Isabelle Sermet-Gaudelus

List of Publications by Year
in descending order

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Version: 2024-02-01

37 papers	2,138 citations	331670 21 h-index	345221 36 g-index
38 all docs	38 docs citations	38 times ranked	2697 citing authors

#	Article	IF	Citations
1	Severe COVIDâ€19 evolving towards organizing pneumonia in a pediatric lung transplant recipient. Pediatric Pulmonology, 2022, 57, 583-585.	2.0	2
2	Correlating genotype with phenotype using CFTRâ€mediated wholeâ€cell Cl^{â”} currents in human nasal epithelial cells. Journal of Physiology, 2022, 600, 1515-1531.	2.9	14
3	Lumacaftor-ivacaftor effects on cystic fibrosis-related liver involvement in adolescents with homozygous F508 del-CFTR. Journal of Cystic Fibrosis, 2022, 21, 212-219.	0.7	19
4	A Phase 3, open-label, 96-week trial to study the safety, tolerability, and efficacy of tezacaftor/ivacaftor in children â‰¥Â6 years of age homozygous for F508del or heterozygous for F508del and a residual function CFTR variant. Journal of Cystic Fibrosis, 2022, 21, 675-683.	0.7	10
5	Reclassifying inconclusive diagnosis after newborn screening for cystic fibrosis. Moving forward. Journal of Cystic Fibrosis, 2022, 21, 448-455.	0.7	9
6	MO1039: 1-Year Follow-Up Data of Arterial Abnormalities Identified in Kidneys Transplanted into Children During the First Covid-19 Pandemic Wave. Nephrology Dialysis Transplantation, 2022, 37, .	0.7	0
7	Clinical response to lumacaftor-ivacaftor in patients with cystic fibrosis according to baseline lung function. Journal of Cystic Fibrosis, 2021, 20, 220-227.	0.7	24
8	Exon identity influences splicing induced by exonic variants and in silico prediction efficacy. Journal of Cystic Fibrosis, 2021, 20, 464-472.	0.7	5
9	Arterial abnormalities identified in kidneys transplanted into children during the COVID-19 pandemic. American Journal of Transplantation, 2021, 21, 1937-1943.	4.7	3
10	Modulators of CFTR. Updates on clinical development and future directions. European Journal of Medicinal Chemistry, 2021, 213, 113195.	5.5	39
11	Sweat Chloride Testing and Nasal Potential Difference (NPD) Are Primary Outcome Parameters in Treatment with Cystic Fibrosis Transmembrane Conductance Regulator (CFTR) Modulators. Journal of Personalized Medicine, 2021, 11, 729.	2.5	12
12	Insights Into Patient Variability During Ivacaftor-Lumacaftor Therapy in Cystic Fibrosis. Frontiers in Pharmacology, 2021, 12, 577263.	3.5	6
13	Antisense oligonucleotide-based drug development for Cystic Fibrosis patients carrying the 3849+10Âkb C-to-T splicing mutation. Journal of Cystic Fibrosis, 2021, 20, 865-875.	0.7	30
14	Pharmacological chaperones improve intra-domain stability and inter-domain assembly via distinct binding sites to rescue misfolded CFTR. Cellular and Molecular Life Sciences, 2021, 78, 7813-7829.	5.4	36
15	Real-Life Safety and Effectiveness of Lumacaftorâ€“Ivacaftor in Patients with Cystic Fibrosis. American Journal of Respiratory and Critical Care Medicine, 2020, 201, 188-197.	5.6	95
16	Insights into the variability of nasal potential difference, a biomarker of CFTR activity. Journal of Cystic Fibrosis, 2020, 19, 620-626.	0.7	14
17	CFTR: New insights into structure and function and implications for modulation by small molecules. Journal of Cystic Fibrosis, 2020, 19, S19-S24.	0.7	16
18	Impact of COVID-19 on people with cystic fibrosis. Lancet Respiratory Medicine,the, 2020, 8, e35-e36.	10.7	114

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19	Airway surface liquid acidification initiates host defense abnormalities in Cystic Fibrosis. Scientific Reports, 2019, 9, 6516.	3.3	61
20	Predictive factors for lumacaftor/ivacaftor clinical response. Journal of Cystic Fibrosis, 2019, 18, 368-374.	0.7	56
21	Might Brushed Nasal Cells Be a Surrogate for CFTR Modulator Clinical Response?. American Journal of Respiratory and Critical Care Medicine, 2019, 199, 123-126.	5.6	38
22	Cisvariants identified in F508del complex alleles modulate CFTR channel rescue by small molecules. Human Mutation, 2018, 39, 506-514.	2.5	28
23	Structure-guided combination therapy to potently improve the function of mutant CFTRs. Nature Medicine, 2018, 24, 1732-1742.	30.7	117
24	Comparative proteomics of respiratory exosomes in cystic fibrosis, primary ciliary dyskinesia and asthma. Journal of Proteomics, 2018, 185, 1-7.	2.4	38
25	Diagnosis of Cystic Fibrosis: Consensus Guidelines from the Cystic Fibrosis Foundation. Journal of Pediatrics, 2017, 181, S4-S15.e1.	1.8	572
26	A multiple reader scoring system for Nasal Potential Difference parameters. Journal of Cystic Fibrosis, 2017, 16, 573-578.	0.7	10
27	Correction of CFTR function in nasal epithelial cells from cystic fibrosis patients predicts improvement of respiratory function by CFTR modulators. Scientific Reports, 2017, 7, 7375.	3.3	134
28	Changes of CFTR functional measurements and clinical improvements in cystic fibrosis patients with non p.Gly551Asp gating mutations treated with ivacaftor. Journal of Cystic Fibrosis, 2017, 16, 45-48.	0.7	12
29	Vaccine coverage in CF children: A French multicenter study. Journal of Cystic Fibrosis, 2015, 14, 615-620.	0.7	12
30	Persistent Bordetella bronchiseptica infection in a child with cystic fibrosis: Relationship to bacterial phenotype. Journal of Cystic Fibrosis, 2015, 14, E13-E15.	0.7	13
31	Biosynthesis of cystic fibrosis transmembrane conductance regulator. International Journal of Biochemistry and Cell Biology, 2014, 52, 26-38.	2.8	46
32	Ataluren for the treatment of nonsense-mutation cystic fibrosis: a randomised, double-blind, placebo-controlled phase 3 trial. Lancet Respiratory Medicine,the, 2014, 2, 539-547.	10.7	301
33	An International Randomized Multicenter Comparison of Nasal Potential Difference Techniques. Chest, 2010, 138, 919-928.	0.8	50
34	Clinical Phenotype and Genotype of Children with Borderline Sweat Test and Abnormal Nasal Epithelial Chloride Transport. American Journal of Respiratory and Critical Care Medicine, 2010, 182, 929-936.	5.6	57
35	Measurement of nasal potential difference in young children with an equivocal sweat test following newborn screening for cystic fibrosis. Thorax, 2010, 65, 539-544.	5.6	41
36	Chloride Transport in Nasal Ciliated Cells of Cystic Fibrosis Heterozygotes. American Journal of Respiratory and Critical Care Medicine, 2005, 171, 1026-1031.	5.6	36

#	Article	IF	Citations
37	Basic protocol for transepithelial nasal potential difference measurements. Journal of Cystic Fibrosis, 2004, 3, 151-155.	0.7	67