Dina Ruano

List of Publications by Year
in descending order

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24 papers	837 citations	687363 13 h-index	610901 24 g-index
25 all docs	25 docs citations	25 times ranked	1802 citing authors

#	Article	IF	Citations
1	Cell-of-origin classification using the Hans and Lymph2Cx algorithms in primary cutaneous large B-cell lymphomas. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2022, 480, 667-675.	2.8	12
2	Targeting pancreatic cancer by TAK-981: a SUMOylation inhibitor that activates the immune system and blocks cancer cell cycle progression in a preclinical model. Gut, 2022, 71, 2266-2283.	12.1	35
3	Prevalence and Prognosis of Lynch Syndrome and Sporadic Mismatch Repair Deficiency in Endometrial Cancer. Journal of the National Cancer Institute, 2021, 113, 1212-1220.	6.3	47
4	Frequent mutated B2M, EZH2, IRF8, and TNFRSF14 in primary bone diffuse large B-cell lymphoma reflect a GCB phenotype. Blood Advances, 2021, 5, 3760-3775.	5.2	11
5	Yield and costs of molecular diagnostics on thyroid cytology slides in the Netherlands, adapting the Bethesda classification. Endocrinology, Diabetes and Metabolism, 2021, 4, e00293.	2.4	7
6	The complexity of screening PMS2 in DNA isolated from formalin-fixed paraffin-embedded material. European Journal of Human Genetics, 2020, 28, 333-338.	2.8	10
7	Identification of a neo-epitope dominating endogenous CD8 T cell responses to MC-38 colorectal cancer. OncoImmunology, 2020, 9, 1673125.	4.6	40
8	Digenic inheritance of <scp>MSH6</scp> and <scp>MUTYH</scp> variants in familial colorectal cancer. Genes Chromosomes and Cancer, 2020, 59, 697-701.	2.8	9
9	Optimizing Mutation and Fusion Detection in NSCLC by Sequential DNA and RNA Sequencing. Journal of Thoracic Oncology, 2020, 15, 1000-1014.	1.1	68
10	Allelic Switching of DLX5, GRB10, and SVOPL during Colorectal Cancer Tumorigenesis. International Journal of Genomics, 2019, 2019, 1-10.	1.6	4
11	Low frequency of POLD1 and POLE exonuclease domain variants in patients with multiple colorectal polyps. Molecular Genetics & Genomic Medicine, 2019, 7, e00603.	1.2	8
12	Revisiting immune escape in colorectal cancer in the era of immunotherapy. British Journal of Cancer, 2019, 120, 815-818.	6.4	30
13	Validation and Implementation of BRCA1/2 Variant Screening in Ovarian Tumor Tissue. Journal of Molecular Diagnostics, 2018, 20, 600-611.	2.8	18
14	Molecular Background of Colorectal Tumors From Patients With Lynch Syndrome Associated With Germline Variants in PMS2. Gastroenterology, 2018, 155, 844-851.	1.3	38
15	Molecular Analysis of Gene Fusions in Bone and Soft Tissue Tumors by Anchored Multiplex PCRâ€“Based Targeted Next-Generation Sequencing. Journal of Molecular Diagnostics, 2018, 20, 653-663.	2.8	85
16	Genomic Characterization of Vulvar (Pre)cancers Identifies Distinct Molecular Subtypes with Prognostic Significance. Clinical Cancer Research, 2017, 23, 6781-6789.	7.0	110
17	Evidence for genetic association between chromosome 1q loci and predisposition to colorectal neoplasia. British Journal of Cancer, 2017, 117, 1215-1223.	6.4	10
18	Distinct Patterns of Somatic Mosaicism in the APC GeneÂinÂNeoplasms From Patients With Unexplained AdenomatousÂPolyposis. Gastroenterology, 2017, 152, 546-549.e3.	1.3	27

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19	Characterization of novel low passage primary and metastatic colorectal cancer cell lines. Oncotarget, 2016, 7, 14499-14509.	1.8	11
20	Whole Gene Capture Analysis of 15 CRC Susceptibility Genes in Suspected Lynch Syndrome Patients. PLoS ONE, 2016, 11, e0157381.	2.5	12
21	Combined mismatch repair and POLE/POLD1 defects explain unresolved suspected Lynch syndrome cancers. European Journal of Human Genetics, 2016, 24, 1089-1092.	2.8	110
22	Recurrent Coding Sequence Variation Explains Only A Small Fraction of the Genetic Architecture of Colorectal Cancer. Scientific Reports, 2015, 5, 16286.	3.3	24
23	Target-Enriched Next-Generation Sequencing Reveals Differences between Primary and Secondary Ovarian Tumors in Formalin-Fixed, Paraffin-Embedded Tissue. Journal of Molecular Diagnostics, 2015, 17, 193-200.	2.8	8
24	Germline variants in POLE are associated with early onset mismatch repair deficient colorectal cancer. European Journal of Human Genetics, 2015, 23, 1080-1084.	2.8	101