

Adrian Pavel Trifa

List of Publications by Year in descending order

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67
papers

772
citations

516215

16
h-index

642321

23
g-index

68
all docs

68
docs citations

68
times ranked

1211
citing authors

#	ARTICLE	IF	CITATIONS
1	Genetic determination of irritable bowel syndrome. <i>World Journal of Gastroenterology</i> , 2008, 14, 6636.	1.4	42
2	How to Diagnose and Treat a Cancer of Unknown Primary Site. <i>Journal of Gastrointestinal and Liver Diseases</i> , 2020, 26, 69-79.	0.5	42
3	<i>CAT, GPX1, MnSOD, GSTM1, GSTT1</i>, and <i>GSTP1</i> Genetic Polymorphisms in Chronic Myeloid Leukemia: A Case-Control Study. <i>Oxidative Medicine and Cellular Longevity</i> , 2014, 2014, 1-6.	1.9	35
4	The role of medical registries, potential applications and limitations. <i>Medicine and Pharmacy Reports</i> , 2019, 92, 7-14.	0.2	33
5	<i>TERT</i> rs2736100 A>C SNP and <i>JAK2</i> 46/1 haplotype significantly contribute to the occurrence of <i>JAK2</i> V617F and <i>CALR</i> mutated myeloproliferative neoplasms â€œ a multicentric study on 529 patients. <i>British Journal of Haematology</i> , 2016, 174, 218-226.	1.2	32
6	<i>MECOM</i>, <i>HBS1Lâ€MYB</i>, <i>THRBâ€RARB, JAK2</i>, and <i>TERT</i> polymorphisms defining the genetic predisposition to myeloproliferative neoplasms: A study on 939 patients. <i>American Journal of Hematology</i> , 2018, 93, 100-106.	2.0	30
7	Exosome-carried microRNA-based signature as a cellular trigger for the evolution of chronic lymphocytic leukemia into Richter syndrome. <i>Critical Reviews in Clinical Laboratory Sciences</i> , 2018, 55, 501-515.	2.7	27
8	The G allele of the JAK2 rs10974944 SNP, part of JAK2 46/1 haplotype, is strongly associated with JAK2 V617F-positive myeloproliferative neoplasms. <i>Annals of Hematology</i> , 2010, 89, 979-983.	0.8	25
9	Polymorphism of <i>XRCC1</i>, <i>XRCC3</i>, and <i>XPD</i> Genes and Risk of Chronic Myeloid Leukemia. <i>BioMed Research International</i> , 2014, 2014, 1-9.	0.9	25
10	From Six Gene Polymorphisms of the Antioxidant System, Only GPX Pro198Leu and GSTP1 Ile105Val Modulate the Risk of Acute Myeloid Leukemia. <i>Oxidative Medicine and Cellular Longevity</i> , 2016, 2016, 1-10.	1.9	25
11	Prevalence of the c.35delG and p.W24X mutations in the GJB2 gene in patients with nonsyndromic hearing loss from North-West Romania. <i>International Journal of Pediatric Otorhinolaryngology</i> , 2010, 74, 351-355.	0.4	22
12	Analysis of <i>CYP2C9*2</i>, <i>CYP2C9*3</i> and <i>VKORC1</i> polymorphisms in a population from South-Eastern Europe. <i>Journal of Cellular and Molecular Medicine</i> , 2012, 16, 2919-2924.	1.6	22
13	<i>XRCC1</i> Arg194Trp and Arg399Gln polymorphisms are significantly associated with shorter survival in acute myeloid leukemia. <i>Leukemia and Lymphoma</i> , 2014, 55, 365-370.	0.6	20
14	Influence of XPC, XPD, XPF, and XPG gene polymorphisms on the risk and the outcome of acute myeloid leukemia in a Romanian population. <i>Tumor Biology</i> , 2016, 37, 9357-9366.	0.8	20
15	Vitamin D receptor polymorphisms and melanoma (Review). <i>Oncology Letters</i> , 2019, 17, 4162-4169.	0.8	20
16	Phytochemicals in Cardiovascular and Respiratory Diseases: Evidence in Oxidative Stress and Inflammation. <i>Oxidative Medicine and Cellular Longevity</i> , 2018, 2018, 1-3.	1.9	19
17	An acenocoumarol dose algorithm based on a South-Eastern European population. <i>European Journal of Clinical Pharmacology</i> , 2013, 69, 1901-1907.	0.8	18
18	Interindividual Variability of Apixaban Plasma Concentrations: Influence of Clinical and Genetic Factors in a Real-Life Cohort of Atrial Fibrillation Patients. <i>Genes</i> , 2020, 11, 438.	1.0	17

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19	Cytokine rs361525, rs1800750, rs1800629, rs1800896, rs1800872, rs1800795, rs1800470, and rs2430561 SNPs in relation with prognostic factors in acute myeloid leukemia. <i>Cancer Medicine</i> , 2019, 8, 5492-5506.	1.3	16
20	<i><sc>CALR</sc> versus <sc>JAK</sc>2</i> mutated essential thrombocythaemia – a report on 141 patients. <i>British Journal of Haematology</i> , 2015, 168, 151-153.	1.2	15
21	Development of a Reliable PCR-RFLP Assay for Investigation of the JAK2 rs10974944 SNP, Which Might Predispose to the Acquisition of Somatic Mutation JAK2^{V617F}. <i>Acta Haematologica</i> , 2010, 123, 84-87.	0.7	14
22	The impact of the CYP2C9 and VKORC1 polymorphisms on acenocoumarol dose requirements in a Romanian population. <i>Blood Cells, Molecules, and Diseases</i> , 2013, 50, 166-170.	0.6	14
23	Absence of BRAF V600E mutation in a cohort of 402 patients with various chronic and acute myeloid neoplasms. <i>Leukemia and Lymphoma</i> , 2012, 53, 2496-2497.	0.6	13
24	The relationship between factor V Leiden, prothrombin G20210A, and MTHFR mutations and the first major thrombotic episode in polycythemia vera and essential thrombocythemia. <i>Annals of Hematology</i> , 2014, 93, 203-209.	0.8	13
25	Persistent Basophilia May Suggest an “Accelerated Phase” in the Evolution of CALR-Positive Primary Myelofibrosis Toward Acute Myeloid Leukemia. <i>Frontiers in Oncology</i> , 2019, 9, 872.	1.3	12
26	: Valuable perspective in the management of chronic diseases. <i>Iranian Journal of Basic Medical Sciences</i> , 2020, 23, 699-713.	1.0	11
27	PAI-1 4G/5G and MTHFR C677T polymorphisms increased the accuracy of two prediction scores for the risk of acute lower extremity deep vein thrombosis. <i>Romanian Journal of Morphology and Embryology</i> , 2014, 55, 153-7.	0.4	11
28	Association of 276G>T adiponectin gene polymorphism to plasma adiponectin and albuminuria in type 2 diabetic patients. <i>International Urology and Nephrology</i> , 2012, 44, 1771-1777.	0.6	10
29	Among a panel of polymorphisms in genes related to oxidative stress, <i>CAT</i>-262 C>T, <i>GPX1</i> Pro198Leu and <i>GSTP1</i> Ile105Val influence the risk of developing <i>BCR-ABL</i> negative myeloproliferative neoplasms. <i>Hematology</i> , 2016, 21, 520-525.	0.7	9
30	Adult acute megakaryoblastic leukemia: rare association with cytopenias of undetermined significance and p210 and p190^{BCR–ABL} transcripts. <i>OncoTargets and Therapy</i> , 2017, Volume 10, 5047-5051.	1.0	9
31	Fibroblast dynamics as an in vitro screening platform for anti-fibrotic drugs in primary myelofibrosis. <i>Journal of Cellular Physiology</i> , 2018, 233, 422-433.	2.0	9
32	Polymorphisms of FDPS, LRP5, SOST and VKORC1 genes and their relation with osteoporosis in postmenopausal Romanian women. <i>PLoS ONE</i> , 2019, 14, e0225776.	1.1	9
33	Exogenous Jaagsiekte Sheep Retrovirus type 2 (exJSRV2) related to ovine pulmonary adenocarcinoma (OPA) in Romania: prevalence, anatomical forms, pathological description, immunophenotyping and virus identification. <i>BMC Veterinary Research</i> , 2020, 16, 296.	0.7	9
34	Presence of copy number aberration and clinical prognostic factors in patients with acute myeloid leukemia: an analysis of effect modification. <i>Polish Archives of Internal Medicine</i> , 2019, 129, 898-906.	0.3	8
35	TheMTHFD1c.1958 G>A polymorphism and recurrent spontaneous abortions. <i>Journal of Maternal-Fetal and Neonatal Medicine</i> , 2011, 24, 189-192.	0.7	7
36	An Exploratory Association Analysis of ABCB1 rs1045642 and ABCB1 rs4148738 with Non-Major Bleeding Risk in Atrial Fibrillation Patients Treated with Dabigatran or Apixaban. <i>Journal of Personalized Medicine</i> , 2020, 10, 133.	1.1	7

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37	Modelling the Effects of MCM7 Variants, Somatic Mutations, and Clinical Features on Acute Myeloid Leukemia Susceptibility and Prognosis. <i>Journal of Clinical Medicine</i> , 2020, 9, 158.	1.0	7
38	VKORC1-1639 G>A Polymorphism and the Risk of Non-Variceal Upper Gastrointestinal Bleeding. <i>Journal of Gastrointestinal and Liver Diseases</i> , 2020, 26, 13-18.	0.5	7
39	TLR1 polymorphisms in Europeans and spontaneous pregnancy loss. <i>Gene</i> , 2012, 494, 109-111.	1.0	6
40	The Importance of Homozygous Polymorphisms of Methylenetetrahydrofolate Reductase Gene in Romanian Patients with Idiopathic Venous Thromboembolism. <i>Balkan Medical Journal</i> , 2013, 30, 197-203.	0.3	6
41	CYP4F2 and VKORC1 Polymorphisms Amplify the Risk of Carotid Plaque Formation. <i>Genes</i> , 2020, 11, 822.	1.0	6
42	TET2 rs1548483 SNP Associating with Susceptibility to Molecularly Annotated Polycythemia Vera and Primary Myelofibrosis. <i>Journal of Personalized Medicine</i> , 2020, 10, 259.	1.1	6
43	Association Analysis of TP53 rs1042522, MDM2 rs2279744, rs3730485, MDM4 rs4245739 Variants and Acute Myeloid Leukemia Susceptibility, Risk Stratification Scores, and Clinical Features: An Exploratory Study. <i>Journal of Clinical Medicine</i> , 2020, 9, 1672.	1.0	6
44	Analysis of the <i><scp>MTHFR</scp></i> (methylenetetrahydrofolate reductase) 677 C>T and 1298 A>C polymorphisms in <i><scp>BCR</scp>–<scp>ABL</scp></i> negative myeloproliferative neoplasms. <i>International Journal of Laboratory Hematology</i> , 2013, 35, e9-12.	0.7	5
45	Familial Essential Thrombocythemia Associated with <i>MPL</i>W515L Mutation in Father and <i>JAK2</i>V617F Mutation in Daughter. <i>Case Reports in Hematology</i> , 2014, 2014, 1-3.	0.3	5
46	The methylenetetrahydrofolate reductase (MTHFR) 677 C>T polymorphism increases the risk of developing chronic myeloid leukemia–a case-control study. <i>Tumor Biology</i> , 2015, 36, 3101-3107.	0.8	5
47	Correlation between ABCB1 gene polymorphisms, antiepileptic drug concentrations and treatment response. <i>Romanian Journal of Laboratory Medicine</i> , 2018, 26, 479-487.	0.1	5
48	Database-Guided Analysis for Immunophenotypic Diagnosis and Follow-Up of Acute Myeloid Leukemia With Recurrent Genetic Abnormalities. <i>Frontiers in Oncology</i> , 2021, 11, 746951.	1.3	5
49	VKORC1 -1639 G>A Polymorphism in Romanian Patients With Deep Vein Thrombosis. <i>Clinical and Applied Thrombosis/Hemostasis</i> , 2016, 22, 760-764.	0.7	4
50	No association between the <i><scp>STAT</scp>5b</i> rs6503691 (C>T) <scp>SNP</scp> and myeloproliferative neoplasms. <i>European Journal of Haematology</i> , 2013, 90, 257-258.	1.1	3
51	Thrombophilia genetic testing in Romanian young women with acute thrombotic events: role of Factor V Leiden, Prothrombin G20210A, MTHFR C677T and A1298C polymorphisms / Evaluarea genetică a trombofililor la femei tinere din România cu evenimente acute trombotice: rolul Factorului V Leiden, Protrombinei G20210A, polimorfismelor MTHFR C677T și A1298C. <i>Romanian Journal of Laboratory Medicine</i> , 2016, 24, 291-305.	0.1	3
52	SERS-Based Assessment of MRD in Acute Promyelocytic Leukemia?. <i>Frontiers in Oncology</i> , 2020, 10, 1024.	1.3	3
53	HFE gene C282Y, H63D and S65C mutations frequency in the Transylvania region, Romania. <i>Journal of Gastrointestinal and Liver Diseases</i> , 2012, 21, 177-80.	0.5	3
54	Association of TLR4 Rs4986791 Polymorphism and TLR9 Haplotypes with Acute Myeloid Leukemia Susceptibility: A Case-Control Study of Adult Patients. <i>Journal of Personalized Medicine</i> , 2022, 12, 409.	1.1	3

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55	JAK2 46/1 haplotype seems not to be associated with lower limb deep venous thrombosis. <i>Blood Cells, Molecules, and Diseases</i> , 2010, 45, 199-200.	0.6	2
56	Concomitant Myeloproliferative and Lymphoid Neoplasms in Two Patients Positive for JAK2 V617F Mutation. Case Report and Literature Review. <i>Indian Journal of Hematology and Blood Transfusion</i> , 2014, 30, 120-123.	0.3	2
57	Modest contribution of JAK2 V617F allele burden to the occurrence of major thrombosis in polycythemia vera and essential thrombocythemia. <i>Blood Cells, Molecules, and Diseases</i> , 2018, 73, 45-46.	0.6	2
58	Presymptomatic diagnosis of CYP24A1-related infantile idiopathic hypercalcemia: A case report. <i>European Journal of Medical Genetics</i> , 2020, 63, 104100.	0.7	2
59	Methylenetetrahydrofolate reductase 677 C> T polymorphism is associated with acute myeloid leukemia. <i>Leukemia and Lymphoma</i> , 2015, 56, 1172-1174.	0.6	1
60	The glucocorticoid receptor A3669G SNP is not associated with polycythemia vera, essential thrombocythemia or primary myelofibrosis. <i>Leukemia and Lymphoma</i> , 2016, 57, 209-211.	0.6	1
61	Haematology patients infected with SARS-CoV-2, pretreated with eculizumab or siltuximab, develop oligosymptomatic disease. <i>European Journal of Hospital Pharmacy</i> , 2022, 29, e8-e8.	0.5	1
62	<i>SH2B3 (LNK)</i> rs3184504 polymorphism is correlated with <i>JAK2</i> V617F-positive myeloproliferative neoplasms. <i>Romanian Journal of Laboratory Medicine</i> , 2020, 28, 267-277.	0.1	1
63	Genotype-phenotype correlations in patients treated with acenocoumarol / CorelaÅ£ii genotip-fenotip la pacienÅ£ii trataÅ£i cu acenocoumarol. <i>Romanian Journal of Laboratory Medicine</i> , 2014, 22, .	0.1	1
64	the Importance of CD36 Expression in Evaluation of Thrombotic Risk in Patients with Myeloproliferative Neoplasms. <i>Blood</i> , 2018, 132, 4989-4989.	0.6	1
65	FGB -455 G>A and GP IIIa PIA1/A2 polymorphisms in a group of Romanian stroke patients. <i>Romanian Journal of Laboratory Medicine</i> , 2016, 24, 45-54.	0.1	0
66	Systemic and Local Factorsâ€™ Influence on the Topological Differences in Deep Vein Thrombosis. <i>Medicina (Lithuania)</i> , 2019, 55, 691.	0.8	0
67	TERT rs2853669 as predictor for overall survival in patients with acute myeloid leukemia. <i>Archives of Medical Science</i> , 2021, 18, 103-111.	0.4	0