

Lale Sever

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/5228515/publications.pdf>

Version: 2024-02-01

19
papers

96
citations

1684188

5
h-index

1588992

8
g-index

19
all docs

19
docs citations

19
times ranked

89
citing authors

#	ARTICLE	IF	CITATIONS
1	Disasters, children and the kidneys. <i>Pediatric Nephrology</i> , 2020, 35, 1381-1393.	1.7	21
2	Renal Crisis in Children during Armed Conflict. <i>Seminars in Nephrology</i> , 2020, 40, 408-420.	1.6	12
3	The Frequency of Familial Congenital Anomalies of the Kidney and Urinary Tract: Should We Screen Asymptomatic First-Degree Relatives Using Urinary Tract Ultrasonography?. <i>Nephron</i> , 2020, 144, 170-175.	1.8	9
4	Factors influencing blood pressure and microalbuminuria in children with type 1 diabetes mellitus: salt or sugar?. <i>Pediatric Nephrology</i> , 2020, 35, 1267-1276.	1.7	8
5	Maintenance Peritoneal Dialysis in Children With Autosomal Recessive Polycystic Kidney Disease: A Comparative Cohort Study of the International Pediatric Peritoneal Dialysis Network Registry. <i>American Journal of Kidney Diseases</i> , 2020, 75, 460-464.	1.9	8
6	Evaluation of non-infectious complications of peritoneal dialysis in children: a multicenter study. <i>Pediatric Nephrology</i> , 2021, 36, 417-423.	1.7	6
7	AGTR1-related Renal Tubular Dysgeneses May Not Be Fatal. <i>Kidney International Reports</i> , 2021, 6, 846-852.	0.8	6
8	Phenotypic Variability in Siblings With Autosomal Recessive Polycystic Kidney Disease. <i>Kidney International Reports</i> , 2022, 7, 1643-1652.	0.8	6
9	A homozygous <i>HOXA11</i> variation as a potential novel cause of autosomal recessive congenital anomalies of the kidney and urinary tract. <i>Clinical Genetics</i> , 2020, 98, 390-395.	2.0	5
10	A splice site mutation in the <i>TSEN2</i> causes a new syndrome with craniofacial and central nervous system malformations, and atypical hemolytic uremic syndrome. <i>Clinical Genetics</i> , 2022, 101, 346-358.	2.0	4
11	Rituximab treatment for difficult-to-treat nephrotic syndrome in children: a multicenter, retrospective study. <i>Turkish Journal of Medical Sciences</i> , 2021, 51, 1781-1790.	0.9	3
12	Different approaches among physicians to treat pediatric stone disease: a survey-based study. <i>Archivos Argentinos De Pediatría</i> , 2021, 119, 83-90.	0.2	3
13	Increased risk for kidney sequelae surrogates in survivors of Wilms tumor. <i>Pediatric Nephrology</i> , 2022, 37, 2415-2426.	1.7	2
14	Anemia after kidney transplantation: Does its basis differ from anemia in chronic kidney disease?. <i>Pediatric Transplantation</i> , 2020, 24, e13818.	1.0	1
15	Natural history of patients with infantile nephrolithiasis: what are the predictors of surgical intervention?. <i>Pediatric Nephrology</i> , 2021, 36, 939-944.	1.7	1
16	Strong mesangial IgA staining “does it always refer to IgA nephropathy in a patient with proteinuria and hematuria? Answers. <i>Pediatric Nephrology</i> , 2021, 36, 2043-2045.	1.7	1
17	A rare cause of proteinuria after kidney transplantation: Answers. <i>Pediatric Nephrology</i> , 2019, 34, 2333-2335.	1.7	0
18	A rare cause of proteinuria after kidney transplantation: Questions. <i>Pediatric Nephrology</i> , 2019, 34, 2331-2332.	1.7	0

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19	Strong mesangial IgA staining“does it always refer to IgA nephropathy in a patient with proteinuria and hematuria? Questions. Pediatric Nephrology, 2021, 36, 2039-2041.	1.7	0