

Michael Wilschanski

List of Publications by Year in descending order

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Version: 2024-02-01

47
papers

2,733
citations

331670

21
h-index

214800

47
g-index

48
all docs

48
docs citations

48
times ranked

2862
citing authors

#	ARTICLE	IF	CITATIONS
1	Tumor necrosis factor- α inhibitor-induced follicular psoriasiform eruption. <i>Pediatric Dermatology</i> , 2022, , .	0.9	3
2	Cystic fibrosis and the gut. <i>Frontline Gastroenterology</i> , 2021, 12, 622-628.	1.8	5
3	Ivacaftor in People with Cystic Fibrosis and a <i>3849+10kb C</i> or <i>D1152H</i> Residual Function Mutation. <i>Annals of the American Thoracic Society</i>, 2021, 18, 433-441.</i></i>	3.2	22
4	Cystic Fibrosis Related Gastrointestinal Manifestations - Moving Forward. <i>Journal of Cystic Fibrosis</i> , 2021, 20, 562-563.	0.7	3
5	International Survey on Severe Acute Respiratory Syndrome Coronavirus 2 and Acute Pancreatitis Co-occurrence in Children. <i>Pancreas</i> , 2021, 50, 1305-1309.	1.1	5
6	Nasal potential difference in suspected cystic fibrosis patients with 5T polymorphism. <i>Journal of Cystic Fibrosis</i> , 2020, 19, 627-631.	0.7	9
7	Insights into the variability of nasal potential difference, a biomarker of CFTR activity. <i>Journal of Cystic Fibrosis</i> , 2020, 19, 620-626.	0.7	14
8	Clinical and Practice Variations in Pediatric Acute Recurrent or Chronic Pancreatitis. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2020, 71, 112-118.	1.8	14
9	Pediatric chronic pancreatitis without prior acute or acute recurrent pancreatitis: A report from the INSPPIRE consortium. <i>Pancreatology</i> , 2020, 20, 781-784.	1.1	8
10	Pancreas Divisum in Pediatric Acute Recurrent and Chronic Pancreatitis. <i>Journal of Clinical Gastroenterology</i> , 2019, 53, e232-e238.	2.2	35
11	Initial Development and Validation of a Transition Readiness Scale for Adolescents with Inflammatory Bowel Disease. <i>Gastroenterology Research and Practice</i> , 2019, 2019, 1-6.	1.5	4
12	Chronic Pancreatitis. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2019, 68, 566-573.	1.8	50
13	Diabetes Mellitus in Children with Acute Recurrent and Chronic Pancreatitis. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2019, 69, 599-606.	1.8	20
14	Risk Factors for Rapid Progression From Acute Recurrent to Chronic Pancreatitis in Children. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2019, 69, 206-211.	1.8	39
15	Celiac Disease and Celiac Antibodies in DM1 Patients: When Are Screening and Biopsy Recommended?. <i>Digestive Diseases and Sciences</i> , 2019, 64, 487-492.	2.3	6
16	EPC/HPSG evidence-based guidelines for the management of pediatric pancreatitis. <i>Pancreatology</i> , 2018, 18, 146-160.	1.1	89
17	CFTR Protein Function Modulation Therapy Is Finally Targeting Cystic Fibrosis-related Gastrointestinal Disease. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2018, 66, 372-373.	1.8	7
18	Cystic fibrosis: a gastrointestinal cancer syndrome. <i>Lancet Oncology</i> , The, 2018, 19, 719-720.	10.7	9

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19	Impact of Obesity on Pediatric Acute Recurrent and Chronic Pancreatitis. <i>Pancreas</i> , 2018, 47, 967-973.	1.1	19
20	International Study Group of Pediatric Pancreatitis: In Search for a CuRE Cohort Study. <i>Pancreas</i> , 2018, 47, 1222-1228.	1.1	36
21	Primary sclerosing cholangitis is associated with abnormalities in CFTR. <i>Journal of Cystic Fibrosis</i> , 2018, 17, 666-671.	0.7	11
22	Recommendations for Diagnosis and Management of Autoimmune Pancreatitis in Childhood. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2018, 67, 232-236.	1.8	35
23	Early-Onset Acute Recurrent and Chronic Pancreatitis Is Associated with PRSS1 or CTFC Gene Mutations. <i>Journal of Pediatrics</i> , 2017, 186, 95-100.	1.8	68
24	A multiple reader scoring system for Nasal Potential Difference parameters. <i>Journal of Cystic Fibrosis</i> , 2017, 16, 573-578.	0.7	10
25	Autoimmune Pancreatitis in Children: Characteristic Features, Diagnosis, and Management. <i>American Journal of Gastroenterology</i> , 2017, 112, 1604-1611.	0.4	70
26	Highlights of the ESPEN-ESPGHAN-ECFS Guidelines on Nutrition Care for Infants and Children With Cystic Fibrosis. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2016, 63, 671-675.	1.8	9
27	International prospective study of distal intestinal obstruction syndrome in cystic fibrosis: Associated factors and outcome. <i>Journal of Cystic Fibrosis</i> , 2016, 15, 531-539.	0.7	51
28	Risk Factors Associated With Pediatric Acute Recurrent and Chronic Pancreatitis. <i>JAMA Pediatrics</i> , 2016, 170, 562.	6.2	205
29	Hepatopulmonary Syndrome in Patients With Cystic Fibrosis and Liver Disease. <i>Chest</i> , 2016, 149, e35-e38.	0.8	13
30	Treatment of cystic fibrosis in low-income countries. <i>Lancet Respiratory Medicine</i> , 2016, 4, 91-92.	10.7	11
31	ESPEN-ESPGHAN-ECFS guidelines on nutrition care for infants, children, and adults with cystic fibrosis. <i>Clinical Nutrition</i> , 2016, 35, 557-577.	5.0	367
32	CFTR: A New Horizon in the Pathomechanism and Treatment of Pancreatitis. <i>Reviews of Physiology, Biochemistry and Pharmacology</i> , 2016, 170, 37-66.	1.6	82
33	Colon Cancer Associated Transcript-1 (CCAT1) Expression in Adenocarcinoma of the Stomach. <i>Journal of Cancer</i> , 2015, 6, 105-110.	2.5	72
34	Preconditioning allows engraftment of mouse and human embryonic lung cells, enabling lung repair in mice. <i>Nature Medicine</i> , 2015, 21, 869-879.	30.7	93
35	Pediatric Chronic Pancreatitis Is Associated with Genetic Risk Factors and Substantial Disease Burden. <i>Journal of Pediatrics</i> , 2015, 166, 890-896.e1.	1.8	165
36	A Novel Familial Mutation in the PCSK1 Gene That Alters the Oxyanion Hole Residue of Proprotein Convertase 1/3 and Impairs Its Enzymatic Activity. <i>PLoS ONE</i> , 2014, 9, e108878.	2.5	19

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37	Novel therapeutic approaches for cystic fibrosis. <i>Discovery Medicine</i> , 2013, 15, 127-33.	0.5	12
38	Class 1 CF Mutations. <i>Frontiers in Pharmacology</i> , 2012, 3, 117.	3.5	15
39	New drugs for cystic fibrosis. <i>Expert Opinion on Investigational Drugs</i> , 2011, 20, 1285-1292.	4.1	9
40	Defining DIOS and Constipation in Cystic Fibrosis With a Multicentre Study on the Incidence, Characteristics, and Treatment of DIOS. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2010, 50, 38-42.	1.8	143
41	Evidence of Intestinal Inflammation in Patients With Cystic Fibrosis. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2010, 51, 304-308.	1.8	146
42	Small Molecules to Treat Cystic Fibrosis. <i>Proceedings of the American Thoracic Society</i> , 2010, 7, 399-403.	3.5	3
43	Genetic Modifiers of Liver Disease in Cystic Fibrosis. <i>JAMA - Journal of the American Medical Association</i> , 2009, 302, 1076.	7.4	256
44	Patterns of gastrointestinal disease associated with mutations of CFTR. <i>Current Gastroenterology Reports</i> , 2008, 10, 316-323.	2.5	20
45	Patterns of GI disease in adulthood associated with mutations in the CFTR gene. <i>Gut</i> , 2007, 56, 1153-1163.	12.1	140
46	A Pilot Study of the Effect of Gentamicin on Nasal Potential Difference Measurements in Cystic Fibrosis Patients Carrying Stop Mutations. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2000, 161, 860-865.	5.6	202
47	Clinical and Genetic Risk Factors for Cystic Fibrosis-related Liver Disease. <i>Pediatrics</i> , 1999, 103, 52-57.	2.1	108