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	Evaluation of a Next-Generation Sequencing Assay for BRCA1 and BRCA2 Mutation Detection. Journal of Molecular Diagnostics, 2018, 20, 87-94.	2.8	11
2	Double somatic SMARCB1 and NF2 mutations in sporadic spinal schwannoma. Journal of Neuro-Oncology, 2018, 137, 33-38.	2.9	16
3	Donor-Specific Anti-HLA Antibodies in Huntington's Disease Recipients of Human Fetal Striatal Grafts. Cell Transplantation, 2015, 24, 811-817.	2.5	12
4	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
5	Characterization of an Italian Founder Mutation in the RING-Finger Domain of BRCA1. PLoS ONE, 2014, 9, e86924.	2.5	24
6	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
7	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
8	Exploring the link between MORF4L1 and risk of breast cancer. Breast Cancer Research, 2011, 13, R40.	5.0	23
9	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
10	A PALB2 germline mutation associated with hereditary breast cancer in Italy. Familial Cancer, 2010, 9, 181-185.	1.9	39
11	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
12	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
13	Lack of association between TNF-Î± polymorphisms and Alzheimerâ€™s disease in an Italian cohort. Neuroscience Letters, 2008, 446, 139-142.	2.1	20
14	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
15	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
16	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