Jos C Jansen

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Defective Lipid Droplet–Lysosome Interaction Causes Fatty Liver Disease as Evidenced by Human Mutations in TMEM199 and CCDC115. Cellular and Molecular Gastroenterology and Hepatology, 2022, 13, 583-597.	4.5	8
2	Screening for abnormal glycosylation in a cohort of adult liver disease patients. Journal of Inherited Metabolic Disease, 2020, 43, 1310-1320.	3.6	6
3	Mutations in the Vâ€ATPase Assembly Factor VMA21 Cause a Congenital Disorder of Glycosylation With Autophagic Liver Disease. Hepatology, 2020, 72, 1968-1986.	7.3	32
4	NAFLD Phenotype in Patients With V-ATPase Proton Pump Assembly Defects. Cellular and Molecular Gastroenterology and Hepatology, 2018, 5, 415-417.e1.	4.5	0
5	CCDC115 Deficiency Causes a Disorder of Golgi Homeostasis with Abnormal Protein Glycosylation. American Journal of Human Genetics, 2016, 98, 310-321.	6.2	88
6	TMEM199 Deficiency Is a Disorder of Golgi Homeostasis Characterized by Elevated Aminotransferases, Alkaline Phosphatase, and Cholesterol and Abnormal Glycosylation. American Journal of Human Genetics, 2016, 98, 322-330.	6.2	73