Dominique Hubert

List of Publications by Year
in descending order

18	741	623574 14	839398 18
papers	citations	h-index	g-index

19	19	19	802
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Clinical response to lumacaftor-ivacaftor in patients with cystic fibrosis according to baseline lung function. Journal of Cystic Fibrosis, 2021, 20, 220-227.	0.3	24
2	Rapid Improvement after Starting Elexacaftorâ€“Tezacaftorâ€“Ivacaftor in Patients with Cystic Fibrosis and Advanced Pulmonary Disease. American Journal of Respiratory and Critical Care Medicine, 2021, 204, 64-73.	2.5	139
3	Real-World Long-Term Ivacaftor for Cystic Fibrosis in France: Clinical Effectiveness and Healthcare Resource Utilization. Pulmonary Therapy, 2021, 7, 455-468.	1.1	8
4	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.	2.5	95
5	Pregnancy after lung and heartâ€“lung transplantation: a French multicentre retrospective study of 39 pregnancies. ERJ Open Research, 2019, 5, 00254-2018.	1.1	17
6	Repaglinide versus insulin for newly diagnosed diabetes in patients with cystic fibrosis: a multicentre, open-label, randomised trial. Lancet Diabetes and Endocrinology,the, 2018, 6, 114-121.	5.5	53
7	Retrospective observational study of French patients with cystic fibrosis and a Gly551Asp-CFTR mutation after 1 and 2 years of treatment with ivacaftor in a real-world setting. Journal of Cystic Fibrosis, 2018, 17, 89-95.	0.3	51
8	A prospective analysis of unplanned patient-initiated contacts in an adult cystic fibrosis centre. Journal of Cystic Fibrosis, 2018, 17, 636-642.	0.3	4
9	Real-life initiation of lumacaftor/ivacaftor combination in adults with cystic fibrosis homozygous for the Phe508del CFTR mutation and severe lung disease. Journal of Cystic Fibrosis, 2017, 16, 388-391.	0.3	81
10	Case series of omalizumab for allergic bronchopulmonary aspergillosis in cystic fibrosis patients. Pediatric Pulmonology, 2017, 52, 190-197.	1.0	33
11	Optimized approach for the identification of highly efficient correctors of nonsense mutations in human diseases. PLoS ONE, 2017, 12, e0187930.	1.1	21
12	Aspergillus fumigatus in the cystic fibrosis lung: pros and cons of azole therapy. Infection and Drug Resistance, 2016, Volume 9, 229-238.	1.1	53
13	Long-term computed tomographic changes in cystic fibrosis patients treated with ivacaftor. European Respiratory Journal, 2016, 48, 249-252.	3.1	30
14	CFTR and/or pancreatitis susceptibility genes mutations as risk factors of pancreatitis in cystic fibrosis patients?. Pancreatology, 2016, 16, 515-522.	0.5	6
15	Causes of death in French cystic fibrosis patients: The need for improvement in transplantation referral strategies!. Journal of Cystic Fibrosis, 2016, 15, 204-212.	0.3	76
16	Maternal and perinatal outcomes of pregnancies in women with cystic fibrosis â€“ A single centre case-control study. Respiratory Medicine, 2016, 113, 22-27.	1.3	22
17	Discussion on genotype and phenotype correlations in patients with cystic fibrosis and pancreatitis. Gastroenterology, 2003, 125, 1286.	0.6	13
18	Management of assisted reproductive technologies in women with cystic fibrosis. Fertility and Sterility, 2001, 76, 1280-1281.	0.5	15