

# Danielle te Vruchte

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/3213707/publications.pdf>

Version: 2024-02-01

9  
papers

376  
citations

1307594

7  
h-index

1474206

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9  
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9  
docs citations

9  
times ranked

652  
citing authors

#	ARTICLE	IF	CITATIONS
1	A novel, highly sensitive and specific biomarker for Niemann-Pick type C1 disease. <i>Orphanet Journal of Rare Diseases</i> , 2015, 10, 78.	2.7	105
2	A novel approach to analyze lysosomal dysfunctions through subcellular proteomics and lipidomics: the case of NPC1 deficiency. <i>Scientific Reports</i> , 2017, 7, 41408.	3.3	93
3	Identification of novel bile acids as biomarkers for the early diagnosis of Niemann-Pick C disease. <i>FEBS Letters</i> , 2016, 590, 1651-1662.	2.8	82
4	Annual severity increment score as a tool for stratifying patients with Niemann-Pick disease type C and for recruitment to clinical trials. <i>Orphanet Journal of Rare Diseases</i> , 2018, 13, 143.	2.7	41
5	Glycosphingolipid metabolism and its role in ageing and Parkinson's disease. <i>Glycoconjugate Journal</i> , 2022, 39, 39-53.	2.7	18
6	NMR analysis reveals significant differences in the plasma metabolic profiles of Niemann Pick C1 patients, heterozygous carriers, and healthy controls. <i>Scientific Reports</i> , 2017, 7, 6320.	3.3	17
7	Glycosphingolipid storage leads to the enhanced degradation of the B cell receptor in Sandhoff disease mice. <i>Journal of Inherited Metabolic Disease</i> , 2010, 33, 261-270.	3.6	12
8	Measuring relative lysosomal volume for monitoring lysosomal storage diseases. <i>Methods in Cell Biology</i> , 2015, 126, 331-347.	1.1	4
9	Urinary excretion and metabolism of miglustat and valproate in patients with Niemann-Pick type C1 disease: One- and two-dimensional solution-state <sup>1</sup> H NMR studies. <i>Journal of Pharmaceutical and Biomedical Analysis</i> , 2016, 117, 276-288.	2.8	4