Ifat Sarouk

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/7270506/publications.pdf

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23 papers	378 citations	1039880 9 h-index	794469 19 g-index
24 all docs	24 docs citations	24 times ranked	704 citing authors

#	Article	IF	Citations
1	Correction of CFTR function in intestinal organoids to guide treatment of cystic fibrosis. European Respiratory Journal, 2021, 57, 1902426.	3.1	71
2	Endocrine abnormalities in ataxia telangiectasia: findings from a national cohort. Pediatric Research, 2016, 79, 889-894.	1.1	63
3	Failure to conceive in women with CF is associated with pancreatic insufficiency and advancing age. Journal of Cystic Fibrosis, 2019, 18, 525-529.	0.3	43
4	Glucose intolerance in cystic fibrosis as a determinant of pulmonary function and clinical status. Diabetes Research and Clinical Practice, 2015, 110, 276-284.	1.1	29
5	The risk for developing cancer in Israeli ATM, BLM, and FANCC heterozygous mutation carriers. Cancer Genetics, 2016, 209, 70-74.	0.2	29
6	Omalizumab in allergic bronchopulmonary aspergillosis in patients with cystic fibrosis. Journal of Asthma and Allergy, 2018, Volume 11, 101-107.	1.5	24
7	Long-term follow-up of distal intestinal obstruction syndrome in cystic fibrosis. World Journal of Gastroenterology, 2015, 21, 318.	1.4	24
8	Ivacaftor for the p.Ser549Arg (S549R) gating mutation â€“ The Israeli experience. Respiratory Medicine, 2017, 131, 225-228.	1.3	19
9	Elevated IgM levels as a marker for a unique phenotype in patients with Ataxia telangiectasia. BMC Pediatrics, 2018, 18, 185.	0.7	15
10	Cough Characteristics and FVC Maneuver in Cystic Fibrosis. Respiratory Care, 2014, 59, 1912-1917.	0.8	10
11	Several siblings with Cystic Fibrosis as a risk factor for poor outcome. Respiratory Medicine, 2015, 109, 74-78.	1.3	7
12	The Short-Term Effect of Breathing Tasks Via an Incentive Spirometer on Lung Function Compared With Autogenic Drainage in Subjects With Cystic Fibrosis. Respiratory Care, 2015, 60, 1819-1825.	0.8	7
13	Tracheal diverticula in cystic fibrosisâ€”A potentially important underreported finding on chest CT. Journal of Cystic Fibrosis, 2016, 15, 503-509.	0.3	7
14	FVC deterioration, airway obstruction determination, and life span in Ataxia telangiectasia. Respiratory Medicine, 2015, 109, 890-896.	1.3	6
15	Nocardia Colonization: A Risk Factor for Lung Deterioration in Cystic Fibrosis Patients?. Medical Science Monitor, 2015, 21, 1889-1894.	0.5	6
16	Long-Term Outcomes of Early Enzyme Replacement Therapy for Mucopolysaccharidosis IV: Clinical Case Studies of Two Siblings. Diagnostics, 2020, 10, 108.	1.3	5
17	Functional parameter measurements in children with ataxia telangiectasia. Developmental Medicine and Child Neurology, 2020, 62, 207-213.	1.1	4
18	The Value of Measuring Inspiratory Capacity in Subjects With Cystic Fibrosis. Respiratory Care, 2018, 63, 981-987.	0.8	3

#	Article	IF	Citations
19	Cough ability measurements and recurrent respiratory symptoms in individuals with Ataxia Telangiectasia. Journal of Asthma, 2016, 53, 37-42.	0.9	2
20	Advanced Lung Disease in Patients with Cystic Fibrosis Is Associated with Low Diffusion capacity. Israel Medical Association Journal, 2020, 22, 770-774.	0.1	2
21	Secondary enuresis and urological manifestations in children with ataxia telangiectasia. European Journal of Paediatric Neurology, 2018, 22, 1118-1123.	0.7	1
22	Phenotypic and molecular characteristics of CF patients carrying the I1234V mutation. Respiratory Medicine, 2020, 170, 106027.	1.3	1
23	Spontaneous pneumothoraxâ€"When do we need to intervene?. Clinical Respiratory Journal, 2021, 15, 967-972.	0.6	0