

Lawrence Copelovitch

List of Publications by Year in descending order

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Version: 2024-02-01

59
papers

1,927
citations

304743

22
h-index

254184

43
g-index

63
all docs

63
docs citations

63
times ranked

2233
citing authors

#	ARTICLE	IF	CITATIONS
1	A novel <i>MBTPS2</i> variant associated with BRESHECK syndrome impairs <i>sterol-regulated</i> transcription and the endoplasmic reticulum stress response. <i>American Journal of Medical Genetics, Part A</i> , 2022, 188, 463-472.	1.2	4
2	Long-term outcome among females with Alport syndrome from a single pediatric center. <i>Pediatric Nephrology</i> , 2021, 36, 945-951.	1.7	7
3	New insights into the pathogenesis of <i>Streptococcus pneumoniae</i> -associated hemolytic uremic syndrome. <i>Pediatric Nephrology</i> , 2020, 35, 1585-1591.	1.7	23
4	Predictors of time to first cannulation for arteriovenous fistula in pediatric hemodialysis patients: Midwest Pediatric Nephrology Consortium study. <i>Pediatric Nephrology</i> , 2020, 35, 287-295.	1.7	7
5	Inflammatory Bowel Diseases Are Associated With an Increased Risk for Chronic Kidney Disease, Which Decreases With Age. <i>Clinical Gastroenterology and Hepatology</i> , 2020, 18, 2262-2268.	4.4	31
6	Prevention of recurrent urinary stone disease. <i>Current Opinion in Pediatrics</i> , 2020, 32, 295-299.	2.0	7
7	Serum Calcification Propensity in Children on Chronic Hemodialysis. <i>Kidney International Reports</i> , 2020, 5, 1528-1531.	0.8	3
8	Medical Expulsive Therapy for Urinary Stone Disease in Children. <i>Indian Pediatrics</i> , 2020, 57, 940-943.	0.4	0
9	Reply. <i>Clinical Gastroenterology and Hepatology</i> , 2020, 19, 1994.	4.4	0
10	Medical Expulsive Therapy for Urinary Stone Disease in Children. <i>Indian Pediatrics</i> , 2020, 57, 940-943.	0.4	0
11	Update on Dent Disease. <i>Pediatric Clinics of North America</i> , 2019, 66, 169-178.	1.8	27
12	Predictors of patency for arteriovenous fistulae and grafts in pediatric hemodialysis patients. <i>Pediatric Nephrology</i> , 2019, 34, 329-339.	1.7	12
13	Uncommon cribfellow: an infant with hypercalcemia, nephrocalcinosis, and acidosis: Answers. <i>Pediatric Nephrology</i> , 2018, 33, 1697-1699.	1.7	6
14	Uncommon cribfellow: an infant with hypercalcemia, nephrocalcinosis, and acidosis: Questions. <i>Pediatric Nephrology</i> , 2018, 33, 1695-1695.	1.7	4
15	Pretreatment of enteral nutrition with sodium polystyrene sulfonate: effective, but beware the high prevalence of electrolyte derangements in clinical practice. <i>CKJ: Clinical Kidney Journal</i> , 2018, 11, 166-171.	2.9	18
16	Salt, sweat, and unclear? Diaphoresis and hypernatremia in end-stage kidney disease: Questions. <i>Pediatric Nephrology</i> , 2018, 33, 251-252.	1.7	1
17	Salt, sweat, and unclear? Diaphoresis and hypernatremia in end-stage kidney disease: Answers. <i>Pediatric Nephrology</i> , 2018, 33, 253-254.	1.7	0
18	Oral Antibiotic Exposure and Kidney Stone Disease. <i>Journal of the American Society of Nephrology: JASN</i> , 2018, 29, 1731-1740.	6.1	109

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19	Assessment of Sex Differences in Fracture Risk Among Patients With Anorexia Nervosa: A Population-Based Cohort Study Using The Health Improvement Network. <i>Journal of Bone and Mineral Research</i> , 2017, 32, 1082-1089.	2.8	47
20	MP95-01 ASSOCIATION BETWEEN ORAL ANTIBIOTICS AND INCIDENT KIDNEY STONES. <i>Journal of Urology</i> , 2017, 197, .	0.4	1
21	Dietary Zinc and Incident Calcium Kidney Stones in Adolescence. <i>Journal of Urology</i> , 2017, 197, 1342-1348.	0.4	16
22	Risk of Urolithiasis in Anorexia Nervosa: A Population-Based Cohort Study Using the Health Improvement Network. <i>European Eating Disorders Review</i> , 2017, 25, 406-410.	4.1	6
23	Variability in measures of mineral metabolism in children on hemodialysis: impact on clinical decision-making. <i>Pediatric Nephrology</i> , 2017, 32, 2311-2318.	1.7	5
24	Glomerular Pathology in Dent Disease and Its Association with Kidney Function. <i>Clinical Journal of the American Society of Nephrology: CJASN</i> , 2016, 11, 2168-2176.	4.5	47
25	Pediatric Kidney Stones—Avoidance and Treatment. <i>Current Treatment Options in Pediatrics</i> , 2016, 2, 104-111.	0.6	24
26	Post-streptococcal glomerulonephritis associated with atypical hemolytic uremic syndrome: to treat or not to treat with eculizumab?. <i>CKJ: Clinical Kidney Journal</i> , 2016, 9, 90-96.	2.9	10
27	Assessing the risk of incident hypertension and chronic kidney disease after exposure to shock wave lithotripsy and ureteroscopy. <i>Kidney International</i> , 2016, 89, 185-192.	5.2	49
28	Annual Incidence of Nephrolithiasis among Children and Adults in South Carolina from 1997 to 2012. <i>Clinical Journal of the American Society of Nephrology: CJASN</i> , 2016, 11, 488-496.	4.5	187
29	Making Every Drop Count for Pediatric Kidney Transplant Patients. <i>Hospital Pediatrics</i> , 2015, 5, 287-289.	1.3	1
30	Intravenous Fluid Management in the Pediatric Hospital Setting: Is Isotonic Fluid the Right Approach for all Patients?. <i>Current Treatment Options in Pediatrics</i> , 2015, 1, 90-99.	0.6	3
31	Risk of Fracture in Urolithiasis. <i>Clinical Journal of the American Society of Nephrology: CJASN</i> , 2014, 9, 2133-2140.	4.5	61
32	Current treatment of atypical hemolytic uremic syndrome. <i>Intractable and Rare Diseases Research</i> , 2014, 3, 34-45.	0.9	50
33	Nephrotic syndrome associated with tyrosine kinase inhibitors for pediatric malignancy: case series and review of the literature. <i>Pediatric Nephrology</i> , 2014, 29, 863-869.	1.7	33
34	Evaluation and Medical Management of Kidney Stones in Children. <i>Journal of Urology</i> , 2014, 192, 1329-1336.	0.4	122
35	Genetics and Urinary Tract Malformations. <i>American Journal of Kidney Diseases</i> , 2014, 63, 183-185.	1.9	0
36	PD28-12 NEPHROLITHIASIS AND FRACTURE RISK OVER THE LIFESPAN: A POPULATION-BASED STUDY USING THE HEALTH IMPROVEMENT NETWORK (THIN). <i>Journal of Urology</i> , 2014, 191, .	0.4	0

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37	Long-term outcomes of Shiga toxin hemolytic uremic syndrome. <i>Pediatric Nephrology</i> , 2013, 28, 2097-2105.	1.7	172
38	Renalâ€hepaticâ€pancreatic dysplasia: A sibship with skeletal and central nervous system anomalies and <i>NPHP3</i> mutation. <i>American Journal of Medical Genetics, Part A</i> , 2013, 161, 1743-1749.	1.2	7
39	Eculizumab treatment of atypical hemolytic uremic syndrome. <i>Expert Opinion on Orphan Drugs</i> , 2013, 1, 167-176.	0.8	1
40	Update on <i>Streptococcus pneumoniae</i> associated hemolytic uremic syndrome. <i>Current Opinion in Pediatrics</i> , 2013, 25, 203-208.	2.0	100
41	Multiple residues in the distal C terminus of the Î±-subunit have roles in modulating human epithelial sodium channel activity. <i>American Journal of Physiology - Renal Physiology</i> , 2012, 303, F220-F228.	2.7	9
42	A time for reappraisal of â€œatypicalâ€ hemolytic uremic syndrome: should all patients be treated the same?. <i>European Journal of Pediatrics</i> , 2012, 171, 1519-1525.	2.7	15
43	Developmental Abnormalities of the Kidneys. , 2012, , 1182-1190.		1
44	Urolithiasis in Children. <i>Pediatric Clinics of North America</i> , 2012, 59, 881-896.	1.8	105
45	Glomerulonephropathies and Disorders of Tubular Function. , 2012, , 1222-1227.		1
46	<i>Streptococcus pneumoniae</i> -associated Hemolytic Uremic Syndrome Among Children in North America. <i>Pediatric Infectious Disease Journal</i> , 2011, 30, 736-739.	2.0	72
47	Insights from the Chronic Kidney Disease in Children (CKiD) Study. <i>Clinical Journal of the American Society of Nephrology: CJASN</i> , 2011, 6, 2047-2053.	4.5	55
48	Multiple Residues in the Distal Câ€terminus of the Alpha Subunit Have Roles in Modulating ENaC Activity. <i>FASEB Journal</i> , 2011, 25, 1041.15.	0.5	0
49	Childhood Nephrotic Syndrome in Cambodia: An Association with Gastrointestinal Parasites. <i>Journal of Pediatrics</i> , 2010, 156, 76-81.	1.8	21
50	Evidence for a recurrent microdeletion at chromosome 16p11.2 associated with congenital anomalies of the kidney and urinary tract (CAKUT) and Hirschsprung disease. <i>American Journal of Medical Genetics, Part A</i> , 2010, 152A, 2618-2622.	1.2	49
51	<i>Streptococcus pneumoniae</i>â€Associated Hemolytic Uremic Syndrome: Classification and the Emergence of Serotype 19A. <i>Pediatrics</i> , 2010, 125, e174-e182.	2.1	87
52	An expanded syndrome of dRTA with hearing loss, hyperoxaluria and beta2-microglobulinuria. CKJ: <i>Clinical Kidney Journal</i> , 2010, 3, 439-442.	2.9	2
53	<i>Streptococcus pneumoniae</i> -associated hemolytic uremic syndrome. <i>Pediatric Nephrology</i> , 2008, 23, 1951-1956.	1.7	126
54	The thrombotic microangiopathies. <i>Pediatric Nephrology</i> , 2008, 23, 1761-1767.	1.7	63

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55	Hypothesis. Clinical Journal of the American Society of Nephrology: CJASN, 2007, 2, 914-918.	4.5	74
56	Renin-angiotensin axis blockade reduces proteinuria in presymptomatic patients with familial FSGS. Pediatric Nephrology, 2007, 22, 1779-1784.	1.7	17
57	Is genetic testing of healthy pre-symptomatic children with possible Alport syndrome ethical?. Pediatric Nephrology, 2006, 21, 455-456.	1.7	8
58	Simplified treatment strategies to fluid therapy in diarrhea. Pediatric Nephrology, 2003, 18, 1152-1156.	1.7	21
59	Eculizumab treatment of atypical hemolytic uremic syndrome. Expert Opinion on Orphan Drugs, 0, , 1-10.	0.8	0