## Giuliana Merati

List of Publications by Year in descending order

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**CILILIANA MEDATI** 

#	Article	IF	CITATIONS
1	Interaction between photosystem I and ferredoxin. Identification by chemical cross-linking of the polypeptide which binds ferredoxin. FEBS Journal, 1987, 169, 143-146.	0.2	174
2	Hypercoagulability in patients with type 2 diabetes mellitus detected by a thrombin generation assay. Journal of Thrombosis and Thrombolysis, 2011, 31, 165-172.	2.1	129
3	Gene Polymorphisms Predicting High Plasma Levels of Coagulation and Fibrinolysis Proteins. Arteriosclerosis, Thrombosis, and Vascular Biology, 1997, 17, 755-759.	2.4	103
4	Circulating microparticles and risk of venous thromboembolism. Thrombosis Research, 2012, 129, 591-597.	1.7	92
5	A 23bp Insertion in the Endothelial Protein C Receptor (EPCR) Gene Impairs EPCR Function. Thrombosis and Haemostasis, 2001, 86, 945-948.	3.4	56
6	Role of Chloride Ions in Modulation of the Interaction between von Willebrand Factor and ADAMTS-13. Journal of Biological Chemistry, 2005, 280, 23295-23302.	3.4	43
7	Mechanisms of the interaction between twoADAMTS13 gene mutations leading to severe deficiency of enzymatic activity. Human Mutation, 2006, 27, 330-336.	2.5	39
8	Chemical cross-linking of ferredoxin to spinach thylakoids. FEBS Letters, 1987, 215, 37-40.	2.8	29
9	Residual platelets are the main determinants of microparticles count in frozen-thawed plasma. Thrombosis Research, 2012, 130, 561-562.	1.7	19
10	Identification and computationally-based structural interpretation of naturally occurring variants of human protein C. Human Mutation, 2007, 28, 345-355.	2.5	17
11	Antioxidant Activity of Ubiquinone-3 in Human Low Density Lipoprotein. Free Radical Research Communications, 1992, 16, 11-17.	1.8	12
12	The G1456 to T Mutation in the Thrombomodulin Gene Is Not Frequent in Patients With Venous Thrombosis. Blood, 1997, 89, 1467-1467.	1.4	12
13	Point Mutations in the Endothelial Protein C Receptor (EPCR) Promoter. Thrombosis and Haemostasis, 2002, 87, 1085-1086.	3.4	4
14	Determination of Lp(a) and APO(a) in an Italian population. Research in Clinic and Laboratory, 1991, 21, 127-134.	0.3	2
15	In Vitro Expression Studies of Two Mutations on the Metalloprotease and First Cub Domains of the ADAMTS-13 Gene Leading to Severe ADAMTS-13 Deficiency and Chronic Recurrent TTP Blood, 2004, 104, 514-514.	1.4	2
16	Point mutations in the endothelial protein C receptor (EPCR) promoter. Thrombosis and Haemostasis, 2002, 87, 1085-6.	3.4	1