Peter Heinz-Erian

List of Publications by Year
in descending order

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	Inflammatory Bowel Disease in Patients with Congenital Chloride Diarrhoea. Journal of Crohn's and Colitis, 2021, 15, 1679-1685.	1.3	14
2	AP1S1 missense mutations cause a congenital enteropathy via an epithelial barrier defect. Human Genetics, 2020, 139, 1247-1259.	3.8	24
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	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
5	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
6	Low sodium status in cystic fibrosisâ€"as assessed by calculating fractional Na + excretionâ€"is associated with decreased growth parameters. Journal of Cystic Fibrosis, 2016, 15, 400-405.	0.7	10
7	Congenital secretory diarrhoea caused by activating germline mutations inGUCY2C. Gut, 2016, 65, 1306-1313.	12.1	74
8	Mutations in SPINT2 Cause a Syndromic Form of Congenital Sodium Diarrhea. American Journal of Human Genetics, 2009, 84, 188-196.	6.2	110