

# Eva Pinti

## List of Publications by Year in descending order

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#	ARTICLE	IF	CITATIONS
1	Diagnostic difficulties and possibilities of NF1-like syndromes in childhood. BMC Pediatrics, 2021, 21, 331.	1.7	6
2	Clinical and genetic findings in Hungarian pediatric patients carrying chromosome 16p copy number variants and a review of the literature. European Journal of Medical Genetics, 2020, 63, 104027.	1.3	4
3	Similar Cause, Different Phenotype: <i>SOX9</i> Enhancer Duplication in a Family. Hormone Research in Paediatrics, 2019, 92, 335-339.	1.8	3
4	What should we consider in the case of combined Down- and 47,XY,+i(X)(q10) Klinefelter syndromes? The unique case of a male newborn and review of the literature. BMC Pediatrics, 2020, 20, 17.	1.7	3
5	Microdeletions in 1q21 and 8q12.1 depict two additional molecular subgroups of Silver-Russell syndrome like phenotypes. Molecular Cytogenetics, 2022, 15, 19.	0.9	3
6	Deletion of 16q22.2q23.3 in a Boy with a Phenotype Reminiscent of Silver-Russell Syndrome. Molecular Syndromology, 2021, 12, 300-304.	0.8	1
7	Chromosome 2q14.3 microdeletion encompassing CNTNAP5 gene in a patient carrying a complex chromosomal rearrangement. Journal of Genetics, 2021, 100, 1.	0.7	1