Anna Laura Putignano

List of Publications by Year
in descending order

16 papers	960 citations	687363 13 h-index	940533 16 g-index
16 all docs	16 docs citations	16 times ranked	2399 citing authors

#	Article	IF	Citations
1	Pathology of Breast and Ovarian Cancers among BRCA1 and BRCA2 Mutation Carriers: Results from the Consortium of Investigators of Modifiers of BRCA1/2 (CIMBA). Cancer Epidemiology Biomarkers and Prevention, 2012, 21, 134-147.	2.5	513
2	FANCMc.5791C>T nonsense mutation (rs144567652) induces exon skipping, affects DNA repair activity and is a familial breast cancer risk factor. Human Molecular Genetics, 2015, 24, 5345-5355.	2.9	91
3	Common alleles at 6q25.1 and 1p11.2 are associated with breast cancer risk for BRCA1 and BRCA2 mutation carriers. Human Molecular Genetics, 2011, 20, 3304-3321.	2.9	68
4	Interleukin-10 promoter polymorphisms influence susceptibility to ulcerative colitis in a gender-specific manner. Scandinavian Journal of Gastroenterology, 2008, 43, 712-718.	1.5	50
5	A PALB2 germline mutation associated with hereditary breast cancer in Italy. Familial Cancer, 2010, 9, 181-185.	1.9	39
6	Founder mutations account for the majority of BRCA1-attributable hereditary breast/ovarian cancer cases in a population from Tuscany, Central Italy. Breast Cancer Research and Treatment, 2009, 117, 497-504.	2.5	31
7	Susceptibility to Refractory Ulcerative Colitis Is Associated with Polymorphism in the hMLH1 Mismatch Repair Gene. Inflammatory Bowel Diseases, 2004, 10, 705-708.	1.9	25
8	Characterization of an Italian Founder Mutation in the RING-Finger Domain of BRCA1. PLoS ONE, 2014, 9, e86924.	2.5	24
9	Exploring the link between MORF4L1 and risk of breast cancer. Breast Cancer Research, 2011, 13, R40.	5.0	23
10	Thymidylate synthase expression and genotype have no major impact on the clinical outcome of colorectal cancer patients treated with 5-fluorouracil. Pharmacological Research, 2011, 64, 242-248.	7.1	21
11	Lack of association between TNF-Î± polymorphisms and Alzheimerâ€™s disease in an Italian cohort. Neuroscience Letters, 2008, 446, 139-142.	2.1	20
12	Double somatic SMARCB1 and NF2 mutations in sporadic spinal schwannoma. Journal of Neuro-Oncology, 2018, 137, 33-38.	2.9	16
13	Germline mutations in MEN1 and BRCA1 genes in a woman with familial multiple endocrine neoplasia type 1 and inherited breastâ€“ovarian cancer syndromes: a case report. Cancer Genetics and Cytogenetics, 2009, 195, 75-79.	1.0	14
14	Donor-Specific Anti-HLA Antibodies in Huntington's Disease Recipients of Human Fetal Striatal Grafts. Cell Transplantation, 2015, 24, 811-817.	2.5	12
15	Evaluation of a Next-Generation Sequencing Assay for BRCA1 and BRCA2 Mutation Detection. Journal of Molecular Diagnostics, 2018, 20, 87-94.	2.8	11
16	Detection of Rearrangements in the NF2 Gene Using Semi-Quantitative Multiplex Fluorescent PCR. Genetic Testing and Molecular Biomarkers, 2005, 9, 14-19.	1.7	2