Aneta BaÅ‚abas

List of Publications by Year
in descending order

16 papers	331 citations	933447 10 h-index	940533 16 g-index
16 all docs	16 docs citations	16 times ranked	832 citing authors

#	Article	IF	Citations
1	Signatures of circulating microRNA in four sarcoma subtypes. Journal of Cancer, 2020, 11, 874-882.	2.5	12
2	Clinical importance of FANCD2, BRIP1, BRCA1, BRCA2 and FANCF expression in ovarian carcinomas. Cancer Biology and Therapy, 2019, 20, 843-854.	3.4	20
3	Wholeâ€exome sequencing identifies novel pathogenic variants across the ATP7B gene and some modifiers of Wilson's disease phenotype. Liver International, 2019, 39, 177-186.	3.9	38
4	Redefining the Practical Utility of Blood Transcriptome Biomarkers in Inflammatory Bowel Diseases. Journal of Crohn's and Colitis, 2019, 13, 626-633.	1.3	22
5	Somatic aberrations of BRCA1 gene are associated with ALDH1, EGFR, and tumor progression in prostate cancer. International Journal of Cancer, 2019, 144, 607-614.	5.1	11
6	GWAS links variants in neuronal development and actin remodeling related loci with pseudoexfoliation syndrome without glaucoma. Experimental Eye Research, 2018, 168, 138-148.	2.6	22
7	IDH1/2 Mutations Predict Shorter Survival in Chondrosarcoma. Journal of Cancer, 2018, 9, 998-1005.	2.5	50
8	A preliminary evaluation of next-generation sequencing as a screening tool for targeted genotyping of erythrocyte and platelet antigens in blood donors. Blood Transfusion, 2018, 16, 285-292.	0.4	23
9	Combination Testing Using a Single MSH5 Variant alongside HLA Haplotypes Improves the Sensitivity of Predicting Coeliac Disease Risk in the Polish Population. PLoS ONE, 2015, 10, e0139197.	2.5	6
10	New recurrent BRCA1/2 mutations in Polish patients with familial breast/ovarian cancer detected by next generation sequencing. BMC Medical Genomics, 2015, 8, 19.	1.5	34
11	Association of the BRCA1 promoter polymorphism rs11655505 with the risk of familial breast and/or ovarian cancer. Familial Cancer, 2013, 12, 691-698.	1.9	2
12	Limited significance of family history for presence of BRCA1 gene mutation in Polish breast and ovarian cancer cases. Familial Cancer, 2012, 11, 351-354.	1.9	20
13	High frequency of BRCA1 founder mutations in Polish women with nonfamilial breast cancer. Familial Cancer, 2012, 11, 623-628.	1.9	20
14	Prevalence of the most frequent BRCA1 mutations in Polish population. Journal of Applied Genetics, 2011, 52, 325-330.	1.9	38
15	Novel germline mutations in BRCA2 gene among breast and breast-ovarian cancer families from Poland. Familial Cancer, 2010, 9, 267-274.	1.9	6
16	Age at Onset of Bilateral Breast Cancer, the Presence of Hereditary BRCA1, BRCA2, CHEK2 Gene Mutations and Positive Family History of Cancer. Oncology Research and Treatment, 2009, 32, 182-188.	1.2	7