Valentina Zampiga

List of Publications by Year
in descending order

13	268	1163117 8	1372567 10
papers	citations	h-index	g-index

13	13	13	957
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Large scale multifactorial likelihood quantitative analysis of BRCA1 and BRCA2 variants: An ENIGMA resource to support clinical variant classification. Human Mutation, 2019, 40, 1557-1578.	2.5	102
2	Multiple-gene panel analysis in a case series of 255 women with hereditary breast and ovarian cancer. Oncotarget, 2017, 8, 47064-47075.	1.8	68
3	Multigene Panel Testing Increases the Number of Loci Associated with Gastric Cancer Predisposition. Cancers, 2019, 11, 1340.	3.7	19
4	Insights into Genetic Susceptibility to Melanoma by Gene Panel Testing: Potential Pathogenic Variants in ACD, ATM, BAP1, and POT1. Cancers, 2020, 12, 1007.	3.7	19
5	Male Breast Cancer: Results of the Application of Multigene Panel Testing to an Italian Cohort of Patients. Diagnostics, 2020, 10, 269.	2.6	14
6	First evidence of a large CHEK2 duplication involved in cancer predisposition in an Italian family with hereditary breast cancer. BMC Cancer, 2014, 14, 478.	2.6	12
7	The Spectrum of FANCM Protein Truncating Variants in European Breast Cancer Cases. Cancers, 2020, 12, 292.	3.7	11
8	BRCA1 p.His1673del is a pathogenic mutation associated with a predominant ovarian cancer phenotype. Oncotarget, 2017, 8, 22640-22648.	1.8	10
9	Results of a population-based screening for hereditary breast cancer in a region of North-Central Italy: contribution of BRCA1/2 germ-line mutations. Breast Cancer Research and Treatment, 2008, 112, 343-349.	2.5	7
10	Disease family history and modification of breast cancer risk in common BRCA2 variants. Oncology Reports, 0, , .	2.6	4
11	Multiple Primary Tumors in a Family with Li-Fraumeni Syndrome with a TP53 Germline Mutation Identified by Next-Generation Sequencing. International Journal of Biological Markers, 2016, 31, 461-465.	1.8	2
12	Characterization of Molecular Alterations of BRCA1/2: Analysis and Interpretation Guidelines. Current Women's Health Reviews, 2012, 8, 4-11.	0.2	0
13	Identification of a novel large EPCAM-MSH2 duplication, concurrently with LOHs in chromosome 20 and X, in a family with Lynch syndrome. International Journal of Colorectal Disease, 2019, 34, 1999-2002.	2.2	0