years of BRCA1 and BRCA2 Molecular Analysis at MMCI â€“ Current Developments for the Classifi cation of Variants. Klinicka Onkologie, 2019, 32, 51-71.	0.1	5
14	Mutational spectrum in a worldwide study of 29,700 families withBRCA1orBRCA2mutations. Human Mutation, 2018, 39, 593-620.	1.1	224
15	Thorough in silico and in vitro cDNA analysis of 21 putativeBRCA1andBRCA2splice variants and a complex tandem duplication inBRCA2allowing the identification of activated cryptic splice donor sites inBRCA2exon 11. Human Mutation, 2018, 39, 515-526.	1.1	5
16	Validation of CZECANCA (CZEch CAncer paNel for Clinical Application) for targeted NGS-based analysis of hereditary cancer syndromes. PLoS ONE, 2018, 13, e0195761.	1.1	31
17	BRCA2 Hypomorphic Missense Variants Confer Moderate Risks of Breast Cancer. Cancer Research, 2017, 77, 2789-2799.	0.4	75
18	Risks of Breast, Ovarian, and Contralateral Breast Cancer for BRCA1 and BRCA2 Mutation Carriers. JAMA - Journal of the American Medical Association, 2017, 317, 2402.	3.8	1,898

#	Article	IF	Citations
19	Identification and Functional Testing of ERCC2 Mutations in a Multi-national Cohort of Patients with Familial Breast- and Ovarian Cancer. PLoS Genetics, 2016, 12, e1006248.	1.5	22
20	Male breast cancer in BRCA1 and BRCA2 mutation carriers: pathology data from the Consortium of Investigators of Modifiers of BRCA1/2. Breast Cancer Research, 2016, 18, 15.	2.2	88
21	Fanconi anemia with biallelic FANCD1/BRCA2 mutations â€“ Case report of a family with three affected children. European Journal of Medical Genetics, 2016, 59, 152-157.	0.7	11
22	BRCA1 Circos: a visualisation resource for functional analysis of missense variants. Journal of Medical Genetics, 2015, 52, 224-230.	1.5	32
23	The AIB1 gene polyglutamine repeat length polymorphism and the risk of breast cancer development. Journal of Cancer Research and Clinical Oncology, 2011, 137, 331-338.	1.2	5
24	Spectrum and characterisation of BRCA1 and BRCA2deleterious mutations in high-risk Czech patients with breast and/or ovarian cancer. BMC Cancer, 2008, 8, 140.	1.1	64
25	High occurrence of BRCA1 intragenic rearrangements in hereditary breast and ovarian cancer syndrome in the Czech Republic. BMC Medical Genetics, 2007, 8, 32.	2.1	45
26	Genetic and Preventive Services for Hereditary Breast and Ovarian Cancer in the Czech Republic. Hereditary Cancer in Clinical Practice, 2006, 4, 3.	0.6	6
27	BRCA1 and BRCA2 mutations in women with familial or early-onset breast/ovarian cancer in the Czech Republic. Human Mutation, 2004, 23, 397-398.	1.1	36
28	Differentiating pathogenic mutations from polymorphic alterations in the splice sites ofBRCA1 andBRCA2. Genes Chromosomes and Cancer, 2003, 37, 314-320.	1.5	78
29	Novel germline BRCA1 and BRCA2 mutations in breast and breast/ovarian cancer families from the Czech Republic. Human Mutation, 2001, 18, 545-545.	1.1	7
30	Mutation analysis of the BRCA1 and BRCA2 genes results in the identification of novel and recurrent mutations in 6/16 Flemish families with breast and/or ovarian cancer but not in 12 sporadic patients with early-onset disease. , 1999, 13, 256-256.		22