Masako Ueda

List of Publications by Year
in descending order

11 papers	426 citations	1163117 8 h-index	1281871 11 g-index
11 all docs	11 docs citations	11 times ranked	682 citing authors

#	Article	IF	Citations
1	Apolipoprotein C-II: New findings related to genetics, biochemistry, and role in triglyceride metabolism. Atherosclerosis, 2017, 267, 49-60.	0.8	148
2	Long-term follow-up of well-treated nephropathic cystinosis patients. Journal of Pediatrics, 2004, 145, 555-560.	1.8	113
3	A Novel Generalized Lipodystrophy-Associated Progeroid Syndrome Due to Recurrent Heterozygous LMNA p.T10I Mutation. Journal of Clinical Endocrinology and Metabolism, 2018, 103, 1005-1014.	3.6	47
4	Coronary Artery and Other Vascular Calcifications in Patients with Cystinosis after Kidney Transplantation. Clinical Journal of the American Society of Nephrology: CJASN, 2006, 1, 555-562.	4.5	40
5	A Novel APOC2 Missense Mutation Causing Apolipoprotein C-II Deficiency With Severe Triglyceridemia and Pancreatitis. Journal of Clinical Endocrinology and Metabolism, 2017, 102, 1454-1457.	3.6	35
6	Familial hypercholesterolemia. Molecular Genetics and Metabolism, 2005, 86, 423-426.	1.1	13
7	Prevention of polyurethane oxidative degradation with phenolic antioxidants covalently attached to the hard segments: Structureâ€“function relationships. Journal of Biomedical Materials Research - Part A, 2010, 94A, 751-759.	4.0	12
8	CD47-dependent molecular mechanisms of blood outgrowth endothelial cell attachment on cholesterol-modified polyurethane. Biomaterials, 2010, 31, 6394-6399.	11.4	9
9	Familial Chylomicronemia Syndrome With a Novel Homozygous LPL Mutation Identified in Three Siblings in Their 50s. Annals of Internal Medicine, 2020, 172, 500.	3.9	4
10	Experimental Therapeutics for Challenging Clinical Care of a Patient with an Extremely Rare Homozygous APOC2 Mutation. Case Reports in Endocrinology, 2020, 2020, 1-6.	0.4	4
11	Rare Diagnosis of Familial Partial Lipodystrophy in a Patient With Life-Threatening Pancreatitis due to Hypertriglyceridemia. AACE Clinical Case Reports, 2022, 8, 11-14.	1.1	1