Hildegunn HÃ¸berg-Vetti

List of Publications by Year
in descending order

11 papers	348 citations	1307594 7 h-index	1281871 11 g-index
11 all docs	11 docs citations	11 times ranked	1026 citing authors

#	Article	IF	Citations
1	Mutational Signature Analysis Reveals NTHL1 Deficiency to Cause a Multi-tumor Phenotype. Cancer Cell, 2019, 35, 256-266.e5.	16.8	123
2	BRCA1/2 testing in newly diagnosed breast and ovarian cancer patients without prior genetic counselling: the DNA-BONus study. European Journal of Human Genetics, 2016, 24, 881-888.	2.8	58
3	BRCA Testing by Single-Molecule Molecular Inversion Probes. Clinical Chemistry, 2017, 63, 503-512.	3.2	46
4	White Blood Cell BRCA1 Promoter Methylation Status and Ovarian Cancer Risk. Annals of Internal Medicine, 2018, 168, 326.	3.9	37
5	Unraveling genetic predisposition to familial or early onset gastric cancer using germline whole-exome sequencing. European Journal of Human Genetics, 2017, 25, 1246-1252.	2.8	34
6	Identifying Needs: a Qualitative Study of women's Experiences Regarding Rapid Genetic Testing for Hereditary Breast and Ovarian Cancer in the DNA BONus Study. Journal of Genetic Counseling, 2017, 26, 182-189.	1.6	16
7	Anxiety and depression symptoms among women attending group-based patient education courses for hereditary breast and ovarian cancer. Hereditary Cancer in Clinical Practice, 2017, 15, 2.	1.5	15
8	Cancer-related distress in unselected women with newly diagnosed breast or ovarian cancer undergoing BRCA1/2 testing without pretest genetic counseling. Acta OncolÃ³gica, 2019, 58, 175-181.	1.8	6
9	The intronic BRCA1 c.5407-25T>A variant causing partly skipping of exon 23â€”a likely pathogenic variant with reduced penetrance?. European Journal of Human Genetics, 2020, 28, 1078-1086.	2.8	6
10	Validation and clinical application of transactivation assays for RUNX1 variant classification. Blood Advances, 2022, , .	5.2	5
11	BRCA1 Norway: comparison of classification for BRCA1 germline variants detected in families with suspected hereditary breast and ovarian cancer between different laboratories. Familial Cancer, 2022, 21, 389-398.	1.9	2