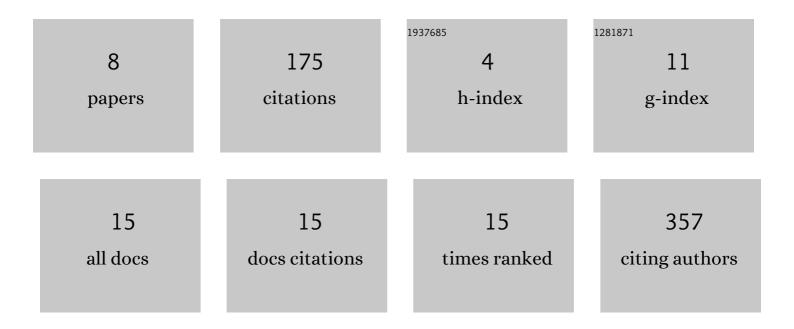


## List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/2172094/publications.pdf Version: 2024-02-01



WENLI

#	Article	IF	CITATIONS
1	<i>DMC1</i> mutation that causes human non-obstructive azoospermia and premature ovarian insufficiency identified by whole-exome sequencing. Journal of Medical Genetics, 2018, 55, 198-204.	3.2	91
2	Noninvasive prenatal testing (NIPT) in twin pregnancies with treatment of assisted reproductive techniques (ART) in a single center. Prenatal Diagnosis, 2016, 36, 672-679.	2.3	49
3	Novel <i><scp>FOXL</scp>2</i> mutations cause blepharophimosisâ€ptosisâ€epicanthus inversus syndrome with premature ovarian insufficiency. Molecular Genetics & Genomic Medicine, 2018, 6, 261-267.	1.2	14
4	Rare partial octosomy and hexasomy of 15q11-q13 associated with intellectual impairment and development delay: report of two cases and review of literature. Molecular Cytogenetics, 2018, 11, 15.	0.9	5
5	Cadmium Removal by Pseudomonas aeruginosa E1. , 2009, , .		4
6	Expanded carrier screening and preimplantation genetic diagnosis in a couple who delivered a baby affected with congenital factor VII deficiency. BMC Medical Genetics, 2018, 19, 15.	2.1	3
7	A case of megalencephalic leukoencephalopathy with subcortical cysts type 1 was identified with a novel compound heterozygous alteration (c.135delC; c.423+2dupT) in China. Clinical Case Reports (discontinued), 2017, 5, 961-967.	0.5	2
8	The Design of a High Efficient DNA In-Situ Synthesis Apparatus Based on the Principle of Typography Technique. , 2009, , .		0