Tiscia Giovanni, L

List of Publications by Year
in descending order

56 papers	1,103 citations	430442 18 h-index	433756 31 g-index
57 all docs	57 docs citations	57 times ranked	1378 citing authors

#	Article	IF	Citations
1	The JAK2 V617F mutation frequently occurs in patients with portal and mesenteric venous thrombosis. Journal of Thrombosis and Haemostasis, 2007, 5, 55-61.	1.9	150
2	Pharmacogenetics of dabigatran etexilate interindividual variability. Thrombosis Research, 2016, 144, 1-5.	0.8	69
3	Haplotype M2 in the annexin A5 (ANXA5) gene and the occurrence of obstetric complications. Thrombosis and Haemostasis, 2009, 102, 309-313.	1.8	63
4	Maternal thrombophilia and the risk of recurrence of preeclampsia. American Journal of Obstetrics and Gynecology, 2009, 200, 46.e1-46.e5.	0.7	62
5	Human Fibrinogen: Molecular and Genetic Aspects of Congenital Disorders. International Journal of Molecular Sciences, 2018, 19, 1597.	1.8	51
6	A new JAK2 gene mutation in patients with polycythemia vera and splanchnic vein thrombosis. Blood, 2007, 110, 2768-2768.	0.6	49
7	Annexin V expression in human placenta is influenced by the carriership of the common haplotype M2. Fertility and Sterility, 2009, 91, 940-942.	0.5	45
8	The COX-2 G/C â˜'765 polymorphism may modulate the occurrence of cerebrovascular ischemia. Blood Coagulation and Fibrinolysis, 2006, 17, 93-96.	0.5	44
9	ABCB1 SNP rs4148738 modulation of apixaban interindividual variability. Thrombosis Research, 2016, 145, 24-26.	0.8	41
10	Outcome of patients with splanchnic venous thrombosis presenting without overt MPN: A role for the JAK2 V617F mutation re-evaluation. Thrombosis Research, 2013, 132, e99-e104.	0.8	34
11	Liquid chromatographyâ€"tandem mass spectrometry method as the golden standard for therapeutic drug monitoring in renal transplant. Journal of Pharmaceutical and Biomedical Analysis, 2013, 86, 123-126.	1.4	34
12	De novo homozygous mutation of the C1 inhibitor gene in a patient with hereditary angioedema. Journal of Allergy and Clinical Immunology, 2013, 132, 748-750.e3.	1.5	28
13	Obstetric complications and pregnancy-related venous thromboembolism: The effect of low-molecular-weight heparin on their prevention in carriers of factor V Leiden or prothrombin G20210A mutation. Thrombosis and Haemostasis, 2012, 107, 477-484.	1.8	25
14	Pregnancy-related venous thrombosis: comparison between spontaneous and ART conception in an Italian cohort. BMJ Open, 2015, 5, e008213.	0.8	24
15	Role of the M2 haplotype within the annexin A5 gene in the occurrence of pregnancy-related venous thromboembolism. American Journal of Obstetrics and Gynecology, 2010, 203, 461.e1-461.e5.	0.7	23
16	Occurrence of the JAK2 V617F mutation in the Buddâ€"Chiari syndrome. Blood Coagulation and Fibrinolysis, 2008, 19, 459-462.	0.5	20
17	The JAK2 rs12343867 CC genotype frequently occurs in patients with splanchnic venous thrombosis without the JAK2V617F mutation: a retrospective study. Journal of Thrombosis and Haemostasis, 2010, 8, 413-416.	1.9	19
18	Risk of obstetric and thromboembolic complications in family members of women with previous adverse obstetric outcomes carrying common inherited thombophilias. Journal of Thrombosis and Haemostasis, 2012, 10, 223-228.	1.9	19

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19	Validation of PLASMIC score and follow-up data in a cohort of patients with suspected microangiopathies from Southern Italy. Journal of Thrombosis and Thrombolysis, 2018, 46, 174-179.	1.0	19
20	Sex modulation of the occurrence of jak2 v617f mutation in patients with splanchnic venous thrombosis. Thrombosis Research, 2011, 128, 233-236.	0.8	18
21	A novel congenital dysprothrombinemia leading to defective prothrombin maturation. Thrombosis Research, 2014, 134, 1135-1141.	0.8	18
22	Findings from a multicentre, observational study on reproductive outcomes in women with unexplained recurrent pregnancy loss: the OTTILIA registry. Human Reproduction, 2021, 36, 2083-2090.	0.4	17
23	The M2 haplotype in the ANXA5 gene is an independent risk factor for idiopathic small-for-gestational age newborns. Molecular Human Reproduction, 2012, 18, 510-513.	1.3	16
24	New TET2 gene mutations in patients with myeloproliferative neoplasms and splanchnic vein thrombosis. Journal of Thrombosis and Haemostasis, 2010, 8, 1142-1144.	1.9	14
25	Management of Patients Taking Oral Anticoagulants Who Need Urgent Surgery for Hip Fracture. Seminars in Thrombosis and Hemostasis, 2019, 45, 164-170.	1.5	14
26	Impact of common thrombophilias and JAK2 V617F on pregnancy outcomes in unselected Italian women. Journal of Thrombosis and Haemostasis, 2011, 9, 496-501.	1.9	13
27	Clinical utility of screening for CALR gene exon 9 mutations in patients with splanchnic venous thrombosis. Thrombosis and Haemostasis, 2015, 113, 1381-1382.	1.8	13
28	Venous thrombosis in oral contraceptive users and the presence of the JAK2 V617F mutation. Thrombosis and Haemostasis, 2008, 99, 640-642.	1.8	12
29	Clinical histories and molecular characterization of two afibrinogenemic patients: insights into clinical management. Haemophilia, 2012, 18, e16-8.	1.0	12
30	Markers of haemostasis and angiogenesis in placentae from gestational vascular complications: Impairment of mechanisms involved in maintaining intervillous blood flow. Thrombosis Research, 2010, 125, 267-271.	0.8	11
31	Low-molecular â€“weight heparin in pregnancies after ART -A retrospective study-. Thrombosis Research, 2014, 134, 336-339.	0.8	11
32	Aspirin and heparin in pregnancy. Expert Opinion on Pharmacotherapy, 2015, 16, 1793-1803.	0.9	11
33	TET2 Mutations in Ph-Negative Myeloproliferative Neoplasms: Identification of Three Novel Mutations and Relationship with Clinical and Laboratory Findings. BioMed Research International, 2013, 2013, 1-5.	0.9	10
34	Genetic variations in the annexin A5 gene and the risk ofÂpregnancyâ€related venous thrombosis. Journal of Thrombosis and Haemostasis, 2015, 13, 409-413.	1.9	10
35	Clinical Pregnancies and Live Births in women approaching ART: A follow-up analysis of 157 women after thrombophilia screening. Thrombosis Research, 2014, 133, 168-172.	0.8	9
36	The haplotype M2 within the ANXA5 gene is independently associated with the occurrence of deep venous thrombosis. Thrombosis and Haemostasis, 2010, 103, 1102-1103.	1.8	8

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37	Influence of the Gly1057Asp variant of the insulin receptor substrate 2 (IRS2) on insulin resistance and relationship with epicardial fat thickness in the elderly. Experimental Gerontology, 2012, 47, 988-993.	1.2	8
38	Functional characterization of annexin A5 gene promoter allelic variants. Thrombosis Research, 2016, 144, 93-99.	0.8	7
39	More on: factor V Leiden and prothrombin G20210A polymorphisms as risk factors for miscarriage during a first-intended pregnancy: the matched case-control 'NOHA first' study. Journal of Thrombosis and Haemostasis, 2006, 4, 709-710.	1.9	6
40	Clinical Utility of Antithrombotic Prophylaxis in ART Procedures: An Italian Experience. PLoS ONE, 2014, 9, e97604.	1.1	6
41	Factor XI gene variants in factor XI-deficient patients of Southern Italy: identification of a novel mutation and genotypeâ€“phenotype relationship. Human Genome Variation, 2017, 4, 17043.	0.4	6
42	Factor VII deficiency: a novel missense variant and genotypeâ€“phenotype correlation in patients from Southern Italy. Human Genome Variation, 2017, 4, 17048.	0.4	6
43	The Prognostic Value of ADAMTS-13 and von Willebrand Factor in COVID-19 Patients: Prospective Evaluation by Care Setting. Diagnostics, 2021, 11, 1648.	1.3	6
44	Obstetric outcomes in pregnant COVID-19 women: the imbalance of von Willebrand factor and ADAMTS13 axis. BMC Pregnancy and Childbirth, 2022, 22, 142.	0.9	6
45	Factor XI deficiency: two novel mutations in asymptomatic Italian patients. Haemophilia, 2010, 16, 767-770.	1.0	4
46	Structural analysis of protein Z gene variants in patients with foetal losses. Thrombosis and Haemostasis, 2013, 110, 534-542.	1.8	3
47	Thromboelastography Parameters in Italian Pregnant Women: Do Antithrombotic Drugs Change Reference Values?. Journal of Investigative Medicine, 2020, 68, 902-905.	0.7	3
48	Modulation of factors involved in placental haemostasis and angiogenesis by low-molecular-weight-heparins. Archives of Gynecology and Obstetrics, 2016, 294, 1323-1329.	0.8	2
49	Postpartum haemorrhage in a woman with essential thrombocythemia carrying calreticulin mutation. Blood Coagulation and Fibrinolysis, 2016, 27, 727-728.	0.5	1
50	<scp>PLASMIC</scp> score: Not intended to replace but rather to prompt the <scp>ADAMTS13</scp> testing. Transfusion, 2020, 60, 3070-3072.	0.8	1
51	C0267 Genotype-phenotype relationship in Italian subjects with congenital FXI deficiency. Thrombosis Research, 2012, 130, S140.	0.8	0
52	C0264 Protein Z mutations in patients with fetal losses: Structural analysis. Thrombosis Research, 2012, 130, S146.	0.8	0
53	C0268 Higher risk of idiopathic small for gestational age newborns in Italian women carrying the annexin A5 M2 haplotype. Thrombosis Research, 2012, 130, S195.	0.8	0
54	Antiphospholipid Antibodies in a General Obstetric Population: Clinical Impact on Pregnancy Outcome and Relationship with the M2 Haplotype in the Annexin A5 (ANXA5) Gene. Hamostaseologie, 2019, 39, 203-207.	0.9	0

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55	Recurrent thrombocytopenia in pregnancy: is it always an obstetric complication?. Internal and Emergency Medicine, 2020, 15, 1271-1274.	1.0	0
56	Focus on Key Issues in Immune Thrombotic Thrombocytopenic Purpura: Italian Experience of Six Centers. Journal of Clinical Medicine, 2021, 10, 5702.	1.0	0