

Elena Tenedini

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/7317273/publications.pdf>

Version: 2024-02-01

12
papers

267
citations

1163065

8
h-index

1281846

11
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12
all docs

12
docs citations

12
times ranked

582
citing authors

#	ARTICLE	IF	CITATIONS
1	Characterization of New ATM Deletion Associated with Hereditary Breast Cancer. <i>Genes</i> , 2021, 12, 136.	2.4	7
2	The Prognostic and Predictive Role of Somatic BRCA Mutations in Ovarian Cancer: Results from a Multicenter Cohort Study. <i>Diagnostics</i> , 2021, 11, 565.	2.6	7
3	Clinicopathologic Profile of Breast Cancer in Germline ATM and CHEK2 Mutation Carriers. <i>Genes</i> , 2021, 12, 616.	2.4	15
4	Ceruloplasmin gene variants are associated with hyperferritinemia and increased liver iron in patients with NAFLD. <i>Journal of Hepatology</i> , 2021, 75, 506-513.	3.7	40
5	Pre-existing cytopenia heralding de novo acute myeloid leukemia: uncommon presentation of NPM1-mutated AML in a single-center study. <i>Leukemia Research</i> , 2021, 111, 106747.	0.8	0
6	P2X7 Receptor Activity Limits Accumulation of T Cells within Tumors. <i>Cancer Research</i> , 2020, 80, 3906-3919.	0.9	36
7	BRCA Detection Rate in an Italian Cohort of Luminal Early-Onset and Triple-Negative Breast Cancer Patients without Family History: When Biology Overcomes Genealogy. <i>Cancers</i> , 2020, 12, 1252.	3.7	15
8	WISP-2 expression induced by Teriparatide treatment affects in vitro osteoblast differentiation and improves in vivo osteogenesis. <i>Molecular and Cellular Endocrinology</i> , 2020, 513, 110817.	3.2	9
9	Hereditary Pancreatic Cancer: A Retrospective Single-Center Study of 5143 Italian Families with History of BRCA-Related Malignancies. <i>Cancers</i> , 2019, 11, 193.	3.7	12
10	Genomic alterations at the basis of treatment resistance in metastatic breast cancer: clinical applications. <i>Oncotarget</i> , 2018, 9, 31606-31619.	1.8	11
11	The chaperone activity of 4PBA ameliorates the skeletal phenotype of Chihuahua, a zebrafish model for dominant osteogenesis imperfecta. <i>Human Molecular Genetics</i> , 2017, 26, 2897-2911.	2.9	68
12	Unravelling the Complexity of Inherited Retinal Dystrophies Molecular Testing: Added Value of Targeted Next-Generation Sequencing. <i>BioMed Research International</i> , 2016, 2016, 1-14.	1.9	47