

# Guido Vogt

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/5972738/publications.pdf>

Version: 2024-02-01

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2258059

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2550090

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#	ARTICLE	IF	CITATIONS
1	Biallelic truncating variants in <i>ATP9A</i> cause a novel neurodevelopmental disorder involving postnatal microcephaly and failure to thrive. <i>Journal of Medical Genetics</i> , 2022, 59, 662-668.	3.2	9
2	Expanding the clinical and molecular spectrum of <i>ATP6V1A</i> related metabolic cutis laxa. <i>Journal of Inherited Metabolic Disease</i> , 2021, 44, 972-986.	3.6	7
3	A CRISPR-Cas9 engineered mouse model for GPI-anchor deficiency mirrors human phenotypes and exhibits hippocampal synaptic dysfunctions. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2021, 118, .	7.1	8