Kazumoto Iijima

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/7361986/publications.pdf

Version: 2024-02-01

| 16 papers | 257 citations | 1040056 9 h-index | 1058476 14 g-index |
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| Papero | | | 5 Muon |
| 16 all docs | 16 docs citations | 16 times ranked | 331 citing authors |

| # | Article | IF | CITATIONS |
|----|--|-----|-----------|
| 1 | Rituximab for nephrotic syndrome in children. Clinical and Experimental Nephrology, 2017, 21, 193-202. | 1.6 | 62 |
| 2 | Rituximab therapy for refractory steroid-resistant nephrotic syndrome in children. Pediatric Nephrology, 2020, 35, 17-24. | 1.7 | 41 |
| 3 | Rituximab in steroid-sensitive nephrotic syndrome: lessons from clinical trials. Pediatric Nephrology, 2018, 33, 1449-1455. | 1.7 | 33 |
| 4 | Biallelic variants in <i>LIG3</i> cause a novel mitochondrial neurogastrointestinal encephalomyopathy. Brain, 2021, 144, 1451-1466. | 7.6 | 28 |
| 5 | Study protocol: multicenter double-blind, randomized, placebo-controlled trial of rituximab for the treatment of childhood-onset early-stage uncomplicated frequently relapsing or steroid-dependent nephrotic syndrome (JSKDC10 trial). BMC Nephrology, 2019, 20, 293. | 1.8 | 13 |
| 6 | Genetic background, recent advances in molecular biology, and development of novel therapy in Alport syndrome. Kidney Research and Clinical Practice, 2020, 39, 402-413. | 2.2 | 13 |
| 7 | An updated view of the pathogenesis of steroid-sensitive nephrotic syndrome. Pediatric Nephrology, 2022, 37, 1957-1965. | 1.7 | 13 |
| 8 | Diversity of renal phenotypes in patients with <i>WDR19</i> mutations: Two case reports. Nephrology, 2017, 22, 566-571. | 1.6 | 12 |
| 9 | Study protocol: mycophenolate mofetil as maintenance therapy after rituximab treatment for childhood-onset, complicated, frequently-relapsing nephrotic syndrome or steroid-dependent nephrotic syndrome: a multicenter double-blind, randomized, placebo-controlled trial (JSKDC07). BMC Nephrology, 2018, 19, 302. | 1.8 | 10 |
| 10 | Involvement of WNT Signaling in the Regulation of Gestational Age-Dependent Umbilical Cord-Derived Mesenchymal Stem Cell Proliferation. Stem Cells International, 2017, 2017, 1-16. | 2.5 | 9 |
| 11 | A Novel Method for Measuring Serum Unbound Bilirubin Levels Using Glucose Oxidase–Peroxidase and Bilirubin-Inducible Fluorescent Protein (UnaG): No Influence of Direct Bilirubin. International Journal of Molecular Sciences, 2020, 21, 6778. | 4.1 | 9 |
| 12 | Clear Evidence of LAMA5 Gene Biallelic Truncating Variants Causing Infantile Nephrotic Syndrome. Kidney360, 2021, 2, 1968-1978. | 2.1 | 8 |
| 13 | Biallelic variants/mutations of IL1RAP in patients with steroid-sensitive nephrotic syndrome. International Immunology, 2020, 32, 283-292. | 4.0 | 3 |
| 14 | Evaluation of suspected autosomal Alport Syndrome synonymous variants. Kidney360, 2022, 3, 10.34067/KID.0005252021. | 2.1 | 3 |
| 15 | Identification of novel OCRL isoforms associated with phenotypic differences between Dent disease-2 and Lowe syndrome. Nephrology Dialysis Transplantation, 2021, , . | 0.7 | O |
| 16 | Preliminary Effectiveness and Safety of High Frequency Oscillation in Addition to Mechanical Insufflation and Exsufflation for Intratracheal Mucus Removal in Patients With Neuromuscular Disease: Protocol for a Prospective Study. JMIR Research Protocols, 2019, 8, e12102. | 1.0 | 0 |