Ugo Cavallari

List of Publications by Year
in descending order

21	984	567247 15	794568 19
papers	citations	h-index	g-index

21	21	21	1478
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	SNPs of the FADS Gene Cluster are Associated with Polyunsaturated Fatty Acids in a Cohort of Patients with Cardiovascular Disease. Lipids, 2008, 43, 289-299.	1.7	218
2	Contribution of Gene Sequence Variations of the Hepatic Cytochrome P450 3A4 Enzyme to Variability in Individual Responsiveness to Clopidogrel. Arteriosclerosis, Thrombosis, and Vascular Biology, 2006, 26, 1895-1900.	2.4	214
3	Lack of association between the P2Y12 receptor gene polymorphism and platelet response to clopidogrel in patients with coronary artery disease. Thrombosis Research, 2005, 116, 491-497.	1.7	137
4	Variability in Platelet Aggregation Following Sustained Aspirin and Clopidogrel Treatment in Patients With Coronary Heart Disease and Influence of the 807 C/T Polymorphism of the Glycoprotein Ia Gene. American Journal of Cardiology, 2005, 96, 1095-1099.	1.6	60
5	Gene sequence variations of the platelet P2Y12 receptor are associated with coronary artery disease. BMC Medical Genetics, 2007, 8, 59.	2.1	53
6	On the association of the oxidised LDL receptor 1 (OLR1) gene in patients with acute myocardial infarction or coronary artery disease. European Journal of Human Genetics, 2006, 14, 127-130.	2.8	45
7	ALOX5AP gene variants and risk of coronary artery disease: an angiography-based study. European Journal of Human Genetics, 2007, 15, 959-966.	2.8	37
8	Combined Effect of Hemostatic Gene Polymorphisms and the Risk of Myocardial Infarction in Patients with Advanced Coronary Atherosclerosis. PLoS ONE, 2008, 3, e1523.	2.5	35
9	Four new cases of double heterozygosity for BRCA1 and BRCA2 gene mutations: clinical, pathological, and family characteristics. Breast Cancer Research and Treatment, 2010, 124, 251-258.	2.5	27
10	Lack of association between gene sequence variations of platelet membrane receptors and aspirin responsiveness detected by the PFA-100 system in patients with coronary artery disease. Platelets, 2006, 17, 586-590.	2.3	26
11	Germline mosaicism in achondroplasia detected in sperm DNA of the father of three affected sibs. American Journal of Medical Genetics, Part A, 2008, 146A, 784-786.	1.2	22
12	Three cases with de novo 6q imbalance and variable prenatal phenotype. American Journal of Medical Genetics, Part A, 2005, 136A, 254-258.	1.2	21
13	Early manifestations in a cohort of children prenatally diagnosed with 47,XYY. Role of multidisciplinary counseling for parental guidance and prevention of aggressive behavior. Italian Journal of Pediatrics, 2012, 38, 52.	2.6	20
14	Biparental expression of ESX1L gene in placentas from normal and intrauterine growth-restricted pregnancies. European Journal of Human Genetics, 2004, 12, 272-278.	2.8	18
15	Triple X syndrome: characteristics of 42 Italian girls and parental emotional response to prenatal diagnosis. European Journal of Pediatrics, 2010, 169, 1255-1261.	2.7	18
16	Influence of the CD14 C260T Promoter Polymorphism on C-Reactive Protein Levels in Patients With Coronary Artery Disease. American Journal of Cardiology, 2006, 98, 1182-1184.	1.6	11
17	Reply to Novelli. European Journal of Human Genetics, 2006, 14, 895-895.	2.8	10
18	Caffeine intake and risk of neural tube defects. Birth Defects Research Part A: Clinical and Molecular Teratology, 2011, 91, 67-67.	1.6	6

#	Article	IF	Citations
19	Array CGH in routine prenatal diagnosis practice. Prenatal Diagnosis, 2012, 32, 708-709.	2.3	6
20	Acro-dermato-ungual-lacrimal-tooth-like syndrome: report of a family with variable expression. Clinical Dysmorphology, 2006, 15, 239-241.	0.3	0
21	ALOX5AP Gene Variants and Risk of Coronary Artery Disease in Italy. An Angiography-Based Study.. Blood, 2006, 108, 1459-1459.	1.4	0