Alessandro CAMA

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Blood Circulating CD133+ Extracellular Vesicles Predict Clinical Outcomes in Patients with Metastatic Colorectal Cancer. Cancers, 2022, 14, 1357.	3.7	13
2	Phenotypic and Proteomic Analysis Identifies Hallmarks of Blood Circulating Extracellular Vesicles in NSCLC Responders to Immune Checkpoint Inhibitors. Cancers, 2021, 13, 585.	3.7	25
3	Exosomes as Pleiotropic Players in Pancreatic Cancer. Biomedicines, 2021, 9, 275.	3.2	14
4	Screening of Benzimidazole-Based Anthelmintics and Their Enantiomers as Repurposed Drug Candidates in Cancer Therapy. Pharmaceuticals, 2021, 14, 372.	3.8	21
5	Protective Effects Induced by a Hydroalcoholic Allium sativum Extract in Isolated Mouse Heart. Nutrients, 2021, 13, 2332.	4.1	15
6	Drug Repurposing, an Attractive Strategy in Pancreatic Cancer Treatment: Preclinical and Clinical Updates. Cancers, 2021, 13, 3946.	3.7	15
7	The Role of Dysfunctional Adipose Tissue in Pancreatic Cancer: A Molecular Perspective. Cancers, 2020, 12, 1849.	3.7	20
8	Bridelia speciosa Müll.Arg. Stem bark Extracts as a Potential Biomedicine: From Tropical Western Africa to the Pharmacy Shelf. Antioxidants, 2020, 9, 128.	5.1	6
9	How Anesthetic, Analgesic and Other Non-Surgical Techniques During Cancer Surgery Might Affect Postoperative Oncologic Outcomes: A Summary of Current State of Evidence. Cancers, 2019, 11, 592.	3.7	50
10	The Benzimidazole-Based Anthelmintic Parbendazole: A Repurposed Drug Candidate That Synergizes with Gemcitabine in Pancreatic Cancer. Cancers, 2019, 11, 2042.	3.7	36
11	Paragangliomas arise through an autonomous vasculo-angio-neurogenic program inhibited by imatinib. Acta Neuropathologica, 2018, 135, 779-798.	7.7	20
12	Cytotoxic effect of a family of peroxisome proliferatorâ€activated receptor antagonists in colorectal and pancreatic cancer cell lines. Chemical Biology and Drug Design, 2017, 90, 1029-1035.	3.2	21
13	Effects of PPARα inhibition in head and neck paraganglioma cells. PLoS ONE, 2017, 12, e0178995.	2.5	30
14	Low AMY1 Gene Copy Number Is Associated with Increased Body Mass Index in Prepubertal Boys. PLoS ONE, 2016, 11, e0154961.	2.5	47
15	Overexpression of PY1289-HER3 in sporadic pulmonary carcinoid from patients bearing MEN1 gene variants. Oncology Letters, 2016, 12, 453-458.	1.8	1
16	Synthesis, inÂvitro evaluation, and molecular modeling investigation of benzenesulfonimide peroxisome proliferator-activated receptors α antagonists. European Journal of Medicinal Chemistry, 2016, 114, 191-200.	5.5	16
17	Correlation between mutations and mRNA expression of APC and MUTYH genes: new insight into hereditary colorectal polyposis predisposition. Journal of Experimental and Clinical Cancer Research, 2015, 34, 131.	8.6	19
18	Association between rs12970134 Near <i>MC4R</i> and Adiposity Indexes in a Homogenous Population of Caucasian Schoolchildren. Hormone Research in Paediatrics, 2014, 82, 187-193.	1.8	1

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19	Integrative genetic, epigenetic and pathological analysis of paraganglioma reveals complex dysregulation of NOTCH signaling. Acta Neuropathologica, 2013, 126, 575-594.	7.7	27
20	Novel insulin receptor substrate 1 and 2 variants in breast and colorectal cancer. Oncology Reports, 2013, 30, 1553-1560.	2.6	6
21	Integrative Analysis of Hereditary Nonpolyposis Colorectal Cancer: the Contribution of Allele-Specific Expression and Other Assays to Diagnostic Algorithms. PLoS ONE, 2013, 8, e81194.	2.5	9
22	Alterations of MEN1 and E-cadherin/β-catenin complex in sporadic pulmonary carcinoids. International Journal of Oncology, 2012, 41, 1221-8.	3.3	10
23	Increased Variance in Germline Allele-Specific Expression of APC Associates With Colorectal Cancer. Gastroenterology, 2012, 142, 71-77.e1.	1.3	24
24	Analysis of gene copy number variations using a method based on lab-on-a-chip technology. Tumori, 2012, 98, 126-36.	1.1	7
25	Transitions at CpG Dinucleotides, Geographic Clustering of TP53 Mutations and Food Availability Patterns in Colorectal Cancer. PLoS ONE, 2009, 4, e6824.	2.5	7
26	Nonfluorescent Denaturing HPLC–Based Primer-Extension Method for Allele-Specific Expression: Application to Analysis of Mismatch Repair Genes. Clinical Chemistry, 2009, 55, 1711-1718.	3.2	3
27	IRS1 G972R polymorphism and type 2 diabetes: a paradigm for the difficult ascertainment of the contribution to disease susceptibility of †low-frequency†"low-risk' variants. Diabetologia, 2009, 52, 1852-1857.	6.3	31
28	<i>P53</i> mutations in colorectal cancer from northern Iran: Relationships with site of tumor origin, microsatellite instability and Kâ€ <i>ras</i> mutations. Journal of Cellular Physiology, 2008, 216, 543-550.	4.1	23
29	Sporadic childhood hepatoblastomas show activation of β-catenin, mismatch repair defects and p53 mutations. Modern Pathology, 2008, 21, 7-14.	5.5	65
30	Methods for routine diagnosis of genomic rearrangements: multiplex PCR-based methods and future perspectives. Expert Review of Molecular Diagnostics, 2008, 8, 41-52.	3.1	8
31	Analysis of extended genomic rearrangements in oncological research. Annals of Oncology, 2007, 18, vi173-vi178.	1.2	11
32	Multicenter Comparative Multimodality Surveillance of Women at Genetic-Familial High Risk for Breast Cancer (HIBCRIT Study): Interim Results. Radiology, 2007, 242, 698-715.	7.3	324
33	High prevalence of BRCA1 deletions in BRCAPRO-positive patients with high carrier probability. Annals of Oncology, 2007, 18, vi86-vi92.	1.2	17
34	Balance between endoscopic and genetic information in the choice of ileorectal anastomosis for familial adenomatous polyposis. Journal of Surgical Oncology, 2007, 95, 28-33.	1.7	24
35	Obesity modifies the effects of the Asp905Tyr variant of PPP1R3A on risk of type 2 diabetes and insulin sensitivity. Diabetes, Obesity and Metabolism, 2007, 9, 759-761.	4.4	1
36	BRCA1 and BRCA2 status in a Central Sudanese series of breast cancer patients: interactions with genetic, ethnic and reproductive factors. Breast Cancer Research and Treatment, 2007, 102, 189-199.	2.5	55

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37	Variation of the insulin receptor substrate gene (IRS-1) in African Pygmies and Bantus. Diabetes Research and Clinical Practice, 2006, 72, 108-109.	2.8	1
38	Combined use of MLPA and nonfluorescent multiplex PCR analysis by high performance liquid chromatography for the detection of genomic rearrangements. Human Mutation, 2006, 27, 1047-1056.	2.5	20
39	Patterns of K-ras mutation in colorectal carcinomas from Iran and Italy (a Gruppo Oncologico) Tj ETQq1 1 0.7843 Annals of Oncology, 2006, 17, vii91-vii96.	14 rgBT / 1.2	Overlock 10 35
40	Genetic evidence that juvenile nasopharyngeal angiofibroma is an integral FAP tumour. Gut, 2005, 54, 1046-1047.	12.1	31
41	The CHEK2 c.1100delC mutation plays an irrelevant role in breast cancer predisposition in Italy. Human Mutation, 2004, 24, 100-101.	2.5	39
42	BRCA1 and BRCA2 mutations in breast/ovarian cancer patients from central Italy. Human Mutation, 2003, 22, 178-179.	2.5	32
43	TGF-β1 modulation of IGF-I signaling pathway in rat thyroid epithelial cells. Experimental Cell Research, 2003, 287, 411-423.	2.6	12
44	A Novel T608R Missense Mutation in Insulin Receptor Substrate-1 Identified in a Subject with Type 2 Diabetes Impairs Metabolic Insulin Signaling. Journal of Clinical Endocrinology and Metabolism, 2003, 88, 1468-1475.	3.6	45
45	Subcellular localization of theBRCA1 gene product in mitotic cells. Genes Chromosomes and Cancer, 2002, 35, 193-203.	2.8	64
46	Human Gene Mutations. Human Genetics, 2002, 110, 294-295.	3.8	0
47	Correlations between Phenotype and Microsatellite Instability in HNPCC: Implications for Genetic Testing. Familial Cancer, 2002, 3, 117-121.	1.9	8
48	Thyroid Carcinoma Usually Occurs in Patients with Familial Adenomatous Polyposis in the Absence of Biallelic Inactivation of the Adenomatous Polyposis Coli Gene. Journal of Clinical Endocrinology and Metabolism, 2001, 86, 427-432.	3.6	23
49	Germline Mutations of the APC Gene in Patients with Familial Adenomatous Polyposis-Associated Thyroid Carcinoma: Results from a European Cooperative Study. Journal of Clinical Endocrinology and Metabolism, 2000, 85, 286-292.	3.6	173
50	Microsatellite instability in thyroid tumours and tumour-like lesions. British Journal of Cancer, 1999, 79, 340-345.	6.4	35
51	Novel deletion at codon 1254 of the BRCA1 gene in an Italian breast cancer kindred. Human Mutation, 1998, 11, S237-S239.	2.5	4
52	Transcripts with splicings of exons 15 and 16 of the hMLH1 gene in normal lymphocytes: implications in RNA-based mutation screening of hereditary non-polyposis colorectal cancer. European Journal of Cancer, 1998, 34, 927-930.	2.8	12
53	Childhood hepatocellular tumors in FAP. Gastroenterology, 1997, 113, 1051-1052.	1.3	16
54	Multiplex PCR analysis and genotype-phenotype correlations of frequentAPC mutations. Human Mutation, 1995, 5, 144-152.	2.5	24

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55	Microsatellite instability and pathological aspects of breast cancer. International Journal of Cancer, 1995, 64, 264-268.	5.1	50
56	Two mutant alleles of the insulin receptor gene in a family with a genetic form of insulin resistance: a 10 base pair deletion in exon 1 and a mutation substituting serine for asparagine-462. Human Genetics, 1995, 95, 174-182.	3.8	22
57	Novel mutations and inactivation of both alleles of the APC gene in desmoid tumors. Human Molecular Genetics, 1995, 4, 1979-1981.	2.9	54
58	A novel deletion in exon 15 of the adenomatous polyposis coli gene in an Italian kindred. Human Mutation, 1994, 3, 301-304.	2.5	15
59	A novel mutation at the splice junction of exon 9 of the APC gene in familial adenomatous polyposis. Human Mutation, 1994, 3, 305-308.	2.5	11
60	Analysis of adenomatous polyposis coli gene in thyroid tumours. British Journal of Cancer, 1994, 70, 1085-1088.	6.4	47
61	Absence of insulin receptor gene mutations in three insulin-resistant women with the polycystic ovary syndrome. Metabolism: Clinical and Experimental, 1994, 43, 1568-1574.	3.4	63
62	Insulin resistance due to mutations of the insulin receptor gene: An overview. Journal of Endocrinological Investigation, 1992, 15, 857-864.	3.3	28
63	Postbinding characterization of five naturally occurring mutations in the human insulin receptor gene: impaired insulin-stimulated c-jun expression and thymidine incorporation despite normal receptor autophosphorylation. Biochemistry, 1992, 31, 9947-9954.	2.5	23
64	GENETIC BASIS OF ENDOCRINE DISEASE 1 Molecular Genetics of Insulin Resistant Diabetes Mellitus. Journal of Clinical Endocrinology and Metabolism, 1991, 73, 1158-1163.	3.6	54
65	Unusual Forms of Insulin Resistance. Annual Review of Medicine, 1991, 42, 373-379.	12.2	19
66	A Mutation in the Tyrosine Kinase Domain of the Insulin Receptor Associated with Insulin Resistance in an Obese Woman*. Journal of Clinical Endocrinology and Metabolism, 1991, 73, 894-901.	3.6	77
67	Two Unrelated Patients with Familial Hyperproinsulinemia due to a Mutation Substituting Histidine for Arginine at Position 65 in the Proinsulin Molecule: Identification of the Mutation by Direct Sequencing of Genomic Deoxyribonucleic Acid Amplified by Polymerase Chain Reaction. Journal of Clinical Endocrinology and Metabolism, 1990, 71, 164-169.	3.6	35
68	The Amino Acid Sequence of the Insulin Receptor Is Normal in an Insulin-Resistant Pima Indian*. Journal of Clinical Endocrinology and Metabolism, 1990, 70, 1155-1166.	3.6	54
69	Mutations in Insulin-Receptor Gene in Insulin-Resistant Patients. Diabetes Care, 1990, 13, 257-279.	8.6	219
70	Mutations of the human insulin receptor gene. Trends in Endocrinology and Metabolism, 1990, 1, 134-139.	7.1	13
71	Immunological Abnormalities in Insulin Receptors on Cultured EBV-Transformed Lymphocytes From Insulin-Resistant Patient With Leprechaunism. Diabetes, 1988, 37, 982-988.	0.6	30
72	Anti-endothelial cell antibodies: detection and characterization in sera from patients with autoimmune hypoparathyroidism Proceedings of the National Academy of Sciences of the United States of America, 1988, 85, 4015-4019.	7.1	51

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73	Tyrosine kinase activity of insulin receptors from an insulin-resistant patient with leprechaunism. Diabetologia, 1987, 30, 631-637.	6.3	14