

Fasano Tommaso

List of Publications by Year in descending order

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Version: 2024-02-01

17
papers

646
citations

840776

11
h-index

1125743

13
g-index

19
all docs

19
docs citations

19
times ranked

1007
citing authors

#	ARTICLE	IF	CITATIONS
1	Spectrum of mutations and phenotypic expression in patients with autosomal dominant hypercholesterolemia identified in Italy. <i>Atherosclerosis</i> , 2013, 227, 342-348.	0.8	128
2	Novel mutations of ABCA1 transporter in patients with Tangier disease and familial HDL deficiency. <i>Molecular Genetics and Metabolism</i> , 2012, 107, 534-541.	1.1	28
3	Lysosomal lipase deficiency: Molecular characterization of eleven patients with Wolman or cholesteryl ester storage disease. <i>Molecular Genetics and Metabolism</i> , 2012, 105, 450-456.	1.1	71
4	Cardiovascular risk profile in a patient with a novel splice site ABCA1 mutation (HDL) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 622 Td (Chel	0.8	0
5	Identification and characterization of novel loss of function mutations in ATP-binding cassette transporter A1 in patients with low plasma high-density lipoprotein cholesterol. <i>Atherosclerosis</i> , 2010, 213, 492-498.	0.8	46
6	Multiple abnormally spliced ABCA1 mRNAs caused by a novel splice site mutation of ABCA1 gene in a patient with Tangier disease. <i>Clinica Chimica Acta</i> , 2010, 411, 524-530.	1.1	15
7	Severe HDL deficiency due to novel defects in the ABCA1 transporter. <i>Journal of Internal Medicine</i> , 2009, 265, 359-372.	6.0	24
8	Degradation of LDLR protein mediated by "gain of function" PCSK9 mutants in normal and ARH cells. <i>Atherosclerosis</i> , 2009, 203, 166-171.	0.8	64
9	A novel mutation of the apolipoprotein A-I gene in a family with familial combined hyperlipidemia. <i>Atherosclerosis</i> , 2008, 198, 145-151.	0.8	9
10	Effect of ezetimibe coadministered with statins in genotype-confirmed heterozygous FH patients. <i>Atherosclerosis</i> , 2007, 194, e116-e122.	0.8	68
11	PO5-152 FUNCTIONAL CHARACTERIZATION OF ABCA1 GENE MUTANTS. <i>Atherosclerosis Supplements</i> , 2007, 8, 55.	1.2	0
12	A Novel Loss of Function Mutation of PCSK9 Gene in White Subjects With Low-Plasma Low-Density Lipoprotein Cholesterol. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2007, 27, 677-681.	2.4	125
13	Coding sequence and intron-exon junctions of the c-myb gene are intact in the chronic phase and blast crisis stages of chronic myeloid leukemia patients. <i>Leukemia Research</i> , 2007, 31, 163-167.	0.8	5
14	Mo-W5:3 Molecular diagnosis and treatment of hypobetalipoproteinemia. <i>Atherosclerosis Supplements</i> , 2006, 7, 19-20.	1.2	0
15	Mo-P6:413 Genotype-phenotype relationship in LCAT deficiencies. <i>Atherosclerosis Supplements</i> , 2006, 7, 136-137.	1.2	0
16	Denaturing high-performance liquid chromatography in the detection of ABCA1 gene mutations in familial HDL deficiency. <i>Journal of Lipid Research</i> , 2005, 46, 817-822.	4.2	16
17	Familial HDL deficiency due to ABCA1 gene mutations with or without other genetic lipoprotein disorders. <i>Atherosclerosis</i> , 2004, 172, 309-320.	0.8	47