

# Ajay S Kasi

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/7389011/publications.pdf>

Version: 2024-02-01

22  
papers

187  
citations

1162367

8  
h-index

1199166

12  
g-index

22  
all docs

22  
docs citations

22  
times ranked

144  
citing authors

#	ARTICLE	IF	CITATIONS
1	Congenital central hypoventilation syndrome: Diagnostic and management challenges. <i>Pediatric Health, Medicine and Therapeutics</i> , 2016, Volume 7, 99-107.	0.7	24
2	Adult With <i>PHOX2B</i> Mutation and Late-Onset Congenital Central Hypoventilation Syndrome. <i>Journal of Clinical Sleep Medicine</i> , 2018, 14, 2079-2081.	1.4	23
3	Three-Generation Family With Congenital Central Hypoventilation Syndrome and Novel <i>PHOX2B</i> Gene Non-Polyalanine Repeat Mutation. <i>Journal of Clinical Sleep Medicine</i> , 2017, 13, 925-927.	1.4	22
4	Congenital Central Hypoventilation Syndrome: Optimizing Care with a Multidisciplinary Approach. <i>Journal of Multidisciplinary Healthcare</i> , 2022, Volume 15, 455-469.	1.1	17
5	Mortality and Outcomes of Pediatric Tracheostomy Dependent Patients. <i>Frontiers in Pediatrics</i> , 2021, 9, 661512.	0.9	15
6	Refining the Application of Microbial Lipids as Tracers of <i>Staphylococcus aureus</i> Growth Rates in Cystic Fibrosis Sputum. <i>Journal of Bacteriology</i> , 2018, 200, .	1.0	13
7	Abnormal Lung Clearance Index in Cystic Fibrosis Screen Positive, Inconclusive Diagnosis (CFSPID) Children with Otherwise Normal FEV1. <i>Lung</i> , 2020, 198, 163-167.	1.4	11
8	Bronchoscopic interventions for plastic bronchitis in children without structural heart disease. <i>European Journal of Pediatrics</i> , 2021, 180, 3547-3554.	1.3	8
9	Variable phenotypes in congenital central hypoventilation syndrome with <i>PHOX2B</i> nonpolyalanine repeat mutations. <i>Journal of Clinical Sleep Medicine</i> , 2021, 17, 2049-2055.	1.4	8
10	Cough. <i>Pediatrics in Review</i> , 2019, 40, 157-167.	0.2	7
11	Tracheostomy decannulation to noninvasive positive pressure ventilation in congenital central hypoventilation syndrome. <i>Sleep and Breathing</i> , 2022, 26, 133-139.	0.9	7
12	Abnormal Lung Clearance Index in Cystic Fibrosis Children with Normal FEV1 and Single-Breath Nitrogen Washout Test. <i>Lung</i> , 2021, 199, 37-41.	1.4	5
13	Heterogeneous Pulmonary Phenotypes in Filamin A Mutation-Related Lung Disease. <i>Pediatric, Allergy, Immunology, and Pulmonology</i> , 2021, 34, 7-14.	0.3	5
14	Recurrent apnoea and respiratory failure in an infant: congenital central hypoventilation syndrome with a novel <i>PHOX2B</i> gene variant. <i>BMJ Case Reports</i> , 2021, 14, e239633.	0.2	5
15	Annual Respiratory Evaluations in Congenital Central Hypoventilation Syndrome and Changes in Ventilatory Management. <i>Pediatric, Allergy, Immunology, and Pulmonology</i> , 2021, 34, 97-101.	0.3	4
16	Positional impairment of gas exchange during diaphragm pacing alleviated by increasing amplitude settings in congenital central hypoventilation syndrome. <i>Journal of Clinical Sleep Medicine</i> , 2020, 16, 459-462.	1.4	4
17	Impaired ventilation during 6â€min walk test in congenital central hypoventilation syndrome. <i>Pediatric Pulmonology</i> , 2022, 57, 1660-1667.	1.0	4
18	Images: Polysomnographic artifacts in a child with congenital central hypoventilation syndrome. <i>Journal of Clinical Sleep Medicine</i> , 2020, 16, 2123-2125.	1.4	2

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19	Ventilator change in children on home mechanical ventilation affected by the Philips respironics trilogy ventilator recall. <i>Pediatric Pulmonology</i> , 2022, 57, 43-48.	1.0	2
20	Can We Prevent COPD by Reaching Out to Children at Risk?. <i>Chest</i> , 2020, 158, 1327-1328.	0.4	1
21	Images: Sleep-disordered breathing and hypoventilation in a child with obesity and hypothalamic dysfunction. <i>Journal of Clinical Sleep Medicine</i> , 2022, 18, 339-342.	1.4	0
22	Hypoxaemia and interstitial lung disease in an infant with hypothyroidism and hypotonia. <i>BMJ Case Reports</i> , 2020, 13, e238466.	0.2	0