Cordula Koerner-Rettberg

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/7634238/publications.pdf

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27	992	516710 16	526287 27
papers	citations	h-index	g-index

28	28	28	1312
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Lung function from school age to adulthood in primary ciliary dyskinesia. European Respiratory Journal, 2022, 60, 2101918.	6.7	17
2	Comparison of the Lung Clearance Index in Preschool Children With Primary Ciliary Dyskinesia and Cystic Fibrosis. Chest, 2022, 162, 534-542.	0.8	11
3	Effects of a Long-Term Monitored Exercise Program on Aerobic Fitness in a Small Group of Children with Cystic Fibrosis. International Journal of Environmental Research and Public Health, 2022, 19, 7923.	2.6	3
4	Association between habitual physical activity (HPA) and sleep quality in patients with cystic fibrosis. Sleep and Breathing, 2021, 25, 609-615.	1.7	8
5	Lung clearance index predicts pulmonary exacerbations in individuals with primary ciliary dyskinesia: a multicentre cohort study. Thorax, 2021, 76, 681-688.	5.6	12
6	Trainability of Health-Related and Motor Performance Fitness in Adults with Cystic Fibrosis within a 12-Month Partially Supervised Exercise Program. Pulmonary Medicine, 2021, 2021, 1-9.	1.9	6
7	Mutations in TP73 cause impaired mucociliary clearance and lissencephaly. American Journal of Human Genetics, 2021, 108, 1318-1329.	6.2	15
8	Health-Related and Motor Performance-Related Fitness and Physical Activity Among Youth With Cystic Fibrosis. Perceptual and Motor Skills, 2021, 128, 2097-2116.	1.3	4
9	Standardised clinical data from patients with primary ciliary dyskinesia: FOLLOW-PCD. ERJ Open Research, 2020, 6, 00237-2019.	2.6	36
10	Effects of a longâ€term exercise program on motor performance in children and adolescents with CF. Pediatric Pulmonology, 2020, 55, 3371-3380.	2.0	8
11	Time trends in diagnostic testing for primary ciliary dyskinesia in Europe. European Respiratory Journal, 2019, 54, 1900528.	6.7	17
12	Prevalence and course of disease after lung resection in primary ciliary dyskinesia: a cohort & nested case-control study. Respiratory Research, 2019, 20, 212.	3.6	23
13	Pulmonary exacerbations in patients with primary ciliary dyskinesia: an expert consensus definition for use in clinical trials. ERJ Open Research, 2019, 5, 00147-2018.	2.6	37
14	Preventive Inhalation of Hypertonic Saline in Infants with Cystic Fibrosis (PRESIS). A Randomized, Double-Blind, Controlled Study. American Journal of Respiratory and Critical Care Medicine, 2019, 199, 1238-1248.	5.6	96
15	25-Hydroxvitamin D concentrations are not lower in children with bronchial asthma, atopic dermatitis, obesity, or attention-deficient/hyperactivity disorder than in healthy children. Nutrition Research, 2018, 52, 39-47.	2.9	23
16	Structural and Functional Lung Impairment in Primary Ciliary Dyskinesia. Assessment with Magnetic Resonance Imaging and Multiple Breath Washout in Comparison to Spirometry. Annals of the American Thoracic Society, 2018, 15, 1434-1442.	3.2	36
17	Lung function in patients with primary ciliary dyskinesia: an iPCD Cohort study. European Respiratory Journal, 2018, 52, 1801040.	6.7	71
18	Comparison of different analysis algorithms to calculate multiple-breath washout outcomes. ERJ Open Research, 2018, 4, 00021-2017.	2.6	3

#	Article	IF	Citations
19	The international primary ciliary dyskinesia cohort (iPCD Cohort): methods and first results. European Respiratory Journal, 2017, 49, 1601181.	6.7	77
20	Alternative inert gas washout outcomes in patients with primary ciliary dyskinesia. European Respiratory Journal, 2017, 49, 1600466.	6.7	21
21	Growth and nutritional status, and their association with lung function: a study from the international Primary Ciliary Dyskinesia Cohort. European Respiratory Journal, 2017, 50, 1701659.	6.7	50
22	Physiological phenotyping of pediatric chronic obstructive airway diseases. Journal of Applied Physiology, 2016, 121, 324-332.	2.5	20
23	Factors Associated with Worse Lung Function in Cystic Fibrosis Patients with Persistent Staphylococcus aureus. PLoS ONE, 2016, 11, e0166220.	2.5	70
24	Further evidence for an association between LCI and FEV1in patients with PCD: FigureÂ1. Thorax, 2015, 70, 896.1-896.	5.6	25
25	High Variability in Oral Glucose Tolerance among 1,128 Patients with Cystic Fibrosis: A Multicenter Screening Study. PLoS ONE, 2014, 9, e112578.	2.5	49
26	Colistimethate sodium for the treatment of chronic pulmonary infection in cystic fibrosis: an evidence-based review of its place in therapy. Core Evidence, 2014, 9, 99.	4.7	22
27	Mutations in CCNO result in congenital mucociliary clearance disorder with reduced generation of multiple motile cilia. Nature Genetics, 2014, 46, 646-651.	21.4	232