

# Jeannette C Bleeker

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/1960745/publications.pdf>

Version: 2024-02-01

10  
papers

353  
citations

1307594

7  
h-index

1372567

10  
g-index

10  
all docs

10  
docs citations

10  
times ranked

539  
citing authors

#	ARTICLE	IF	CITATIONS
1	Disorders of mitochondrial long-chain fatty acid oxidation and the carnitine shuttle. <i>Reviews in Endocrine and Metabolic Disorders</i> , 2018, 19, 93-106.	5.7	215
2	Impact of newborn screening for very long-chain acyl-CoA dehydrogenase deficiency on genetic, enzymatic, and clinical outcomes. <i>Journal of Inherited Metabolic Disease</i> , 2019, 42, 414-423.	3.6	36
3	Nutritional ketosis improves exercise metabolism in patients with very long-chain acyl-CoA dehydrogenase deficiency. <i>Journal of Inherited Metabolic Disease</i> , 2020, 43, 787-799.	3.6	26
4	Electrophysiological Abnormalities in VLCAD Deficient hiPSC-Cardiomyocytes Can Be Improved by Lowering Accumulation of Fatty Acid Oxidation Intermediates. <i>International Journal of Molecular Sciences</i> , 2020, 21, 2589.	4.1	24
5	Proposal for an individualized dietary strategy in patients with very long-chain acyl-CoA dehydrogenase deficiency. <i>Journal of Inherited Metabolic Disease</i> , 2019, 42, 159-168.	3.6	21
6	Subclinical effects of long-chain fatty acid $\beta$ -oxidation deficiency on the adult heart: A case-control magnetic resonance study. <i>Journal of Inherited Metabolic Disease</i> , 2020, 43, 969-980.	3.6	11
7	Exploring the metabolic fate of medium-chain triglycerides in healthy individuals using a stable isotope tracer. <i>Clinical Nutrition</i> , 2021, 40, 1396-1404.	5.0	8
8	Severe Fat Accumulation in Multiple Organs in Pediatric Autopsies. <i>Pediatric and Developmental Pathology</i> , 2017, 20, 269-276.	1.0	5
9	Electrophysiological Abnormalities in VLCAD Deficient hiPSC-Cardiomyocytes Do not Improve with Carnitine Supplementation. <i>Frontiers in Pharmacology</i> , 2020, 11, 616834.	3.5	5
10	Proposal for an individualized dietary strategy in patients with very long-chain acyl-CoA dehydrogenase deficiency. <i>Journal of Inherited Metabolic Disease</i> , 2019, 42, 159.	3.6	2