

# Maria F Bedeschi

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/7264097/publications.pdf>

Version: 2024-02-01

9  
papers

209  
citations

1162889

8  
h-index

1474057

9  
g-index

10  
all docs

10  
docs citations

10  
times ranked

464  
citing authors

#	ARTICLE	IF	CITATIONS
1	Moebius syndrome: clinical features, diagnosis, management and early intervention. Italian Journal of Pediatrics, 2016, 42, 56.	1.0	62
2	Clinical Characterization, Genetics, and Long-Term Follow-up of a Large Cohort of Patients With Agenesis of the Corpus Callosum. Journal of Child Neurology, 2017, 32, 60-71.	0.7	34
3	Phenotype delineation of <i>ZNF462</i> related syndrome. American Journal of Medical Genetics, Part A, 2019, 179, 2075-2082.	0.7	23
4	Childhood-onset dystonia-causing KMT2B variants result in a distinctive genomic hypermethylation profile. Clinical Epigenetics, 2021, 13, 157.	1.8	22
5	Biallelic MADD variants cause a phenotypic spectrum ranging from developmental delay to a multisystem disorder. Brain, 2020, 143, 2437-2453.	3.7	21
6	A Genetics-First Approach Revealed Monogenic Disorders in Patients With ARM and VACTERL Anomalies. Frontiers in Pediatrics, 2020, 8, 310.	0.9	17
7	The spatiotemporal organization of episodic memory and its disruption in a neurodevelopmental disorder. Scientific Reports, 2019, 9, 18447.	1.6	13
8	(Epi)genetic profiling of extraembryonic and postnatal tissues from female monozygotic twins discordant for Beckwith-Wiedemann syndrome. Molecular Genetics & Genomic Medicine, 2020, 8, e1386.	0.6	9
9	STAR syndrome plus: The first description of a female patient with the lethal form. American Journal of Medical Genetics, Part A, 2017, 173, 3226-3230.	0.7	8