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List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/837810/publications.pdf

Version: 2024-02-01

		1478505	1474206
12	99	6	9
papers	citations	h-index	g-index
12	12	12	176
all docs	docs citations	times ranked	citing authors

#	Article	IF	CITATIONS
1	A teenager presenting with anuric acute renal failure and metabolic acidosis with a high anion gap: Questions. Pediatric Nephrology, 2020, 35, 2253-2255.	1.7	O
2	A teenager presenting with anuric acute renal failure and metabolic acidosis with a high anion gap: Answers. Pediatric Nephrology, 2020, 35, 2257-2258.	1.7	1
3	Kidney involvement and associated risk factors in children with Duchenne muscular dystrophy. Pediatric Nephrology, 2020, 35, 1953-1958.	1.7	8
4	Delayed diagnosis of primary vesicoureteral reflux in children with recurrent urinary tract infections: Diagnostic approach and renal outcomes. Turkish Journal of Urology, 2018, 44, 498-502.	1.3	6
5	A Rare Cause of Gross Hematuria in Childhood: Renal Lymphangiectasia. Journal of Pediatric Research, 2018, 5, 234-236.	0.2	О
6	Assessment of left ventricular function by tissue Doppler echocardiography in pediatric chronic kidney disease. Renal Failure, 2015, 37, 1094-1099.	2.1	14
7	Unilateral multicystic dysplastic kidney in children. Turkish Journal of Pediatrics, 2014, 56, 75-9.	0.6	16
8	Prevalence and significance of the MEFV gene mutations in childhood Henoch–Schönlein purpura without FMF symptoms. Rheumatology International, 2013, 33, 377-380.	3.0	33
9	Effect of Renal Graft on Longitudinal Growth in Prepubertal Children. Experimental and Clinical Transplantation, 2013, 11, 315-319.	0.5	O
10	Renal function and linear growth of children with nephrocalcinosis: a retrospective single-center study. Turkish Journal of Pediatrics, 2013, 55, 58-62.	0.6	3
11	Renal outcome of children with unilateral renal agenesis. Turkish Journal of Pediatrics, 2013, 55, 612-5.	0.6	11
12	A novel splice site mutation of the beta subunit gene of epithelial sodium channel (ENaC) in one Turkish patient with a systemic form of pseudohypoaldosteronism type 1. Journal of Pediatric Endocrinology and Metabolism, 2012, 25, 1035-9.	0.9	7