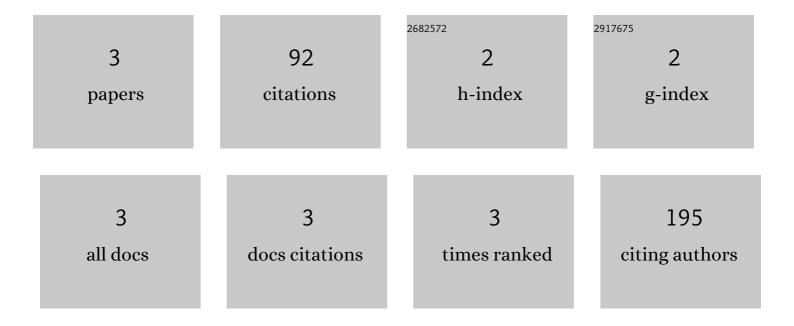
Vaksha Patel

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/2143661/publications.pdf Version: 2024-02-01



#	Article	IF	CITATIONS
1	A Founder Mutation in EHD1 Presents with Tubular Proteinuria and Deafness. Journal of the American Society of Nephrology: JASN, 2022, 33, 732-745.	6.1	7
2	Quantification of FAM20A in human milk and identification of calcium metabolism proteins. Physiological Reports, 2021, 9, e15150.	1.7	1
3	Nephrocalcinosis (Enamel Renal Syndrome) Caused by Autosomal Recessive FAM20A Mutations. Nephron Physiology, 2013, 122, 1-6.	1.2	84