

# Chao-Chun Zou

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/4629985/publications.pdf>

Version: 2024-02-01

15  
papers

101  
citations

1478505

6  
h-index

1474206

9  
g-index

20  
all docs

20  
docs citations

20  
times ranked

89  
citing authors

| #  | ARTICLE   | IF  | CITATIONS |
|----|---|-----|-----------|
| 1  | Application of next generation sequencing in the screening of monogenic diseases in China, 2021: a consensus among Chinese newborn screening experts. <i>World Journal of Pediatrics</i> , 2022, 18, 235-242. | 1.8 | 16        |
| 2  | Microdeletion of 4p16.2 in Children: A Case Report and Literature Review. <i>Case Reports in Genetics</i> , 2022, 2022, 1-4.  | 0.2 | 0         |
| 3  | Do patients with Prader-Willi syndrome have favorable glucose metabolism?. <i>Orphanet Journal of Rare Diseases</i> , 2022, 17, 187.  | 2.7 | 4         |
| 4  | Recommendations for the diagnosis and management of childhood Prader-Willi syndrome in China. <i>Orphanet Journal of Rare Diseases</i> , 2022, 17, .  | 2.7 | 8         |
| 5  | 6q25.1-q25.3 Microdeletion in a Chinese Girl. <i>JCRPE Journal of Clinical Research in Pediatric Endocrinology</i> , 2021, 13, 109-113.   | 0.9 | 1         |
| 6  | Regulatory roles and mechanisms of alternative RNA splicing in adipogenesis and human metabolic health. <i>Cell and Bioscience</i> , 2021, 11, 66.  | 4.8 | 26        |
| 7  | Prader-Willi Syndrome Coincident with DiGeorge Syndrome. <i>Indian Journal of Pediatrics</i> , 2020, 87, 471-472.   | 0.8 | 1         |
| 8  | Clinical Features of Patients With 7p22.1 Microdeletion. <i>Indian Pediatrics</i> , 2020, 57, 580-581.  | 0.4 | 0         |
| 9  | Prader-Willi Syndrome: Molecular Mechanism and Epigenetic Therapy. <i>Current Gene Therapy</i> , 2020, 20, 36-43.   | 2.0 | 7         |
| 10 | Clinical Features of Patients With 7p22.1 Microdeletion. <i>Indian Pediatrics</i> , 2020, 57, 580-581.  | 0.4 | 0         |
| 11 | Quality of life in caregivers of young children with Prader-Willi syndrome. <i>World Journal of Pediatrics</i> , 2019, 15, 506-510.   | 1.8 | 9         |
| 12 | Clinical features and genetic analysis of childhood sitosterolemia. <i>Medicine (United States)</i> , 2019, 98, e15013.   | 1.0 | 13        |
| 13 | 13q22.1-q32.1 Microdeletion Syndrome. <i>Indian Journal of Pediatrics</i> , 2019, 86, 303-305.  | 0.8 | 0         |
| 14 | Clinical Features in Patients with Microdeletion at 6q14.1-q15. <i>Indian Journal of Pediatrics</i> , 2017, 84, 883-886.  | 0.8 | 1         |
| 15 | Glucose metabolism disorder in obese children assessed by continuous glucose monitoring system. <i>World Journal of Pediatrics</i> , 2008, 4, 26-30.  | 1.8 | 13        |