Fasano Tommaso

List of Publications by Year
in descending order

17	646	840776 11	1125743 13
papers	citations	h-index	g-index

19	19	19	1007
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Spectrum of mutations and phenotypic expression in patients with autosomal dominant hypercholesterolemia identified in Italy. Atherosclerosis, 2013, 227, 342-348.	0.8	128
2	A Novel Loss of Function Mutation of PCSK9 Gene in White Subjects With Low-Plasma Low-Density Lipoprotein Cholesterol. Arteriosclerosis, Thrombosis, and Vascular Biology, 2007, 27, 677-681.	2.4	125
3	Lysosomal lipase deficiency: Molecular characterization of eleven patients with Wolman or cholesteryl ester storage disease. Molecular Genetics and Metabolism, 2012, 105, 450-456.	1.1	71
4	Effect of ezetimibe coadministered with statins in genotype-confirmed heterozygous FH patients. Atherosclerosis, 2007, 194, e116-e122.	0.8	68
5	Degradation of LDLR protein mediated by â€˜gain of functionâ€™ PCSK9 mutants in normal and ARH cells. Atherosclerosis, 2009, 203, 166-171.	0.8	64
6	Familial HDL deficiency due to ABCA1 gene mutations with or without other genetic lipoprotein disorders. Atherosclerosis, 2004, 172, 309-320.	0.8	47
7	Identification and characterization of novel loss of function mutations in ATP-binding cassette transporter A1 in patients with low plasma high-density lipoprotein cholesterol. Atherosclerosis, 2010, 213, 492-498.	0.8	46
8	Novel mutations of ABCA1 transporter in patients with Tangier disease and familial HDL deficiency. Molecular Genetics and Metabolism, 2012, 107, 534-541.	1.1	28
9	Severe HDL deficiency due to novel defects in the ABCA1 transporter. Journal of Internal Medicine, 2009, 265, 359-372.	6.0	24
10	Denaturing high-performance liquid chromatography in the detection of ABCA1 gene mutations in familial HDL deficiency. Journal of Lipid Research, 2005, 46, 817-822.	4.2	16
11	Multiple abnormally spliced ABCA1 mRNAs caused by a novel splice site mutation of ABCA1 gene in a patient with Tangier disease. Clinica Chimica Acta, 2010, 411, 524-530.	1.1	15
12	A novel mutation of the apolipoprotein A-I gene in a family with familial combined hyperlipidemia. Atherosclerosis, 2008, 198, 145-151.	0.8	9
13	Coding sequence and intronâ€“exon junctions of the c-myb gene are intact in the chronic phase and blast crisis stages of chronic myeloid leukemia patients. Leukemia Research, 2007, 31, 163-167.	0.8	5
14	Mo-W5:3 Molecular diagnosis and treatment of hypobetalipoproteinemia. Atherosclerosis Supplements, 2006, 7, 19-20.	1.2	0
15	Mo-P6:413 Genotype-phenotype relationship in LCAT deficiencies. Atherosclerosis Supplements, 2006, 7, 136-137.	1.2	0
16	PO5-152 FUNCTIONAL CHARACTERIZATION OF ABCA1 GENE MUTANTS. Atherosclerosis Supplements, 2007, 8, 55.	1.2	0
17	Cardiovascular risk profile in a patient with a novel splice site ABCA1 mutation (HDL) Tj ETQq1 1 0.784314 rgBT /Overlock 10 Tf 50 10	0.8	0