Cristina Rodriguez-Antona

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Influence of cytochrome P450 polymorphisms on drug therapies: Pharmacogenetic, pharmacoepigenetic and clinical aspects. , 2007, 116, 496-526.		990
2	Cytochrome P450 pharmacogenetics and cancer. Oncogene, 2006, 25, 1679-1691.	5.9	492
3	Exome sequencing identifies MAX mutations as a cause of hereditary pheochromocytoma. Nature Genetics, 2011, 43, 663-667.	21.4	478
4	Identification of 12 new susceptibility loci for different histotypes of epithelial ovarian cancer. Nature Genetics, 2017, 49, 680-691.	21.4	356
5	Cytochrome P450 expression in human hepatocytes and hepatoma cell lines: molecular mechanisms that determine lower expression in cultured cells. Xenobiotica, 2002, 32, 505-520.	1.1	340
6	Tumoral and tissueâ€specific expression of the major human βâ€ŧubulin isotypes. Cytoskeleton, 2010, 67, 214-223.	2.0	221
7	Single nucleotide polymorphism associations with response and toxic effects in patients with advanced renal-cell carcinoma treated with first-line sunitinib: a multicentre, observational, prospective study. Lancet Oncology, The, 2011, 12, 1143-1150.	10.7	217
8	The miR-200 family controls Â-tubulin III expression and is associated with paclitaxel-based treatment response and progression-free survival in ovarian cancer patients. Endocrine-Related Cancer, 2010, 18, 85-95.	3.1	188
9	Research Resource: Transcriptional Profiling Reveals Different Pseudohypoxic Signatures in SDHB and VHL-Related Pheochromocytomas. Molecular Endocrinology, 2010, 24, 2382-2391.	3.7	179
10	Whole-Exome Sequencing Identifies MDH2 as a New Familial Paraganglioma Gene. Journal of the National Cancer Institute, 2015, 107, .	6.3	143
11	Tumoral EPAS1 (HIF2A) mutations explain sporadic pheochromocytoma and paraganglioma in the absence of erythrocytosis. Human Molecular Genetics, 2013, 22, 2169-2176.	2.9	142
12	The Variant rs1867277 in FOXE1 Gene Confers Thyroid Cancer Susceptibility through the Recruitment of USF1/USF2 Transcription Factors. PLoS Genetics, 2009, 5, e1000637.	3.5	140
13	Cytochrome P-450 mRNA Expression in Human Liver and Its Relationship with Enzyme Activity. Archives of Biochemistry and Biophysics, 2001, 393, 308-315.	3.0	129
14	Long-term expression of differentiated functions in hepatocytes cultured in three-dimensional collagen matrix. , 1998, 177, 553-562.		125
15	Genetics of Pheochromocytoma and Paraganglioma in Spanish Patients. Journal of Clinical Endocrinology and Metabolism, 2009, 94, 1701-1705.	3.6	120
16	Polymorphisms in cytochromes P450 2C8 and 3A5 are associated with paclitaxel neurotoxicity. Pharmacogenomics Journal, 2011, 11, 121-129.	2.0	112
17	Pazopanib in pretreated advanced neuroendocrine tumors: a phase II, open-label trial of the Spanish Task Force Group for Neuroendocrine Tumors (GETNE). Annals of Oncology, 2015, 26, 1987-1993.	1.2	112
18	Molecular genetics and epigenetics of the cytochrome P450 gene family and its relevance for cancer risk and treatment. Human Genetics, 2010, 127, 1-17.	3.8	110

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19	Overexpression and activation of EGFR and VEGFR2 in medullary thyroid carcinomas is related to metastasis. Endocrine-Related Cancer, 2010, 17, 7-16.	3.1	108
20	Pharmacogenetics of drug-metabolizing enzymes: implications for a safer and more effective drug therapy. Philosophical Transactions of the Royal Society B: Biological Sciences, 2005, 360, 1563-1570.	4.0	106
21	Recommendations for somatic and germline genetic testing of single pheochromocytoma and paraganglioma based on findings from a series of 329 patients. Journal of Medical Genetics, 2015, 52, 647-656.	3.2	102
22	Tumor-specific expression of the novel cytochrome P450 enzyme, CYP2W1. Biochemical and Biophysical Research Communications, 2006, 341, 451-458.	2.1	98
23	Transcriptional Regulation of Human CYP3A4 Basal Expression by CCAAT Enhancer-Binding Protein α and Hepatocyte Nuclear Factor-3γ. Molecular Pharmacology, 2003, 63, 1180-1189.	2.3	97
24	Quantitative RT-PCR Measurement of Human Cytochrome P-450s: Application to Drug Induction Studies. Archives of Biochemistry and Biophysics, 2000, 376, 109-116.	3.0	93
25	Phenotype–genotype variability in the human CYP3A locus as assessed by the probe drug quinine and analyses of variant CYP3A4 alleles. Biochemical and Biophysical Research Communications, 2005, 338, 299-305.	2.1	93
26	Identification of Tissue microRNAs Predictive of Sunitinib Activity in Patients with Metastatic Renal Cell Carcinoma. PLoS ONE, 2014, 9, e86263.	2.5	76
27	CYP3A5 and ABCB1 Polymorphisms as Predictors for Sunitinib Outcome in Metastatic Renal Cell Carcinoma. European Urology, 2015, 68, 621-629.	1.9	75
28	GrossSDHB deletions in patients with paraganglioma detected by multiplex PCR: A possible hot spot?. Genes Chromosomes and Cancer, 2006, 45, 213-219.	2.8	73
29	Prospective study assessing hypoxia-related proteins as markers for the outcome of treatment with sunitinib in advanced clear-cell renal cell carcinoma. Annals of Oncology, 2013, 24, 2409-2414.	1.2	73
30	Gain-of-function mutations in DNMT3A in patients with paraganglioma. Genetics in Medicine, 2018, 20, 1644-1651.	2.4	73
31	Prognostic and Predictive Value of PBRM1 in Clear Cell Renal Cell Carcinoma. Cancers, 2020, 12, 16.	3.7	72
32	Deregulated miRNAs in Hereditary Breast Cancer Revealed a Role for miR-30c in Regulating KRAS Oncogene. PLoS ONE, 2012, 7, e38847.	2.5	71
33	Molecular characterisation of a common SDHB deletion in paraganglioma patients. Journal of Medical Genetics, 2007, 45, 233-238.	3.2	69
34	Genome-wide association study identifies ephrin type A receptors implicated in paclitaxel induced peripheral sensory neuropathy. Journal of Medical Genetics, 2013, 50, 599-605.	3.2	67
35	Novel copy-number variations in pharmacogenes contribute to interindividual differences in drug pharmacokinetics. Genetics in Medicine, 2018, 20, 622-629.	2.4	66
36	PheoSeq. Journal of Molecular Diagnostics, 2017, 19, 575-588.	2.8	63

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37	Expression and induction of a large set of drug-metabolizing enzymes by the highly differentiated human hepatoma cell line BC2. FEBS Journal, 2001, 268, 1448-1459.	0.2	62
38	Regulatory Polymorphisms in β-Tubulin IIa Are Associated with Paclitaxel-Induced Peripheral Neuropathy. Clinical Cancer Research, 2012, 18, 4441-4448.	7.0	61
39	Whole-Exome Sequencing Reveals Defective <i>CYP3A4</i> Variants Predictive of Paclitaxel Dose-Limiting Neuropathy. Clinical Cancer Research, 2015, 21, 322-328.	7.0	61
40	Characterization of novel CYP2C8 haplotypes and their contribution to paclitaxel and repaglinide metabolism. Pharmacogenomics Journal, 2008, 8, 268-277.	2.0	59
41	Pharmacogenomic biomarkers for personalized cancer treatment. Journal of Internal Medicine, 2015, 277, 201-217.	6.0	57
42	Effect of the Most Relevant CYP3A4 and CYP3A5 Polymorphisms on the Pharmacokinetic Parameters of 10 CYP3A Substrates. Biomedicines, 2020, 8, 94.	3.2	57
43	A Transcriptome-Wide Association Study Among 97,898 Women to Identify Candidate Susceptibility Genes for Epithelial Ovarian Cancer Risk. Cancer Research, 2018, 78, 5419-5430.	0.9	54
44	Integrative multi-omics analysis identifies a prognostic miRNA signature and a targetable miR-21-3p/TSC2/mTOR axis in metastatic pheochromocytoma/paraganglioma. Theranostics, 2019, 9, 4946-4958.	10.0	54
45	DNA Methylation Profiling in Pheochromocytoma and Paraganglioma Reveals Diagnostic and Prognostic Markers. Clinical Cancer Research, 2015, 21, 3020-3030.	7.0	53
46	Identification and phenotype characterization of two haplotypes causing different enzymatic capacity in fetal livers. Clinical Pharmacology and Therapeutics, 2005, 77, 259-270.	4.7	52
47	Integrative analysis of miRNA and mRNA expression profiles in pheochromocytoma and paraganglioma identifies genotype-specific markers and potentially regulated pathways. Endocrine-Related Cancer, 2013, 20, 477-493.	3.1	52
48	Recurrent Germline DLST Mutations in Individuals with Multiple Pheochromocytomas and Paragangliomas. American Journal of Human Genetics, 2019, 104, 651-664.	6.2	51
49	Clinical Pharmacogenetics Implementation Consortium Guideline for the Use of Aminoglycosides Based on <i>MTâ€RNR1</i> Genotype. Clinical Pharmacology and Therapeutics, 2022, 111, 366-372.	4.7	50
50	Genetic Data from Nearly 63,000 Women of European Descent Predicts DNA Methylation Biomarkers and Epithelial Ovarian Cancer Risk. Cancer Research, 2019, 79, 505-517.	0.9	49
51	High frequency and founder effect of the CYP3A4*20 loss-of-function allele in the Spanish population classifies CYP3A4 as a polymorphic enzyme. Pharmacogenomics Journal, 2015, 15, 288-292.	2.0	48
52	Role of MDH2 pathogenic variant in pheochromocytoma and paraganglioma patients. Genetics in Medicine, 2018, 20, 1652-1662.	2.4	45
53	Transcriptional Regulation of the Human CYP2A6 Gene. Journal of Pharmacology and Experimental Therapeutics, 2005, 313, 814-822.	2.5	44
54	Improving pharmacovigilance in Europe: TPMT genotyping and phenotyping in the UK and Spain. European Journal of Human Genetics, 2009, 17, 991-998.	2.8	43

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55	Pharmacogenomics of paclitaxel. Pharmacogenomics, 2010, 11, 621-623.	1.3	43
56	Development and Validation of the Gene Expression Predictor of High-grade Serous Ovarian Carcinoma Molecular SubTYPE (PrOTYPE). Clinical Cancer Research, 2020, 26, 5411-5423.	7.0	43
57	Expression of CYP3A4 as a predictor of response to chemotherapy in peripheral T-cell lymphomas. Blood, 2007, 110, 3345-3351.	1.4	42
58	Loss of the actin regulator HSPC300 results in clear cell renal cell carcinoma protection in Von Hippel-Lindau patients. Human Mutation, 2007, 28, 613-621.	2.5	41
59	Rationalization of Genetic Testing in Patients with Apparently Sporadic Pheochromocytoma/Paraganglioma. Hormone and Metabolic Research, 2009, 41, 672-675.	1.5	41
60	Detection of the first gross CDC73 germline deletion in an HPTâ€JT syndrome family. Genes Chromosomes and Cancer, 2011, 50, 922-929.	2.8	41
61	Oxaliplatin induced-neuropathy in digestive tumors. Critical Reviews in Oncology/Hematology, 2014, 89, 166-178.	4.4	40
62	Deep sequencing reveals microRNAs predictive of antiangiogenic drug response. JCI Insight, 2016, 1, e86051.	5.0	39
63	<i>SDHC</i> mutation in an elderly patient without familial antecedents. Clinical Endocrinology, 2008, 69, 906-910.	2.4	37
64	Allelic variant at â^'79 (C>T) in CDKN1B (p27Kip1) confers an increased risk of thyroid cancer and alters mRNA levels. Endocrine-Related Cancer, 2010, 17, 317-328.	3.1	35
65	Differential Gene Expression of Medullary Thyroid Carcinoma Reveals Specific Markers Associated with Genetic Conditions. American Journal of Pathology, 2013, 182, 350-362.	3.8	35
66	IL8 polymorphisms and overall survival in pazopanib- or sunitinib-treated patients with renal cell carcinoma. British Journal of Cancer, 2015, 112, 1190-1198.	6.4	35
67	Clinical and pathological associations of PTEN expression in ovarian cancer: a multicentre study from the Ovarian Tumour Tissue Analysis Consortium. British Journal of Cancer, 2020, 123, 793-802.	6.4	35
68	Cytochrome P450 3A5 is highly expressed in normal prostate cells but absent in prostate cancer. Endocrine-Related Cancer, 2007, 14, 645-654.	3.1	34
69	<i>CYP2D6</i> genotyping for psychiatric patients treated with risperidone: considerations for cost–effectiveness studies. Pharmacogenomics, 2009, 10, 685-699.	1.3	34
70	Cytochrome P450 2D6 Genotyping. CNS Drugs, 2009, 23, 181-191.	5.9	33
71	A nicotineC-oxidase gene (CYP2A6) polymorphism important for promoter activity. Human Mutation, 2004, 23, 258-266.	2.5	32
72	Hsaâ€miRâ€139â€5p is a prognostic thyroid cancer marker involved in HNRNPFâ€mediated alternative splicing. International Journal of Cancer, 2020, 146, 521-530.	5.1	29

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73	Molecular characterization of chromophobe renal cell carcinoma reveals mTOR pathway alterations in patients with poor outcome. Modern Pathology, 2020, 33, 2580-2590.	5.5	29
74	Polymorphic NF-Y dependent regulation of human nicotine C-oxidase (CYP2A6). Pharmacogenetics and Genomics, 2004, 14, 369-379.	5.7	28
75	Genetic variation in the <i>SLC19A1</i> gene and methotrexate toxicity in rheumatoid arthritis patients. Pharmacogenomics, 2012, 13, 1583-1594.	1.3	27
76	Impact of chemotherapy on telomere length in sporadic and familial breast cancer patients. Breast Cancer Research and Treatment, 2015, 149, 385-394.	2.5	27
77	Polymorphisms associated with everolimus pharmacokinetics, toxicity and survival in metastatic breast cancer. PLoS ONE, 2017, 12, e0180192.	2.5	27
78	Hematologic Î ² -Tubulin VI Isoform Exhibits Genetic Variability That Influences Paclitaxel Toxicity. Cancer Research, 2012, 72, 4744-4752.	0.9	26
79	Functional and in silico assessment of MAX variants of unknown significance. Journal of Molecular Medicine, 2015, 93, 1247-1255.	3.9	25
80	Genetic polymorphisms of SCN9A are associated with oxaliplatin-induced neuropathy. BMC Cancer, 2017, 17, 63.	2.6	25
81	Collaboration with other firms and customers: innovation's secret weapon. Strategy and Leadership, 2004, 32, 16-20.	0.5	24
82	Polygenic risk modeling for prediction of epithelial ovarian cancer risk. European Journal of Human Genetics, 2022, 30, 349-362.	2.8	23
83	A Novel Polymorphic Cytochrome P450 Formed by Splicing of CYP3A7 and the Pseudogene CYP3AP1. Journal of Biological Chemistry, 2005, 280, 28324-28331.	3.4	22
84	Impact of genetic polymorphisms in <i>CYP2C8</i> and rosiglitazone intake on the urinary excretion of dihydroxyeicosatrienoic acids. Pharmacogenomics, 2008, 9, 277-288.	1.3	22
85	Sunitinib-induced hypertension in CYP3A4 rs4646437 A-allele carriers with metastatic renal cell carcinoma. Pharmacogenomics Journal, 2017, 17, 42-46.	2.0	21
86	Concomitant Medications and Risk of Chemotherapy-Induced Peripheral Neuropathy. Oncologist, 2019, 24, e784-e792.	3.7	20
87	Pâ€Glycoprotein Inhibition Exacerbates Paclitaxel Neurotoxicity in Neurons and Patients With Cancer. Clinical Pharmacology and Therapeutics, 2020, 108, 671-680.	4.7	20
88	Association of single nucleotide polymorphisms in IL8 and IL13 with sunitinib-induced toxicity in patients with metastatic renal cell carcinoma. European Journal of Clinical Pharmacology, 2015, 71, 1477-1484.	1.9	19
89	The hematopoietic-specific l²1-tubulin is naturally resistant to 2-Methoxyestradiol and protects patients from drug-induced myelosuppression. Cell Cycle, 2009, 8, 3914-3924.	2.6	18
90	Molecular Markers to Predict Response to Therapy. Seminars in Oncology, 2013, 40, 444-458.	2.2	18

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91	VEGF, VEGFR3, and PDGFRB Protein Expression Is Influenced by <i>RAS</i> Mutations in Medullary Thyroid Carcinoma. Thyroid, 2014, 24, 1251-1255.	4.5	18
92	Biallelic <i>TSC2</i> Mutations in a Patient With Chromophobe Renal Cell Carcinoma Showing Extraordinary Response to Temsirolimus. Journal of the National Comprehensive Cancer Network: JNCCN, 2018, 16, 352-358.	4.9	18
93	Influence of RET mutations on the expression of tyrosine kinases in medullary thyroid carcinoma. Endocrine-Related Cancer, 2013, 20, 611-619.	3.1	17
94	Role of cytochrome P450 <i>2C8*3</i> (<i>CYP2C8*3</i>) in paclitaxel metabolism and paclitaxel-induced neurotoxicity. Pharmacogenomics, 2015, 16, 929-937.	1.3	17
95	Exceptional Response to Temsirolimus in a Metastatic Clear Cell Renal Cell Carcinoma With an Early Novel MTOR -Activating Mutation. Journal of the National Comprehensive Cancer Network: JNCCN, 2017, 15, 1310-1315.	4.9	16
96	Targeted Sequencing Reveals Low-Frequency Variants in <i>EPHA</i> Genes as Markers of Paclitaxel-Induced Peripheral Neuropathy. Clinical Cancer Research, 2017, 23, 1227-1235.	7.0	16
97	A Pilot, Phase II, Randomized, Open-Label Clinical Trial Comparing the Neurotoxicity of Three Dose Regimens of Nab-Paclitaxel to That of Solvent-Based Paclitaxel as the First-Line Treatment for Patients with Human Epidermal Growth Factor Receptor Type 2-Negative Metastatic Breast Cancer. Oncologist, 2019, 24, e1024-e1033.	3.7	16
98	MicroRNAs Targeting HIF-2α, VEGFR1 and/or VEGFR2 as Potential Predictive Biomarkers for VEGFR Tyrosine Kinase and HIF-2α Inhibitors in Metastatic Clear-Cell Renal Cell Carcinoma. Cancers, 2021, 13, 3099.	3.7	16
99	Identification of candidate SNPs for drug induced toxicity from differentially expressed genes in associated tissues. Gene, 2012, 506, 62-68.	2.2	15
100	Assessment of moderate coffee consumption and risk of epithelial ovarian cancer: a Mendelian randomization study. International Journal of Epidemiology, 2018, 47, 450-459.	1.9	15
101	Determination of CYP2D6 gene copy number by multiplex polymerase chain reaction analysis. Analytical Biochemistry, 2009, 389, 74-76.	2.4	14
102	Association between SLC19A1 gene polymorphism and high dose methotrexate toxicity in childhood acute lymphoblastic leukaemia and non Hodgkin malignant lymphoma: introducing a haplotype based approach. Radiology and Oncology, 2017, 51, 455-462.	1.7	14
103	PTEN expression and mutations in TSC1 , TSC2 and MTOR are associated with response to rapalogs in patients with renal cell carcinoma. International Journal of Cancer, 2020, 146, 1435-1444.	5.1	14
104	Analysis of Telomere Maintenance Related Genes Reveals NOP10 as a New Metastatic-Risk Marker in Pheochromocytoma/Paraganglioma. Cancers, 2021, 13, 4758.	3.7	14
105	Influence of donor liver CYP3A4*20 loss-of-function genotype on tacrolimus pharmacokinetics in transplanted patients. Pharmacogenetics and Genomics, 2018, 28, 41-48.	1.5	13
106	Advanced sporadic renal epithelioid angiomyolipoma: case report of an extraordinary response to sirolimus linked to TSC2 mutation. BMC Cancer, 2018, 18, 561.	2.6	13
107	Progress in pharmacogenetics: consortiums and new strategies. Drug Metabolism and Personalized Therapy, 2016, 31, 17-23.	0.6	12
108	Prognostic and predictive biomarkers for somatostatin analogs, peptide receptor radionuclide therapy and serotonin pathway targets in neuroendocrine tumours. Cancer Treatment Reviews, 2018, 70, 209-222.	7.7	12

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109	Sunitinib and Evofosfamide (<scp>TH</scp> -302) in Systemic Treatment-NaÃ ⁻ ve Patients with Grade 1/2 Metastatic Pancreatic Neuroendocrine Tumors: The <scp>GETNE</scp> -1408 Trial. Oncologist, 2021, 26, 941-949.	3.7	12
110	Constitutional genetic variants as predictors of antiangiogenic therapy outcome in renal cell carcinoma. Pharmacogenomics, 2012, 13, 1621-1633.	1.3	11
111	Renal carcinoma pharmacogenomics and predictors of response: Steps toward treatment individualization. Urologic Oncology: Seminars and Original Investigations, 2015, 33, 179-186.	1.6	10
112	Human genetics: international projects and personalized medicine. Drug Metabolism and Personalized Therapy, 2016, 31, 3-8.	0.6	10
113	Circulating Levels of the Interferon-γ-Regulated Chemokines CXCL10/CXCL11, IL-6 and HGF Predict Outcome in Metastatic Renal Cell Carcinoma Patients Treated with Antiangiogenic Therapy. Cancers, 2021, 13, 2849.	3.7	10
114	Clinical pharmacogenetic analysis in 5,001 individuals with diagnostic Exome Sequencing data. Npj Genomic Medicine, 2022, 7, 12.	3.8	10
115	Replication of Genetic Polymorphisms Reported to Be Associated with Taxane-Related Sensory Neuropathy in Patients with Early Breast Cancer Treated with Paclitaxel—Letter. Clinical Cancer Research, 2015, 21, 3092-3093.	7.0	9
116	Description of the EuroTARGET cohort: A European collaborative project on TArgeted therapy in renal cell cancer—GEnetic- and tumor-related biomarkers for response and toxicity. Urologic Oncology: Seminars and Original Investigations, 2017, 35, 529.e9-529.e16.	1.6	9
117	MicroRNAs Possibly Involved in the Development of Bone Metastasis in Clear-Cell Renal Cell Carcinoma. Cancers, 2021, 13, 1554.	3.7	9
118	Prevalence of pathogenic germline variants in patients with metastatic renal cell carcinoma. Genetics in Medicine, 2021, 23, 698-704.	2.4	9
119	SNPs associated with activity and toxicity of cabazitaxel in patients with advanced urothelial cell carcinoma. Pharmacogenomics, 2016, 17, 463-471.	1.3	8
120	A Genetic Polymorphism in <i>CTLA-4</i> Is Associated with Overall Survival in Sunitinib-Treated Patients with Clear Cell Metastatic Renal Cell Carcinoma. Clinical Cancer Research, 2018, 24, 2350-2356.	7.0	7
121	Genes and variants in hematopoiesis-related pathways are associated with gemcitabine/carboplatin-induced thrombocytopenia. Pharmacogenomics Journal, 2020, 20, 179-191.	2.0	7
122	Meta-analysis on the association of <i>VEGFR1</i> genetic variants with sunitinib outcome in metastatic renal cell carcinoma patients. Oncotarget, 2017, 8, 1204-1212.	1.8	6
123	Evaluation of KDR rs34231037 as a predictor of sunitinib efficacy in patients with metastatic renal cell carcinoma. Pharmacogenetics and Genomics, 2017, 27, 227-231.	1.5	5
124	Genetic association of gemcitabine/carboplatin-induced leukopenia and neutropenia in non-small cell lung cancer patients using whole-exome sequencing. Lung Cancer, 2020, 147, 106-114.	2.0	5
125	Novel DNMT3A Germline Variant in a Patient with Multiple Paragangliomas and Papillary Thyroid Carcinoma. Cancers, 2020, 12, 3304.	3.7	5
126	CD133 Expression in Medullary Thyroid Cancer Cells Identifies Patients with Poor Prognosis. Journal of Clinical Endocrinology and Metabolism, 2020, 105, 3548-3561.	3.6	5

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127	MicroRNA expression profiles in molecular subtypes of clear-cell renal cell carcinoma are associated with clinical outcome and repression of specific mRNA targets. PLoS ONE, 2020, 15, e0238809.	2.5	5
128	A Prospective Observational Study for Assessment and Outcome Association of Circulating Endothelial Cells in Clear Cell Renal Cell Carcinoma Patients Who Show Initial Benefit from First-line Treatment. The CIRCLES (CIRCuLating Endothelial cellS) Study (SOGUG-CEC-2011-01). European Urology Focus, 2017, 3, 430-436.	3.1	4
129	Overexpression of miRâ€483â€5p is confined to metastases and linked to high circulating levels in patients with metastatic pheochromocytoma/paraganglioma. Clinical and Translational Medicine, 2020, 10, e260.	4.0	4
130	mTOR Pathway Mutations and Response to Rapalogs in RCC—Letter. Clinical Cancer Research, 2017, 23, 5320-5320.	7.0	3
131	Impact of the combination of durvalumab (MEDI4736) plus olaparib (AZD2281) administered prior to surgery in the molecular profile of resectable urothelial bladder cancer: NEODURVARIB Trial Journal of Clinical Oncology, 2019, 37, TPS503-TPS503.	1.6	3
132	The role of pharmacogenetics and pharmacogenomics in 21st-century medicine: state of the art and new challenges discussed in the VII Conference of the Spanish Pharmacogenetics and Pharmacogenomics Society (SEFF). Drug Metabolism and Personalized Therapy, 2016, 31, 1-2.	0.6	2
133	MicroRNAs as potential predictors of extreme response to tyrosine kinase inhibitors in renal cell cancer. Urologic Oncology: Seminars and Original Investigations, 2020, 38, 640.e23-640.e29.	1.6	2
134	Genome-Wide Meta-Analysis Identifies Variants in DSCAM and PDLIM3 That Correlate with Efficacy Outcomes in Metastatic Renal Cell Carcinoma Patients Treated with Sunitinib. Cancers, 2022, 14, 2838.	3.7	1
135	Microtubule-targeting drugs and personalization of cancer treatment. Pharmacogenomics, 2011, 12, 449-451.	1.3	Ο
136	A Novel Approach for the Identification of Pharmacogenetic Variants in MT-RNR1 through Next-Generation Sequencing Off-Target Data. Journal of Clinical Medicine, 2020, 9, 2082.	2.4	0
137	Abstract P3-07-40: Pharmacogenetic study of exemestane and everolimus in metastatic breast cancer patients progressing on prior non-steroidal aromatase inhibitors. , 2016, , .		0
138	Title is missing!. , 2020, 15, e0238809.		0
139	Title is missing!. , 2020, 15, e0238809.		Ο
140	Title is missing!. , 2020, 15, e0238809.		0
141	Title is missing!. , 2020, 15, e0238809.		0
142	The need of the clinical implementation of pharmacogenetics in European health services for routine drug prescription. What's next? An urgent clinical unmet need for patients. Drug Metabolism and Drug Interactions, 2020, 35, .	0.3	0