## Donato Gemmati

List of Publications by Year in descending order

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84 papers 3,186 citations

32 h-index 53 g-index

84 all docs

84 docs citations

84 times ranked 4283 citing authors

#	Article	IF	CITATIONS
1	COVID-19 and Individual Genetic Susceptibility/Receptivity: Role of ACE1/ACE2 Genes, Immunity, Inflammation and Coagulation. Might the Double X-Chromosome in Females Be Protective against SARS-CoV-2 Compared to the Single X-Chromosome in Males?. International Journal of Molecular Sciences, 2020, 21, 3474.	1.8	290
2	Risk of venous thromboembolism associated with single and combined effects of Factor V Leiden, Prothrombin 20210A and Methylenetethraydrofolate reductase C677T: a meta-analysis involving over 11,000 cases and 21,000 controls. European Journal of Epidemiology, 2013, 28, 621-647.	2.5	141
3	Value of Platelet Reactivity in Predicting Response to Treatment and Clinical Outcome in Patients Undergoing Primary Coronary Intervention. Journal of the American College of Cardiology, 2006, 48, 2178-2185.	1.2	140
4	Effects of Electromagnetic Fields on Proteoglycan Metabolism of Bovine Articular Cartilage Explants. Connective Tissue Research, 2003, 44, 154-159.	1.1	116
5	A common mutation in the gene for coagulation factor XIII-A (VAL34Leu): A risk factor for primary intracerebral hemorrhage is protective against atherothrombotic diseases. American Journal of Hematology, 2001, 67, 183-188.	2.0	97
6	Gene polymorphisms in folate metabolizing enzymes in adult acute lymphoblastic leukemia: effects on methotrexate-related toxicity and survival. Haematologica, 2009, 94, 1391-1398.	1.7	96
7	Genetic Association and Altered Gene Expression of Mir-155 in Multiple Sclerosis Patients. International Journal of Molecular Sciences, 2011, 12, 8695-8712.	1.8	93
8	Hemochromatosis C282Y gene mutation increases the risk of venous leg ulceration. Journal of Vascular Surgery, 2005, 42, 309-314.	0.6	89
9	Poor Responsiveness to Clopidogrel: Drug-Specific or Class-Effect Mechanism?. Journal of the American College of Cardiology, 2007, 50, 1132-1137.	1.2	82
10	Redox metals homeostasis in multiple sclerosis and amyotrophic lateral sclerosis: a review. Cell Death and Disease, 2018, 9, 348.	2.7	82
11	Proteoglycan synthesis in bovine articular cartilage explants exposed to different low-frequency low-energy pulsed electromagnetic fields. Osteoarthritis and Cartilage, 2007, 15, 163-168.	0.6	78
12	Effects of physical stimulation with electromagnetic field and insulin growth factor-I treatment on proteoglycan synthesis of bovine articular cartilage. Osteoarthritis and Cartilage, 2004, 12, 793-800.	0.6	75
13	Low Folate Levels and Thermolabile Methylenetetrahydrofolate Reductase as Primary Determinant of Mild Hyperhomocystinemia in Normal and Thromboembolic Subjects. Arteriosclerosis, Thrombosis, and Vascular Biology, 1999, 19, 1761-1767.	1.1	70
14	Detection of two missense mutations and characterization of a repeat polymorphism in the factor VII gene (F7). Human Genetics, 1992, 89, 497-502.	1.8	66
15	"Bridging the Gap―Everything that Could Have Been Avoided If We Had Applied Gender Medicine, Pharmacogenetics and Personalized Medicine in the Gender-Omics and Sex-Omics Era. International Journal of Molecular Sciences, 2020, 21, 296.	1.8	63
16	Common gene polymorphisms in the metabolic folate and methylation pathway and the risk of acute lymphoblastic leukemia and non-Hodgkin's lymphoma in adults. Cancer Epidemiology Biomarkers and Prevention, 2004, $13$ , $787-94$ .	1.1	56
17	Tissue Factor and Coagulation Factor VII Levels During Acute Myocardial Infarction. Arteriosclerosis, Thrombosis, and Vascular Biology, 2006, 26, 2800-2806.	1.1	53
18	Methylenetetrahydrofolate reductase C677T and A1298C gene variants in adult non-Hodgkin's lymphoma patients: association with toxicity and survival. Haematologica, 2007, 92, 478-485.	1.7	53

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19	Genetic Hypothesis and Pharmacogenetics Side of Renin-Angiotensin-System in COVID-19. Genes, 2020, 11, 1044.	1.0	52
20	DNA-array of gene variants in venous leg ulcers: Detection of prognostic indicators. Journal of Vascular Surgery, 2009, 50, 1444-1451.	0.6	49
21	Review: Interplay of Iron Metallobiology, Metalloproteinases, and FXIII, and Role of Their Gene Variants in Venous Leg Ulcer. International Journal of Lower Extremity Wounds, 2010, 9, 166-179.	0.6	48
22	Sustainable Agriculture through Multidisciplinary Seed Nanopriming: Prospects of Opportunities and Challenges. Cells, 2021, 10, 2428.	1.8	48
23	Prognostic role of factor XIII gene variants in nonhealing venous leg ulcers. Journal of Vascular Surgery, 2006, 44, 815-819.	0.6	45
24	Investigation of in vitro cytotoxicity of the redox state of ionic iron in neuroblastoma cells. Journal of Neurosciences in Rural Practice, 2012, 03, 301-310.	0.3	45
25	Effects of electromagnetic fields on proteoglycan metabolism of bovine articular cartilage explants. Connective Tissue Research, 2003, 44, 154-9.	1.1	44
26	Factor XIII V34L polymorphism modulates the risk of chronic venous leg ulcer progression and extension. Wound Repair and Regeneration, 2004, 12, 512-517.	1.5	43
27	Influence of gene polymorphisms in ulcer healing process after superficial venous surgery. Journal of Vascular Surgery, 2006, 44, 554-562.	0.6	43
28	Serum Iron and Matrix Metalloproteinase-9 Variations in Limbs Affected by Chronic Venous Disease and Venous Leg Ulcers. Dermatologic Surgery, 2005, 31, 644-649.	0.4	42
29	Polymorphisms in the genes coding for iron binding and transporting proteins are associated with disability, severity, and early progression in multiple sclerosis. BMC Medical Genetics, 2012, 13, 70.	2.1	42
30	Gene-gene interactions among coding genes of iron-homeostasis proteins and APOE-alleles in cognitive impairment diseases. PLoS ONE, 2018, 13, e0193867.	1.1	40
31	Predictive role of coagulation-balance gene polymorphisms in the efficacy of photodynamic therapy with verteporfin for classic choroidal neovascularization secondary to age-related macular degeneration. Pharmacogenetics and Genomics, 2007, 17, 1039-1046.	0.7	38
32	Coagulation Gene Predictors of Photodynamic Therapy for Occult Choroidal Neovascularization in Age-Related Macular Degeneration. , 2008, 49, 3100.		38
33	Factor XIII Contrasts the Effects of Metalloproteinases in Human Dermal Fibroblast Cultured Cells. Vascular and Endovascular Surgery, 2004, 38, 431-438.	0.3	37
34	Clinical implications of gene polymorphisms in venous leg ulcer: A model in tissue injury and reparative process. Thrombosis and Haemostasis, 2007, 98, 131-137.	1.8	34
35	Coagulation Factor XIIIA (F13A1): Novel Perspectives in Treatment and Pharmacogenetics. Current Pharmaceutical Design, 2016, 22, 1449-1459.	0.9	34
36	Factor XIIIA-V34L and Factor XIIIB-H95R Gene Variants: Effects on Survival in Myocardial Infarction Patients. Molecular Medicine, 2007, 13, 112-120.	1.9	32

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37	Functional variations modulating PRKCA expression and alternative splicing predispose to multiple sclerosis. Human Molecular Genetics, 2014, 23, 6746-6761.	1.4	32
38	Serum Iron and Matrix Metalloproteinase-9 Variations in Limbs Affected by Chronic Venous Disease and Venous Leg Ulcers. Dermatologic Surgery, 2006, 31, 644-649.	0.4	31
39	TRAIL, OPG, and TWEAK in kidney disease: biomarkers or therapeutic targets?. Clinical Science, 2019, 133, 1145-1166.	1.8	30
40	Nanobiomaterials for vascular biology and wound management: A review. Veins and Lymphatics, 2018, $7$ , .	0.1	27
41	Traditional Herbal Remedies with a Multifunctional Therapeutic Approach as an Implication in COVID-19 Associated Co-Infections. Coatings, 2020, 10, 761.	1.2	27
42	Predictive role of C677T <i>MTHFR</i> polymorphism in variable efficacy of photodynamic therapy for neovascular age-related macular degeneration. Pharmacogenomics, 2009, 10, 81-95.	0.6	25
43	Sex/Gender-Specific Imbalance in CVD: Could Physical Activity Help to Improve Clinical Outcome Targeting CVD Molecular Mechanisms in Women?. International Journal of Molecular Sciences, 2020, 21, 1477.	1.8	24
44	Factor XIII-A dynamics in acute myocardial infarction: a novel prognostic biomarker?. Thrombosis and Haemostasis, 2015, 114, 123-132.	1.8	23
45	Sudden Sensorineural Hearing Loss and Polymorphisms in Iron Homeostasis Genes: New Insights from a Case-Control Study. BioMed Research International, 2015, 2015, 1-10.	0.9	23
46	Recent Advances in Plant Nanobionics and Nanobiosensors for Toxicology Applications. Current Nanoscience, 2020, 16, 27-41.	0.7	23
47	Meta-Analysis of Multiple Sclerosis Microarray Data Reveals Dysregulation in RNA Splicing Regulatory Genes. International Journal of Molecular Sciences, 2015, 16, 23463-23481.	1.8	22
48	Genetics and Epigenetics of One-Carbon Metabolism Pathway in Autism Spectrum Disorder: A Sex-Specific Brain Epigenome?. Genes, 2021, 12, 782.	1.0	22
49	DHFR 19â€bp insertion/deletion polymorphism and MTHFR C677T in adult acute lymphoblastic leukaemia: Is the risk reduction due to intracellular folate unbalancing?. American Journal of Hematology, 2009, 84, 526-529.	2.0	21
50	Custom CGH array profiling of copy number variations (CNVs) on chromosome 6p21.32 (HLA locus) in patients with venous malformations associated with multiple sclerosis. BMC Medical Genetics, 2010, 11, 64.	2.1	21
51	F13A1 Gene Variant (V34L) and Residual Circulating FXIIIA Levels Predict Short- and Long-Term Mortality in Acute Myocardial Infarction after Coronary Angioplasty. International Journal of Molecular Sciences, 2018, 19, 2766.	1.8	21
52	Crosstalk Between Adipokines and Paraoxonase 1: A New Potential Axis Linking Oxidative Stress and Inflammation. Antioxidants, 2019, 8, 287.	2,2	19
53	Time- and Dose-Dependent Effects of Chronic Wound Fluid on Human Adult Dermal Fibroblasts. Dermatologic Surgery, 2008, 34, 347-356.	0.4	18
54	p53/NF-kB Balance in SARS-CoV-2 Infection: From OMICs, Genomics and Pharmacogenomics Insights to Tailored Therapeutic Perspectives (COVIDomics). Frontiers in Pharmacology, 0, 13, .	1.6	18

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55	A Modified Functional Global Test to Measure Protein C, Protein S Activities and the Activated Protein C-Resistance Phenotype. Thrombosis Research, 1998, 92, 141-148.	0.8	15
56	Clinical implications of gene polymorphisms in venous leg ulcer: a model in tissue injury and reparative process. Thrombosis and Haemostasis, 2007, 98, 131-7.	1.8	15
57	Prevalence of factor XIII Val34Leu polymorphism in patients affected by spontaneous subconjunctival hemorrhage. American Journal of Ophthalmology, 2004, 138, 481-484.	1.7	14
58	Recurrent episodes of spontaneous subconjunctival hemorrhage in patients with factor XIII Val34Leu mutation. American Journal of Ophthalmology, 2002, 134, 927-929.	1.7	13
59	Inherited genetic predispositions in F13A1 and F13B genes predict abdominal adhesion formation: identification of gender prognostic indicators. Scientific Reports, 2018, 8, 16916.	1.6	13
60	C6orf10 Low-Frequency and Rare Variants in Italian Multiple Sclerosis Patients. Frontiers in Genetics, 2019, 10, 573.	1.1	13
61	Time- and Dose-Dependent Effects of Chronic Wound Fluid on Human Adult Dermal Fibroblasts. Dermatologic Surgery, 2008, 34, 347-356.	0.4	12
62	Impact of Coagulation-Balance Gene Predictors on Efficacy of Photodynamic Therapy for Choroidal Neovascularization in Pathologic Myopia. Ophthalmology, 2010, 117, 517-523.	2.5	12
63	The Active Metabolite of Warfarin (3'-Hydroxywarfarin) and Correlation with INR, Warfarin and Drug Weekly Dosage in Patients under Oral Anticoagulant Therapy: A Pharmacogenetics Study. PLoS ONE, 2016, 11, e0162084.	1.1	12
64	Pharmacogenetic Aspects in Therapeutic Management of Subfoveal Choroidal Neovascularisation: Role of Factor XIII-A 185 T-Allele. Current Drug Targets, 2011, 12, 138-148.	1.0	12
65	Predictive Role of Gene Polymorphisms Affecting Thrombin-Generation Pathway in Variable Efficacy of Photodynamic Therapy for Neovascular Age-Related Macular Degeneration. Recent Patents on DNA & Gene Sequences, 2009, 3, 114-122.	0.7	10
66	Maternal Haplotypes in DHFR Promoter and MTHFR Gene in Tuning Childhood Acute Lymphoblastic Leukemia Onset-Latency: Genetic/Epigenetic Mother/Child Dyad Study (GEMCDS). Genes, 2019, 10, 634.	1.0	10
67	Profiling the mutational landscape of coagulation factor V deficiency. Haematologica, 2020, 105, e180-e185.	1.7	10
68	Nanoengineering Approaches to Design Advanced Dental Materials for Clinical Applications. Journal of Bionanoscience, 2010, 4, 53-65.	0.4	9
69	Genetic Predictors of Response to Photodynamic Therapy. Molecular Diagnosis and Therapy, 2011, 15, 195-210.	1.6	9
70	Cis-Segregation of c.1171C>T Stop Codon (p.R391*) in SERPINC1 Gene and c.1691G>A Transition (p.R506Q) in F5 Gene and Selected GWAS Multilocus Approach in Inherited Thrombophilia. Genes, 2021, 12, 934.	1.0	9
71	F9 missense mutations impairing factor IX activation are associated with pleiotropic plasma phenotypes. Journal of Thrombosis and Haemostasis, 2022, 20, 69-81.	1.9	9
72	A photometric method for the dosage of factor XIII applied to the study of chronic hepatopathies. Thrombosis Research, 1995, 78, 451-456.	0.8	8

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73	Genetic variants associated with methotrexate-induced mucositis in cancer treatment: A systematic review and meta-analysis. Critical Reviews in Oncology/Hematology, 2021, 161, 103312.	2.0	8
74	Can estrogens protect against COVID-19? The COVID-19 puzzling and gender medicine. Minerva Ginecologica, 2020, 72, 178-179.	0.8	8
75	Vision Loss after PDT. Ophthalmology, 2006, 113, 157.e1-157.e4.	2.5	7
76	Changes in Adipose Tissue Distribution and Association between Uric Acid and Bone Health during Menopause Transition. International Journal of Molecular Sciences, 2019, 20, 6321.	1.8	7
77	Effect of Factor XIII-A G185T Polymorphism on Visual Prognosis after Photodynamic Therapy for Neovascular Macular Degeneration. International Journal of Molecular Sciences, 2015, 16, 19796-19811.	1.8	5
78	Impact of methylenetetrahydrofolate reductase C677T polymorphism on the efficacy of photodynamic therapy in patients with neovascular age-related macular degeneration. Scientific Reports, 2019, 9, 2614.	1.6	5
79	Assessment of the interlaboratory variability and robustness of <i>JAK2</i> V617F mutation assays: A study involving a consortium of 19 Italian laboratories. Oncotarget, 2017, 8, 32608-32617.	0.8	5
80	Asymptomatic carriership of factor V Leiden and genotypes of the fibrinogen gene cluster. British Journal of Haematology, 2003, 121, 632-638.	1.2	4
81	Reference materials (RMs) for analysis of the human factor II (prothrombin) gene G20210A mutation. Clinical Chemistry and Laboratory Medicine, 2005, 43, 862-8.	1.4	3
82	Temporal and Genotype-Driven Variation of Factor VII Levels in Patients With Acute Myocardial Infarction. Clinical and Applied Thrombosis/Hemostasis, 2009, 15, 119-122.	0.7	3
83	Multiple Strokes in a Newborn. Ophthalmology, 2009, 116, 812-813.e3.	2.5	3
84	Thrombophilia in the Occurrence of Retinal Vascular Infarction After Photodynamic Therapy With Verteporfin Using the Standard Protocol. JAMA Ophthalmology, 2010, 128, 1632.	2.6	2