Nadia Nathan

List of Publications by Year
in descending order

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90 papers	2,824 citations	186265 28 h-index	182427 51 g-index
98 all docs	98 docs citations	98 times ranked	3819 citing authors

#	Article	IF	Citations
1	Diffuse Parenchymal Lung Disease in Early Childhood. , 2022, , 229-243.		0
2	Childhood Interstitial Lung Diseases (chILD) Recognition: When Epidemiology Increases a Rare Disease Incidence. Archivos De Bronconeumologia, 2022, , .	0.8	2
3	Paediatric sarcoidosis. , 2022, , 285-294.		0
4	French national cohort of neuroendocrine cell hyperplasia of infancy (FRENCHI) study: diagnosis and initial management. European Journal of Pediatrics, 2022, 181, 3067-3073.	2.7	2
5	COPA syndrome, 5 years after: Where are we?. Joint Bone Spine, 2021, 88, 105070.	1.6	21
6	Interstitial lung diseases. , 2021, , 588-601.		0
7	Syndrome COPA, quoi de neuf cinq ans aprÃ¨sÂ?. Revue Du Rhumatisme (Edition Francaise), 2021, 88, 183-189.	0.0	0
8	Interstitial lung diseases in the neonatal period. , 2021, , 213-230.		2
9	Severe Acute Respiratory Syndrome Coronavirus 2 Variant Delta Infects All 6 Siblings but Spares Comirnaty (BNT162b2, BioNTech/Pfizer)-Vaccinated Parents. Journal of Infectious Diseases, 2021, 224, 1984-1986.	4.0	6
10	Exposure to inorganic particles in paediatric sarcoidosis: the PEDIASARC study. Thorax, 2021, , thoraxjnl-2021-217870.	5.6	10
11	Benefits and risks of bronchoalveolar lavage in severe asthma in children. ERJ Open Research, 2021, 7, 00332-2021.	2.6	2
12	Usefulness of bronchoalveolar lavage in a French pediatric cohort with hypersensitivity pneumonitis. Pediatric Pulmonology, 2020, 55, 136-140.	2.0	5
13	Use of ruxolitinib in COPA syndrome manifesting as life-threatening alveolar haemorrhage. Thorax, 2020, 75, 92-95.	5.6	36
14	Childâ€“Adult Transition in Sarcoidosis: A Series of 52 Patients. Journal of Clinical Medicine, 2020, 9, 2097.	2.4	13
15	Mutations in COPA lead to abnormal trafficking of STING to the Golgi and interferon signaling. Journal of Experimental Medicine, 2020, 217, .	8.5	130
16	Functional assessment and phenotypic heterogeneity of SFTPA1 and SFTPA2 mutations in interstitial lung diseases and lung cancer. European Respiratory Journal, 2020, 56, 2002806.	6.7	23
17	The Wide Spectrum of COVID-19 Clinical Presentation in Children. Journal of Clinical Medicine, 2020, 9, 2950.	2.4	28
18	Dramatic improvement after tocilizumab of severe <scp>COVID</scp>â€19 in a child with sickle cell disease and acute chest syndrome. American Journal of Hematology, 2020, 95, E192-E194.	4.1	56

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19	Interstitial lung diseases in children. Presse Medicale, 2020, 49, 103909.	1.9	26
20	Atypical presentation of COVID-19 in young infants. Lancet, The, 2020, 395, 1481.	13.7	78
21	Paediatric sarcoidosis. Paediatric Respiratory Reviews, 2019, 29, 53-59.	1.8	31
22	Pulmonary Fibrosis in Children. Journal of Clinical Medicine, 2019, 8, 1312.	2.4	19
23	Healthâ€related quality of life in infants and children with interstitial lung disease. Pediatric Pulmonology, 2019, 54, 828-836.	2.0	13
24	A Nonsmoker Man in His 40s With a Diagnosis of Genetic-Related Idiopathic Pulmonary Fibrosis (Surfactant-Protein C Gene Mutation). Chest, 2019, 155, e91-e96.	0.8	4
25	Bi-allelic missense ABCA3 mutations in a patient with childhood ILD who reached adulthood. ERJ Open Research, 2019, 5, 00066-2019.	2.6	22
26	OP0107â€…HETEROZYGOUS MUTATIONS IN COPA ARE ASSOCIATED WITH ENHANCED TYPE I INTERFERON SIGNALLING. , 2019, , .		0
27	Pilot experience of multidisciplinary team discussion dedicated to inherited pulmonary fibrosis. Orphanet Journal of Rare Diseases, 2019, 14, 280.	2.7	24
28	A survey of childrenâ€™s interstitial lung disease (ChILD) databases across the EU and an ability to identify pan-registry clinical trial cohorts. , 2019, , .		0
29	Health-related quality of life in children interstitial lung disease. , 2019, , .		0
30	Genetic causes and clinical management of pediatric interstitial lung diseases. Current Opinion in Pulmonary Medicine, 2018, 24, 253-259.	2.6	30
31	Management of antiplatelet therapy in patients undergoing elective invasive procedures: Proposals from the French Working Group on perioperative hemostasis (GIHP) and the French Study Group on thrombosis and hemostasis (GFHT). In collaboration with the French Society for Anesthesia and Intensive Care (SFAR). Archives of Cardiovascular Diseases, 2018, 111, 210-223.	1.6	22
32	Whole exome sequencing in three families segregating a pediatric case of sarcoidosis. BMC Medical Genomics, 2018, 11, 23.	1.5	26
33	International management platform for childrenâ€™s interstitial lung disease (chILD-EU). Thorax, 2018, 73, 231-239.	5.6	64
34	Chronic interstitial lung diseases in children: diagnosis approaches. Expert Review of Respiratory Medicine, 2018, 12, 1051-1060.	2.5	16
35	MUC5B Promoter Variant and Rheumatoid Arthritis with Interstitial Lung Disease. New England Journal of Medicine, 2018, 379, 2209-2219.	27.0	326
36	Pulmonary hemosiderosis in children with Down syndrome: a national experience. Orphanet Journal of Rare Diseases, 2018, 13, 60.	2.7	32

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37	Contribution of mutations in genes of the surfactant system to idiopathic interstitial pneumonia (IIP). , 2018, , .		1
38	Down syndrome and pulmonary hemosiderosis: an under-recognized association. , 2018, , .		1
39	OP0284â€…Muc5b promoter variant rs35705950 is a risk factor for rheumatoid arthritis â€“ interstitial lung disease. , 2018, , .		0
40	TERT/TERC mutations in a Greek cohort of suspected genetic pulmonary fibrosis patients. , 2018, , .		0
41	COPA syndrome restricted to life-threatening alveolar hemorrhages: clinical, pathological, molecular and biological characterization. , 2018, , .		1
42	Multidisciplinary team dedicated to suspected heritable pulmonary fibrosis. , 2018, , .		1
43	Lung disease caused by non-null ABCA3 mutations: long-term follow-up. , 2018, , .		0
44	Functional assessment of newly identified SFTPA1 and SFTPA2 mutations in patients with idiopathic interstitial pneumonia (IIP) and lung cancer. , 2018, , .		0
45	Shared genetic predisposition in rheumatoid arthritis-interstitial lung disease and familial pulmonary fibrosis. European Respiratory Journal, 2017, 49, 1602314.	6.7	154
46	Management of suspected monogenic lung fibrosis in a specialised centre. European Respiratory Review, 2017, 26, 160122.	7.1	54
47	Inborn Errors of Metabolism: The Achilles' Heel of the Respiratory System. Respiration, 2017, 94, 14-15.	2.6	0
48	Heterogeneity of lung disease associated with NK2 homeobox 1 mutations. Respiratory Medicine, 2017, 129, 16-23.	2.9	54
49	AB0007â€…Shared genetic predisposition in rheumatoid arthritisâ€“interstitial lung disease and familial pulmonary fibrosis. , 2017, , .		7
50	SFTPA mutations in interstitial lung disease (ILD) and lung cancer. , 2017, , .		0
51	A genetic landscape of familial predisposition to sarcoidosis identified by whole exome sequencing. , 2017, , .		0
52	Ultrasonography and Computed Tomographic Manifestations of Abdominal Sarcoidosis in Children. Journal of Pediatric Gastroenterology and Nutrition, 2016, 63, 195-199.	1.8	3
53	Eosinophilic pneumonias in children: A review of the epidemiology, diagnosis, and treatment. Pediatric Pulmonology, 2016, 51, 203-216.	2.0	31
54	Management of children with interstitial lung diseases: the difficult issue of acute exacerbations. European Respiratory Journal, 2016, 48, 1559-1563.	6.7	33

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55	Long-term effects of azithromycin in patients with cystic fibrosis. Respiratory Medicine, 2016, 117, 1-6.	2.9	42
56	Nonsteroidal Anti-Inflammatory Drug without Antibiotics for Acute Viral Infection Increases the Empyema Risk in Children: A Matched Case-Control Study. Journal of Pediatrics, 2016, 175, 47-53.e3.	1.8	58
57	Surfactant protein A: A key player in lung homeostasis. International Journal of Biochemistry and Cell Biology, 2016, 81, 151-155.	2.8	50
58	Respiratory Distress, Congenital Hypothyroidism and Hypotonia in a Newborn. Respiration, 2016, 92, 188-191.	2.6	10
59	Familial vs. sporadic sarcoidosis: BTNL2 polymorphisms, clinical presentations, and outcomes in a French cohort. Orphanet Journal of Rare Diseases, 2016, 11, 165.	2.7	27
60	GermlineSFTPA1mutation in familial idiopathic interstitial pneumonia and lung cancer. Human Molecular Genetics, 2016, 25, 1457-1467.	2.9	119
61	GermlineSFTPA1mutation in familial idiopathic interstitial pneumonia and lung cancer. , 2016, , .		1
62	Phenotype heterogeneity in a familial â€œbrain lung thyroid syndromeâ€•related to a novelNKX-2.1mutation. , 2016, , .		0
63	Biomarkers in Interstitial lung diseases. Paediatric Respiratory Reviews, 2015, 16, 219-224.	1.8	20
64	Lung sarcoidosis in children: update on disease expression and management. Thorax, 2015, 70, 537-542.	5.6	49
65	Familial pulmonary fibrosis. Revue Des Maladies Respiratoires, 2015, 32, 413-434.	1.7	39
66	Work of breathing in children with diffuse parenchymal lung disease. Respiratory Physiology and Neurobiology, 2015, 206, 45-52.	1.6	8
67	Inorganic exposome in pediatric sarcoidosis: The PEDIASARC study. , 2015, , .		1
68	Genetic testing in idiopathic interstitial pneumonia. , 2015, , .		0
69	An idiopathic congenital chylothorax: surgery or conservative management?. BMJ Case Reports, 2014, 2014, bcr2014204147-bcr2014204147.	0.5	3
70	Idiopathic eosinophilic pneumonia in children: the French experience. Orphanet Journal of Rare Diseases, 2014, 9, 28.	2.7	19
71	Alveolar epithelial cells: Master regulators of lung homeostasis. International Journal of Biochemistry and Cell Biology, 2013, 45, 2568-2573.	2.8	187
72	New insights into pediatric idiopathic pulmonary hemosiderosis: the French RespiRareÂ® cohort. Orphanet Journal of Rare Diseases, 2013, 8, 161.	2.7	95

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73	Respiratory Outcome in Children with Scimitar Syndrome. Journal of Pediatrics, 2013, 162, 275-279.e1.	1.8	5
74	Loss-of-Function Mutations in RSPH1 Cause Primary Ciliary Dyskinesia with Central-Complex and Radial-Spoke Defects. American Journal of Human Genetics, 2013, 93, 561-570.	6.2	148
75	Endobronchial avium mycobacteria infection in an immunocompetent child. BMJ Case Reports, 2013, 2013, bcr2013200776-bcr2013200776.	0.5	5
76	Diffuse parenchymal lung disease caused by surfactant deficiency: dramatic improvement by azithromycin. BMJ Case Reports, 2013, 2013, bcr2013009988-bcr2013009988.	0.5	32
77	Pulmonary sarcoid-like granulomatous disease in an 11-month-old girl. BMJ Case Reports, 2013, 2013, bcr2012008024-bcr2012008024.	0.5	2
78	A national internet-linked based database for pediatric interstitial lung diseases: the French network. Orphanet Journal of Rare Diseases, 2012, 7, 40.	2.7	79
79	A rare CFTR intronic mutation related to a mild CF disease in a 12-year-old girl. BMJ Case Reports, 2012, 2012, bcr2012006918-bcr2012006918.	0.5	6
80	Macrolides: New therapeutic perspectives in lung diseases. International Journal of Biochemistry and Cell Biology, 2011, 43, 1241-1246.	2.8	21
81	Search For Interstitial Lung Disease Etiology In Children: A Step By Step Approach. , 2011, , .		0
82	Interstitial lung disease: Physiopathology in the context of lung growth. Paediatric Respiratory Reviews, 2011, 12, 216-222.	1.8	25
83	Azithromycin In Interstitial Lung Disease Associated With Surfactant Metabolism Disorders. , 2010, , .		0
84	Efficacy of Blebs Detection for Preventive Surgery in Childrenâ€™s Idiopathic Spontaneous Pneumothorax. World Journal of Surgery, 2010, 34, 185-189.	1.6	36
85	A National French Web Database For Children Rare Lung Diseases. , 2010, , .		0
86	Interstitial lung diseases in children. Orphanet Journal of Rare Diseases, 2010, 5, 22.	2.7	112
87	Chronic eosinophilic pneumonia in a 13-year-old child. European Journal of Pediatrics, 2008, 167, 1203-1207.	2.7	14
88	Physiologic Benefits of Mechanical Insufflation-Exsufflation in Children With Neuromuscular Diseases. Chest, 2008, 133, 161-168.	0.8	130
89	Glucocorticoid receptor gene polymorphisms associated with progression of lung disease in young patients with cystic fibrosis. Respiratory Research, 2007, 8, 88.	3.6	28
90	SARS-CoV-2 B.1.1.529 (Omicron) Variant Causes an Unprecedented Surge in Children Hospitalizations and Distinct Clinical Presentation Compared to the SARS-CoV-2 B.1.617.2 (Delta) Variant. Frontiers in Pediatrics, 0, 10, .	1.9	14