# Alina German

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/7628952/publications.pdf

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|---|---|---|---|
| 12 papers | 126 citations | 1307594 7 h-index | 1372567 10 g-index |
| 12 all docs | 12 docs citations | 12 times ranked | 203 citing authors |

| # | Article | IF | Citations |
|---|---|---|---|
| 1 | Alarming increase in ketoacidosis in children and adolescents with newly diagnosed type 1 diabetes during the first wave of the <scp>COVID</scp> â€19 pandemic in Israel. Pediatric Diabetes, 2022, 23, 10-18. | 2.9 | 26 |
| 2 | Human Type 1 Iodothyronine Deiodinase (DIO1) Mutations Cause Abnormal Thyroid Hormone Metabolism. Thyroid, 2021, 31, 202-207. | 4.5 | 25 |
| 3 | Combined Gestational Age- and Birth Weightâ€"Adjusted Cutoffs for Newborn Screening of Congenital Adrenal Hyperplasia. Journal of Clinical Endocrinology and Metabolism, 2019, 104, 3172-3180. | 3.6 | 22 |
| 4 | Global Application of the Assessment of Communication Skills of Paediatric Endocrinology Fellows in the Management of Differences in Sex Development Using the ESPE E-Learning.Org Portal. Hormone Research in Paediatrics, 2017, 88, 127-139. | 1.8 | 13 |
| 5 | Environmental Rather than Genetic Factors Determine the Variation inÂtheÂAge of the Infancy to Childhood Transition: AÂTwins Study. Journal of Pediatrics, 2015, 166, 731-735. | 1.8 | 10 |
| 6 | Sexual Dimorphism of Size Ontogeny and Life History. Frontiers in Pediatrics, 2020, 8, 387. | 1.9 | 9 |
| 7 | Prediction of Adult Height by Machine Learning Technique. Journal of Clinical Endocrinology and Metabolism, 2021, 106, e2700-e2710. | 3.6 | 8 |
| 8 | Primary Ovarian Insufficiency Nationwide Incidence Rate and Etiology Among Israeli Adolescents. Journal of Adolescent Health, 2020, 66, 603-609. | 2.5 | 7 |
| 9 | Periorbital inflammation associated with craniofacial fibrous dysplasia: Report of three cases and review of the literature. Bone, 2021, 153, 116157. | 2.9 | 4 |
| 10 | OR01-01 Human Type 1 Iodothyronine Deiodinase (DIO1) Mutations Cause Abnormal Thyroid Hormone Metabolism. Journal of the Endocrine Society, 2020, 4, . | 0.2 | 2 |
| 11 | A novel truncating variant in the FGD1 gene associated with Aarskogâ€"Scott syndrome in a family previously diagnosed with Tel Hashomer camptodactyly. American Journal of Medical Genetics, Part A, 2021, 185, 3161-3166. | 1.2 | 0 |
| 12 | Family Size and the Age at Infancy-Childhood Transition Determine a Childâ€™s Compromised Growth in Large Families. Frontiers in Pediatrics, 2022, 10, 821048. | 1.9 | 0 |