

# Andrea Mancikova

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/9329831/publications.pdf>

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8  
papers

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1684129  
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#	ARTICLE	IF	CITATIONS
1	Functional Characterization of Rare Variants in OAT1/SLC22A6 and OAT3/SLC22A8 Urate Transporters Identified in a Gout and Hyperuricemia Cohort. <i>Cells</i> , 2022, 11, 1063.	4.1	9
2	A homozygous deletion in the SLC19A1 gene as a cause of folate-dependent recurrent megaloblastic anemia. <i>Blood</i> , 2020, 135, 2427-2431.	1.4	13
3	Clinical and Functional Characterization of a Novel URAT1 Dysfunctional Variant in a Pediatric Patient with Renal Hypouricemia. <i>Applied Sciences (Switzerland)</i> , 2019, 9, 3479.	2.5	2
4	Functional Characterization of Clinically-Relevant Rare Variants in ABCG2 Identified in a Gout and Hyperuricemia Cohort. <i>Cells</i> , 2019, 8, 363.	4.1	46
5	Folate-Dependent Normocytic Anemia Caused By a Hypomorphic Mutation in SLC19A1 gene. <i>Blood</i> , 2018, 132, 502-502.	1.4	0
6	Functional analysis of novel allelic variants in URAT1 and GLUT9 causing renal hypouricemia type 1 and 2. <i>Clinical and Experimental Nephrology</i> , 2016, 20, 578-584.	1.6	35
7	SP013IDIOPATHIC RENAL HYPOURICEMIA: IDENTIFICATION AND CHARACTERIZATION OF SLC22A12 MUTATIONS IN SPANISH PATIENTS. <i>Nephrology Dialysis Transplantation</i> , 2015, 30, iii385-iii385.	0.7	0
8	Complex Analysis of Urate Transporters SLC2A9, SLC22A12 and Functional Characterization of Non-Synonymous Allelic Variants of GLUT9 in the Czech Population: No Evidence of Effect on Hyperuricemia and Gout. <i>PLoS ONE</i> , 2014, 9, e107902.	2.5	33