## **Bahriye Atmis**

List of Publications by Year in descending order

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1937685 1720034 64 19 4 7 citations h-index g-index papers 19 19 19 109 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	MEFV gene variants in children with Henoch-SchĶnlein purpura and association with clinical manifestations: a single-center Mediterranean experience. Postgraduate Medicine, 2019, 131, 68-72.	2.0	14
2	Evaluation of phenotypic and genotypic features of children with distal kidney tubular acidosis. Pediatric Nephrology, 2020, 35, 2297-2306.	1.7	12
3	Kidney disease profile and encountered problems during follow-up in Syrian refugee children: a multicenter retrospective study. Pediatric Nephrology, 2022, 37, 393-402.	1.7	9
4	An infant with hyponatremia, hyperkalemia, and metabolic acidosis associated with urinary tract infection: Answers. Pediatric Nephrology, 2019, 34, 1739-1741.	1.7	5
5	From infancy to adulthood: challenges in congenital nephrogenic diabetes insipidus. Journal of Pediatric Endocrinology and Metabolism, 2020, 33, 1019-1025.	0.9	5
6	Phenotypic variability in two patients with tumor necrosis factor receptor associated periodic fever syndrome emphasizes a rare manifestation: Immunoglobulin A nephropathy. European Journal of Medical Genetics, 2020, 63, 103780.	1.3	4
7	Bedside sonographic assessments for predicting predialysis fluid overload in children with end-stage kidney disease. European Journal of Pediatrics, 2021, 180, 3191-3200.	2.7	4
8	More than tubular dysfunction: cystinosis and kidney outcomes. Journal of Nephrology, 2022, 35, 831-840.	2.0	3
9	Community-acquired Pediatric Urinary Tract Infections Caused by Morganella Morganii. Journal of Pediatric Research, 2020, 7, 121-125.	0.2	3
10	Time-averaged hemoglobin values, not hemoglobin cycling, have an impact on outcomes in pediatric dialysis patients. Pediatric Nephrology, 2018, 33, 2143-2150.	1.7	2
11	A rare manifestation of renal osteodystrophy in a non-compliant child on hemodialysis: Answers. Pediatric Nephrology, 2016, 31, 1451-1453.	1.7	1
12	An infant with hyponatremia, hyperkalemia, and metabolic acidosis associated with urinary tract infection: Questions. Pediatric Nephrology, 2019, 34, 1737-1737.	1.7	1
13	Acquired Bartter-like Syndrome Presenting with Polyuria and Reversible Hypokalemia Associated with Colistin Use in a Critically Ill Pediatric Patient. Indian Journal of Critical Care Medicine, 2021, 25, 822-824.	0.9	1
14	A rare manifestation of renal osteodystrophy in a non-compliant hemodialysis child: Questions. Pediatric Nephrology, 2016, 31, 1449-1450.	1.7	0
15	P314â€Renal tract anomalies in children with congenital heart disease detected during the procedure of cardiac catheterization. , 2017, , .		O
16	Proptosis in a child with chronic kidney disease: Questions. Pediatric Nephrology, 2020, 35, 787-788.	1.7	0
17	Proptosis in a child with chronic kidney disease: Answers. Pediatric Nephrology, 2020, 35, 789-791.	1.7	О
18	Recurrent macroscopic hematuria in a pediatric patient: is it early to diagnose as having type I hereditary C2 deficiency?. CEN Case Reports, 2020, 9, 344-346.	0.9	0

#	Article	lF	CITATIONS
19	Henoch SchĶnlein Purpurası tanılı çocuklarda Kompleman C2 gen polimorfizmleri. Cukurova Medical Journal, 2020, 45, 89-95.	0.2	0