# Eva Pinti

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/5135993/publications.pdf

Version: 2024-02-01

| 7 | 21 | 2258059<br>3 | 2053705<br>5 |
|---|---|---|---|
| papers | citations | h-index | g-index |

| 9 | 9 | 9 | 7 |
|---|---|---|---|
| all docs | docs citations | times ranked | citing authors |

| # | Article | IF | Citations |
|---|---|---|---|
| 1 | 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 |
| 2 | 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 |
| 3 | Diagnostic difficulties and possibilities of NF1-like syndromes in childhood. BMC Pediatrics, 2021, 21, 331. | 1.7 | 6 |
| 4 | Chromosome 2q14.3 microdeletion encompassing CNTNAP5 gene in a patient carrying a complex chromosomal rearrangement. Journal of Genetics, 2021, 100, 1. | 0.7 | 1 |
| 5 | 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 |
| 6 | 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 |
| 7 | Similar Cause, Different Phenotype: SOX9 Enhancer Duplication in a Family. Hormone Research in Paediatrics, 2019, 92, 335-339. | 1.8 | 3 |