

# Chee Y Ooi

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/1010377/publications.pdf>

Version: 2024-02-01

120  
papers

4,247  
citations

94433

37  
h-index

133252

59  
g-index

123  
all docs

123  
docs citations

123  
times ranked

3410  
citing authors

#	ARTICLE	IF	CITATIONS
1	Definitions of Pediatric Pancreatitis and Survey of Present Clinical Practices. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2012, 55, 261-265.	1.8	354
2	Type of CFTR Mutation Determines Risk of Pancreatitis in Patients With Cystic Fibrosis. <i>Gastroenterology</i> , 2011, 140, 153-161.	1.3	226
3	Risk Factors Associated With Pediatric Acute Recurrent and Chronic Pancreatitis. <i>JAMA Pediatrics</i> , 2016, 170, 562.	6.2	205
4	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
5	Cystic fibrosis from the gastroenterologist's perspective. <i>Nature Reviews Gastroenterology and Hepatology</i> , 2016, 13, 175-185.	17.8	112
6	Inconclusive Diagnosis of Cystic Fibrosis After Newborn Screening. <i>Pediatrics</i> , 2015, 135, e1377-e1385.	2.1	105
7	Impact of CFTR modulation with Ivacaftor on Gut Microbiota and Intestinal Inflammation. <i>Scientific Reports</i> , 2018, 8, 17834.	3.3	99
8	Cystic fibrosis transmembrane conductance regulator (CFTR) gene mutations in pancreatitis. <i>Journal of Cystic Fibrosis</i> , 2012, 11, 355-362.	0.7	94
9	Features of Severe Liver Disease With Portal Hypertension in Patients With Cystic Fibrosis. <i>Clinical Gastroenterology and Hepatology</i> , 2016, 14, 1207-1215.e3.	4.4	94
10	Intestinal Inflammation and Impact on Growth in Children With Cystic Fibrosis. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2015, 60, 521-526.	1.8	87
11	Disrupted progression of the intestinal microbiota with age in children with cystic fibrosis. <i>Scientific Reports</i> , 2016, 6, 24857.	3.3	85
12	Gut Microbiota in Children With Cystic Fibrosis: A Taxonomic and Functional Dysbiosis. <i>Scientific Reports</i> , 2019, 9, 18593.	3.3	84
13	Causal Evaluation of Acute Recurrent and Chronic Pancreatitis in Children. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2017, 64, 95-103.	1.8	73
14	Summary and recommendations from the Australasian guidelines for the management of pancreatic exocrine insufficiency. <i>Pancreatology</i> , 2016, 16, 164-180.	1.1	71
15	Serum Lipase as an Early Predictor of Severity in Pediatric Acute Pancreatitis. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2013, 56, 602-608.	1.8	70
16	Autoimmune Pancreatitis in Children: Characteristic Features, Diagnosis, and Management. <i>American Journal of Gastroenterology</i> , 2017, 112, 1604-1611.	0.4	70
17	Early-Onset Acute Recurrent and Chronic Pancreatitis Is Associated with PRSS1 or CTSC Gene Mutations. <i>Journal of Pediatrics</i> , 2017, 186, 95-100.	1.8	68
18	Updated guidance on the management of children with cystic fibrosis transmembrane conductance regulator-related metabolic syndrome/cystic fibrosis screen positive, inconclusive diagnosis (CRMS/CFSPID). <i>Journal of Cystic Fibrosis</i> , 2021, 20, 810-819.	0.7	62

#	ARTICLE	IF	CITATIONS
19	Design and Implementation of INSPPIRE. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2014, 59, 360-364.	1.8	60
20	International survey of enteral nutrition protocols used in children with Crohn's disease. <i>Journal of Digestive Diseases</i> , 2012, 13, 107-112.	1.5	58
21	Dietary intake of energy-dense, nutrient-poor and nutrient-dense food sources in children with cystic fibrosis. <i>Journal of Cystic Fibrosis</i> , 2018, 17, 804-810.	0.7	58
22	Eosinophilic esophagitis in children with celiac disease. <i>Journal of Gastroenterology and Hepatology (Australia)</i> , 2008, 23, 1144-1148.	2.8	55
23	Thiopurine metabolite monitoring in paediatric inflammatory bowel disease. <i>Alimentary Pharmacology and Therapeutics</i> , 2007, 25, 941-947.	3.7	53
24	The Enigmatic Gut in Cystic Fibrosis: Linking Inflammation, Dysbiosis, and the Increased Risk of Malignancy. <i>Current Gastroenterology Reports</i> , 2017, 19, 6.	2.5	53
25	Fecal Biomarkers of Intestinal Health and Disease in Children. <i>Frontiers in Pediatrics</i> , 2014, 2, 6.	1.9	52
26	Differences in Outcomes between Early and Late Diagnosis of Cystic Fibrosis in the Newborn Screening Era. <i>Journal of Pediatrics</i> , 2017, 181, 137-145.e1.	1.8	52
27	Early Posthepatoportoenterostomy Predictors of Native Liver Survival in Biliary Atresia. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2017, 64, 203-209.	1.8	50
28	Chronic Pancreatitis. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2019, 68, 566-573.	1.8	50
29	Direct Costs of Acute Recurrent and Chronic Pancreatitis in Children in the INSPPIRE Registry. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2016, 62, 443-449.	1.8	49
30	Update of Faecal Markers of Inflammation in Children with Cystic Fibrosis. <i>Mediators of Inflammation</i> , 2012, 2012, 1-6.	3.0	46
31	Does extensive genotyping and nasal potential difference testing clarify the diagnosis of cystic fibrosis among patients with single-organ manifestations of cystic fibrosis?. <i>Thorax</i> , 2014, 69, 254-260.	5.6	45
32	Therapeutic Endoscopic Retrograde Cholangiopancreatography in Pediatric Patients With Acute Recurrent and Chronic Pancreatitis. <i>Pancreas</i> , 2017, 46, 764-769.	1.1	45
33	Comparing the American and European diagnostic guidelines for cystic fibrosis: same disease, different language?. <i>Thorax</i> , 2012, 67, 618-624.	5.6	43
34	Age-dependent variation of fecal calprotectin in cystic fibrosis and healthy children. <i>Journal of Cystic Fibrosis</i> , 2017, 16, 631-636.	0.7	43
35	Pancreatitis and pancreatic cystosis in Cystic Fibrosis. <i>Journal of Cystic Fibrosis</i> , 2017, 16, S79-S86.	0.7	43
36	Paediatric Patients with Coeliac Disease on a Gluten-Free Diet: Nutritional Adequacy and Macro- and Micronutrient Imbalances. <i>Current Gastroenterology Reports</i> , 2018, 20, 2.	2.5	40

#	ARTICLE	IF	CITATIONS
37	Toxicâ€metabolic Risk Factors in Pediatric Pancreatitis. Journal of Pediatric Gastroenterology and Nutrition, 2016, 62, 609-617.	1.8	39
38	Risk Factors for Rapid Progression From Acute Recurrent to Chronic Pancreatitis in Children. Journal of Pediatric Gastroenterology and Nutrition, 2019, 69, 206-211.	1.8	39
39	Prevalence of meconium ileus marks the severity of mutations of the Cystic Fibrosis Transmembrane Conductance Regulator (CFTR) gene. Genetics in Medicine, 2016, 18, 333-340.	2.4	37
40	INternational Study Group of Pediatric Pancreatitis: In Search for a CuRE Cohort Study. Pancreas, 2018, 47, 1222-1228.	1.1	36
41	Recommendations for Diagnosis and Management of Autoimmune Pancreatitis in Childhood. Journal of Pediatric Gastroenterology and Nutrition, 2018, 67, 232-236.	1.8	35
42	Pancreas Divisum in Pediatric Acute Recurrent and Chronic Pancreatitis. Journal of Clinical Gastroenterology, 2019, 53, e232-e238.	2.2	35
43	Probiotics in paediatric gastrointestinal diseases. Journal of Paediatrics and Child Health, 2007, 43, 331-336.	0.8	29
44	Are Thiopurines Always Contraindicated After Thiopurineâ€Induced Pancreatitis in Inflammatory Bowel Disease?. Journal of Pediatric Gastroenterology and Nutrition, 2013, 57, 583-586.	1.8	29
45	Early and Peri-operative Prognostic Indicators in Infants Undergoing Hepatic Portoenterostomy for Biliary Atresia: a Review. Current Gastroenterology Reports, 2017, 19, 16.	2.5	29
46	Thrombotic events after pediatric liver transplantation. Pediatric Transplantation, 2010, 14, 476-482.	1.0	28
47	Gastrointestinal complications in children with acute myeloid leukemia. Leukemia and Lymphoma, 2010, 51, 768-777.	1.3	27
48	What Do We Know about the Microbiome in Cystic Fibrosis? Is There a Role for Probiotics and Prebiotics?. Nutrients, 2022, 14, 480.	4.1	27
49	Contrasts and comparisons between childhood and adult onset acute pancreatitis. Pancreatology, 2013, 13, 429-435.	1.1	26
50	Elevated fecal <math>M</math>-pyruvate kinase in children with cystic fibrosis: A clue to the increased risk of intestinal malignancy in adulthood?. Journal of Gastroenterology and Hepatology (Australia), 2015, 30, 866-871.	2.8	26
51	Peak OGTT glucose is associated with lower lung function in young children with cystic fibrosis. Journal of Cystic Fibrosis, 2020, 19, 305-309.	0.7	26
52	Genetic Testing in Pancreatitis. Gastroenterology, 2010, 138, 2202-2206.e1.	1.3	25
53	Early glucose abnormalities are associated with pulmonary inflammation in young children with cystic fibrosis. Journal of Cystic Fibrosis, 2019, 18, 869-873.	0.7	25
54	Role of Cystic Fibrosis Transmembrane Conductance Regulator in Patients With Chronic Sinopulmonary Disease. Chest, 2012, 142, 996-1004.	0.8	23

#	ARTICLE	IF	CITATIONS
55	Fecal Human Î²-Defensin 2 in Children with Cystic Fibrosis: Is There a Diminished Intestinal Innate Immune Response?. <i>Digestive Diseases and Sciences</i> , 2015, 60, 2946-2952.	2.3	23
56	Glucose abnormalities detected by continuous glucose monitoring are common in young children with Cystic Fibrosis. <i>Journal of Cystic Fibrosis</i> , 2020, 19, 700-703.	0.7	23
57	Diagnosing acute pancreatitis in children: What is the diagnostic yield and concordance for serum pancreatic enzymes and imaging within 96Âh of presentation?. <i>Pancreatology</i> , 2014, 14, 251-256.	1.1	21
58	Clinical significance of liver histology on outcomes in biliary atresia. <i>Journal of Paediatrics and Child Health</i> , 2017, 53, 252-256.	0.8	21
59	Probiotics for people with cystic fibrosis. <i>The Cochrane Library</i> , 2020, 1, CD012949.	2.8	21
60	Predicting severe acute pancreatitis in children based on serum lipase and calcium: A multicentre retrospective cohort study. <i>Pancreatology</i> , 2016, 16, 529-534.	1.1	20
61	Immunoreactive trypsinogen levels in newborn screened infants with an inconclusive diagnosis of cystic fibrosis. <i>BMC Pediatrics</i> , 2019, 19, 369.	1.7	20
62	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
63	Ursodeoxycholic acid and liver disease associated with cystic fibrosis: A multicenter cohort study. <i>Journal of Cystic Fibrosis</i> , 2022, 21, 220-226.	0.7	20
64	Intestinal Inflammation and Alterations in the Gut Microbiota in Cystic Fibrosis: A Review of the Current Evidence, Pathophysiology and Future Directions. <i>Journal of Clinical Medicine</i> , 2022, 11, 649.	2.4	20
65	Impact of Obesity on Pediatric Acute Recurrent and Chronic Pancreatitis. <i>Pancreas</i> , 2018, 47, 967-973.	1.1	19
66	High ambient temperature and risk of intestinal obstruction in cystic fibrosis. <i>Journal of Paediatrics and Child Health</i> , 2016, 52, 430-435.	0.8	18
67	Diagnosing cystic fibrosis-related diabetes: current methods and challenges. <i>Expert Review of Respiratory Medicine</i> , 2016, 10, 799-811.	2.5	18
68	Factors Associated With Frequent Opioid Use in Children With Acute Recurrent and Chronic Pancreatitis. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2020, 70, 106-114.	1.8	18
69	Web-based cognitive-behavioral intervention for pain in pediatric acute recurrent and chronic pancreatitis: Protocol of a multicenter randomized controlled trial from the study of chronic pancreatitis, diabetes and pancreatic cancer (CPDPC). <i>Contemporary Clinical Trials</i> , 2020, 88, 105898.	1.8	18
70	<i>Saccharomyces boulardii</i> in a child with recurrent <i>Clostridium difficile</i>. <i>Pediatrics International</i> , 2009, 51, 156-158.	0.5	17
71	Predicting a biliary aetiology in paediatric acute pancreatitis. <i>Archives of Disease in Childhood</i> , 2013, 98, 965-969.	1.9	17
72	Ursodeoxycholic acid in cystic fibrosis-associated liver disease. <i>Journal of Cystic Fibrosis</i> , 2012, 11, 72-73.	0.7	16

#	ARTICLE	IF	CITATIONS
73	Advancing nutritional therapy: A novel polymeric formulation attenuates intestinal inflammation in a murine colitis model and suppresses pro-inflammatory cytokine production in ex-vivo cultured inflamed colonic biopsies. <i>Clinical Nutrition</i> , 2017, 36, 497-505.	5.0	16
74	Age-related levels of fecal M2-pyruvate kinase in children with cystic fibrosis and healthy children 0 to 10 years old. <i>Journal of Cystic Fibrosis</i> , 2018, 17, 109-113.	0.7	16
75	Resuscitating Cardiopulmonary Resuscitation Training in a Virtual Reality: Prospective Interventional Study. <i>Journal of Medical Internet Research</i> , 2021, 23, e22920.	4.3	16
76	Elevated plasma dihydroorotate in Miller syndrome: Biochemical, diagnostic and clinical implications, and treatment with uridine. <i>Molecular Genetics and Metabolism</i> , 2016, 119, 83-90.	1.1	15
77	Resolution of Intestinal Histopathology Changes in Cystic Fibrosis after Treatment with Ivacaftor. <i>Annals of the American Thoracic Society</i> , 2016, 13, 297-298.	3.2	15
78	Early Feeding in Acute Pancreatitis in Children: A Randomized Controlled Trial. <i>Pediatrics</i> , 2020, 146, .	2.1	15
79	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
80	Cystic Fibrosis-Related Liver Disease is Associated With Increased Disease Burden and Endocrine Comorbidities. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2020, 70, 796-800.	1.8	14
81	Diagnosis and treatment of exocrine pancreatic insufficiency in chronic pancreatitis: An international expert survey and case vignette study. <i>Pancreatology</i> , 2022, 22, 457-465.	1.1	14
82	Differences in clinical outcomes of paediatric cystic fibrosis patients with and without meconium ileus. <i>Journal of Cystic Fibrosis</i> , 2019, 18, 857-862.	0.7	13
83	The Exocrine Pancreas in Cystic Fibrosis in the Era of CFTR Modulation: A Mini Review. <i>Frontiers in Pediatrics</i> , 0, 10, .	1.9	13
84	Does Integration of Various Ion Channel Measurements Improve Diagnostic Performance in Cystic Fibrosis?. <i>Annals of the American Thoracic Society</i> , 2014, 11, 562-570.	3.2	12
85	The role, yield and cost of paediatric faecal elastase-1 testing. <i>Pancreatology</i> , 2016, 16, 551-554.	1.1	11
86	Is there a role for stool metabolomics in cystic fibrosis?. <i>Pediatrics International</i> , 2016, 58, 808-811.	0.5	11
87	Practical approach to the gastrointestinal manifestations of cystic fibrosis. <i>Journal of Paediatrics and Child Health</i> , 2018, 54, 609-619.	0.8	11
88	The intestinal virome in children with cystic fibrosis differs from healthy controls. <i>PLoS ONE</i> , 2020, 15, e0233557.	2.5	11
89	Micronutrient intake in children with cystic fibrosis in Sydney, Australia. <i>Journal of Cystic Fibrosis</i> , 2020, 19, 146-152.	0.7	10
90	Using HbA1c as a screening tool for Cystic Fibrosis Related Diabetes. <i>Journal of Cystic Fibrosis</i> , 2016, 15, 263-264.	0.7	9

#	ARTICLE	IF	CITATIONS
91	What can the gut microbiome teach us about the connections between child physical and mental health? A systematic review. <i>Developmental Psychobiology</i> , 2019, 61, 700-713.	1.6	9
92	Liver Transplantation for Massive Hepatic Lymphangiomas in a Child. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2011, 52, 366-369.	1.8	8
93	Pediatric chronic pancreatitis without prior acute or acute recurrent pancreatitis: A report from the INSPIRE consortium. <i>Pancreatology</i> , 2020, 20, 781-784.	1.1	8
94	Molecular Dynamics and Therotyping in Airway and Gut Organoids Reveal R352Q-CFTR Conductance Defect. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2022, 67, 99-111.	2.9	8
95	Molecular dynamics and functional characterization of I37R-CFTR lasso mutation provide insights into channel gating activity. <i>IScience</i> , 2022, 25, 103710.	4.1	6
96	Colonic Atresia Presenting as Neonatal Bowel Obstruction in Cystic Fibrosis. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2014, 58, e37-8.	1.8	5
97	Fecal calprotectin concentrations in young children with cystic fibrosis: Authors response. <i>Journal of Cystic Fibrosis</i> , 2018, 17, e10-e11.	0.7	5
98	Demographics and risk factors for pediatric recurrent acute pancreatitis. <i>Current Opinion in Gastroenterology</i> , 2021, 37, 491-497.	2.3	5
99	Vascular Complications in Pediatric Pancreatitis: A Case Series. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2021, 73, e94-e97.	1.8	5
100	Paediatric clinical exposure for medical students: Are they seeing enough?. <i>Journal of Paediatrics and Child Health</i> , 2016, 52, 1086-1089.	0.8	4
101	Influence of Dietitians in Preventing Parenteral Nutrition Prescription Errors in Children. <i>Journal of Parenteral and Enteral Nutrition</i> , 2018, 42, 607-612.	2.6	4
102	Enteritis with pneumatosis intestinalis following rotavirus immunisation in an infant with short bowel syndrome. <i>BMJ Case Reports</i> , 2017, 2017, bcr-2017-219482.	0.5	4
103	Evaluating the Alimentary and Respiratory Tracts in Health and disease (EARTH) research programme: a protocol for prospective, longitudinal, controlled, observational studies in children with chronic disease at an Australian tertiary paediatric hospital. <i>BMJ Open</i> , 2020, 10, e033916.	1.9	4
104	Are children on jejunal feeds at risk of iron deficiency?. <i>World Journal of Gastroenterology</i> , 2015, 21, 5751.	3.3	4
105	Probiotics for people with cystic fibrosis. <i>The Cochrane Library</i> , 0, , .	2.8	3
106	Paediatric pancreatic diseases. <i>Journal of Paediatrics and Child Health</i> , 2020, 56, 1694-1701.	0.8	3
107	A systematic cochrane review of probiotics for people with cystic fibrosis. <i>Paediatric Respiratory Reviews</i> , 2020, 39, 61-64.	1.8	3
108	Health-Related Quality of Life in Pediatric Acute Recurrent or Chronic Pancreatitis. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2022, 74, 636-642.	1.8	3

#	ARTICLE	IF	CITATIONS
109	Children With Cystic Fibrosis Have Elevated Levels of Fecal Chitinase-3-like-1. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2022, 75, 48-51.	1.8	3
110	Intestinal dysbiosis and inflammation in cystic fibrosis impacts gut and multi-organ axes. <i>Medicine in Microecology</i> , 2022, 13, 100057.	1.6	3
111	Case report: Cholecystoduodenostomy for cholestatic liver disease in a premature infant with cystic fibrosis and short gut syndrome. <i>BMC Pediatrics</i> , 2019, 19, 78.	1.7	2
112	Coeliac disease in Chinese children. <i>Journal of Paediatrics and Child Health</i> , 2015, 51, 566-566.	0.8	1
113	An unusual case of haemolytic anaemia and failure to thrive in a Burmese refugee baby. <i>Journal of Paediatrics and Child Health</i> , 2017, 53, 500-502.	0.8	1
114	Authors' Response. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2014, 58, e42.	1.8	0
115	Author's Response: Re: Stratifying Cystic Fibrosis Risk for Newborn Screen Infants With Equivocal Sweat Chloride Levels. <i>Pediatrics</i> , 2015, 136, e1490-e1491.	2.1	0
116	Ode to the exocrine pancreas. <i>Journal of Cystic Fibrosis</i> , 2016, 15, 557-558.	0.7	0
117	Differences in Outcomes Between Early and Late Diagnosis of Cystic Fibrosis in the Newborn Screening Era. <i>Obstetrical and Gynecological Survey</i> , 2017, 72, 328-330.	0.4	0
118	Hepatobiliary and Pancreatic: Hepatic arterioportal fistula: A novel and treatable feature of Alagille syndrome. <i>Journal of Gastroenterology and Hepatology (Australia)</i> , 2019, 34, 633-633.	2.8	0
119	Cystic Fibrosis—Pancreas and Intestine. , 2020, , 780-789.		0
120	Pancreatic Disease, <i>Pediatric</i> . , 2020, , 39-54.		0