

Onur Cil

List of Publications by Year in descending order

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papers

643
citations

623574

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times ranked

823
citing authors

#	ARTICLE	IF	CITATIONS
1	Small-molecule inhibitor of intestinal anion exchanger SLC26A3 for treatment of hyperoxaluria and nephrolithiasis. JCI Insight, 2022, 7, .	2.3	8
2	Lubiprostone is a Non-Selective Activator of cAMP-Gated Ion Channels and Chloride Channel Protein 2 (Clc-2) Has a Minor Role in its Prosecretory Effect in Intestinal Epithelial Cells. Molecular Pharmacology, 2022, 102, 106-115.	1.0	6
3	SLC26A6-selective inhibitor identified in a small-molecule screen blocks fluid absorption in small intestine. JCI Insight, 2021, 6, .	2.3	9
4	A small molecule inhibitor of the chloride channel TMEM16A blocks vascular smooth muscle contraction and lowers blood pressure in spontaneously hypertensive rats. Kidney International, 2021, 100, 311-320.	2.6	23
5	Repurposing calcium sensing receptor agonist cinacalcet for treatment of CFTR-mediated secretory diarrheas. JCI Insight, 2021, 6, .	2.3	4
6	Î±-Lipoic Acid (ALA) Improves Cystine Solubility in Cystinuria: Report of 2 Cases. Pediatrics, 2020, 145, e20192951.	1.0	10
7	4,8-Dimethylcoumarin Inhibitors of Intestinal Anion Exchanger slc26a3 (Downregulated in Adenoma) for Anti-Absorptive Therapy of Constipation. Journal of Medicinal Chemistry, 2019, 62, 8330-8337.	2.9	14
8	Slowed gastric emptying and improved oral glucose tolerance produced by a nanomolarâ€”potency inhibitor of calciumâ€”activated chloride channel TMEM16A. FASEB Journal, 2019, 33, 11247-11257.	0.2	14
9	Inhibition of CFTRâ€”mediated intestinal chloride secretion as potential therapy for bile acid diarrhea. FASEB Journal, 2019, 33, 10924-10934.	0.2	10
10	Intestinal epithelial potassium channels and CFTR chloride channels activated in ErbB tyrosine kinase inhibitor diarrhea. JCI Insight, 2019, 4, .	2.3	34
11	Nanomolar-Potency 1,2,4-Triazoloquinoline Inhibitors of the Kidney Urea Transporter UT-A1. Journal of Medicinal Chemistry, 2018, 61, 3209-3217.	2.9	18
12	SLC26A3 inhibitor identified in small molecule screen blocks colonic fluid absorption and reduces constipation. JCI Insight, 2018, 3, .	2.3	36
13	Monogenic Causes of Proteinuria in Children. Frontiers in Medicine, 2018, 5, 55.	1.2	17
14	High-Potency Phenylquinoxalinone Cystic Fibrosis Transmembrane Conductance Regulator (CFTR) Activators. Journal of Medicinal Chemistry, 2017, 60, 2401-2410.	2.9	27
15	Substituted 2-Acylaminocycloalkylthiophene-3-carboxylic Acid Arylamides as Inhibitors of the Calcium-Activated Chloride Channel Transmembrane Protein 16A (TMEM16A). Journal of Medicinal Chemistry, 2017, 60, 4626-4635.	2.9	31
16	Phenylquinoxalinone CFTR activator as potential prosecretory therapy for constipation. Translational Research, 2017, 182, 14-26.e4.	2.2	15
17	Benzopyrimidoâ€”pyrroloâ€”oxazineâ€”dione CFTR inhibitor (R)â€”BPOâ€”27 for antisecretory therapy of diarrheas caused by bacterial enterotoxins. FASEB Journal, 2017, 31, 751-760.	0.2	43
18	Small-Molecule Inhibitors of Pendrin Potentiate the Diuretic Action of Furosemide. Journal of the American Society of Nephrology: JASN, 2016, 27, 3706-3714.	3.0	37

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19	Su1580 A Small-Molecule CFTR Activator Increases Intestinal Fluid Secretion and Normalizes Stool Output in a Mouse Model of Constipation. <i>Gastroenterology</i> , 2016, 150, S532.	0.6	0
20	CFTR Activator Increases Intestinal Fluid Secretion and Normalizes Stool Output in a Mouse Model of Constipation. <i>Cellular and Molecular Gastroenterology and Hepatology</i> , 2016, 2, 317-327.	2.3	60
21	A Turkish BCS1L mutation causes GRACILE-like disorder. <i>Turkish Journal of Pediatrics</i> , 2016, 58, 658-661.	0.3	5
22	Salt-sparing diuretic action of a water-soluble urea analog inhibitor of urea transporters UT-A and UT-B in rats. <i>Kidney International</i> , 2015, 88, 311-320.	2.6	19
23	Genetic abnormalities and prognosis in patients with congenital and infantile nephrotic syndrome. <i>Pediatric Nephrology</i> , 2015, 30, 1279-1287.	0.9	29
24	<i>MCP1</i>2518 A/G polymorphism affects progression of childhood focal segmental glomerulosclerosis. <i>Renal Failure</i> , 2015, 37, 1435-1439.	0.8	9
25	A novel mutation in protein C gene (<i>PROC</i>) causing severe phenotype in neonatal period. <i>Pediatric Blood and Cancer</i> , 2014, 61, 763-764.	0.8	6
26	Diuresis and reduced urinary osmolality in rats produced by smallâ€molecule UTâ€Aâ€selective urea transport inhibitors. <i>FASEB Journal</i> , 2014, 28, 3878-3890.	0.2	18
27	Role of CXCR1 (CKR-1) in Inflammation of Experimental Mesangioproliferative Glomerulonephritis. <i>Renal Failure</i> , 2013, 35, 380-385.	0.8	2
28	The diuretic effect of urea analog dimethylthiourea in female Wistar rats. <i>Human and Experimental Toxicology</i> , 2012, 31, 1050-1055.	1.1	9
29	Endothelial Dysfunction and Increased Responses to Renal Nerve Stimulation in Rat Kidneys during Rhabdomyolysis-Induced Acute Renal Failure: Role of Hydroxyl Radical. <i>Renal Failure</i> , 2012, 34, 211-220.	0.8	8
30	Evidence for Pathogenicity of Atypical Splice Mutations in Autosomal Dominant Polycystic Kidney Disease. <i>Clinical Journal of the American Society of Nephrology: CJASN</i> , 2009, 4, 442-449.	2.2	12
31	Family History of Renal Disease Severity Predicts the Mutated Gene in ADPKD. <i>Journal of the American Society of Nephrology: JASN</i> , 2009, 20, 1833-1838.	3.0	110