

Peter Heinz-Erian

List of Publications by Year in descending order

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Version: 2024-02-01

8
papers

265
citations

1307594

7
h-index

1588992

8
g-index

8
all docs

8
docs citations

8
times ranked

481
citing authors

#	ARTICLE	IF	CITATIONS
1	Mutations in SPINT2 Cause a Syndromic Form of Congenital Sodium Diarrhea. American Journal of Human Genetics, 2009, 84, 188-196.	6.2	110
2	Congenital secretory diarrhoea caused by activating germline mutations in <i>GUCY2C</i> . Gut, 2016, 65, 1306-1313.	12.1	74
3	SPINT2 (HAI-2) missense variants identified in congenital sodium diarrhea/tufting enteropathy affect the ability of HAI-2 to inhibit prostasin but not matriptase. Human Molecular Genetics, 2019, 28, 828-841.	2.9	25
4	AP1S1 missense mutations cause a congenital enteropathy via an epithelial barrier defect. Human Genetics, 2020, 139, 1247-1259.	3.8	24
5	Inflammatory Bowel Disease in Patients with Congenital Chloride Diarrhoea. Journal of Crohn's and Colitis, 2021, 15, 1679-1685.	1.3	14
6	Low sodium status in cystic fibrosis as assessed by calculating fractional Na + excretion is associated with decreased growth parameters. Journal of Cystic Fibrosis, 2016, 15, 400-405.	0.7	10
7	Reduced NHE3 activity results in congenital diarrhea and can predispose to inflammatory bowel disease. American Journal of Physiology - Regulatory Integrative and Comparative Physiology, 2017, 312, R311-R311.	1.8	7
8	Isolated choanal and gut atresias: pathogenetic role of serine protease inhibitor type 2 (SPINT2) gene mutations unlikely. European Journal of Medical Research, 2018, 23, 13.	2.2	1