

Nadia Nathan

List of Publications by Year in descending order

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Version: 2024-02-01

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	<i>MUC5B</i> Promoter Variant and Rheumatoid Arthritis with Interstitial Lung Disease. <i>New England Journal of Medicine</i> , 2018, 379, 2209-2219.	27.0	326
2	Alveolar epithelial cells: Master regulators of lung homeostasis. <i>International Journal of Biochemistry and Cell Biology</i> , 2013, 45, 2568-2573.	2.8	187
3	Shared genetic predisposition in rheumatoid arthritis-interstitial lung disease and familial pulmonary fibrosis. <i>European Respiratory Journal</i> , 2017, 49, 1602314.	6.7	154
4	Loss-of-Function Mutations in RSPH1 Cause Primary Ciliary Dyskinesia with Central-Complex and Radial-Spoke Defects. <i>American Journal of Human Genetics</i> , 2013, 93, 561-570.	6.2	148
5	Physiologic Benefits of Mechanical Insufflation-Exsufflation in Children With Neuromuscular Diseases. <i>Chest</i> , 2008, 133, 161-168.	0.8	130
6	Mutations in <i>COPA</i> lead to abnormal trafficking of STING to the Golgi and interferon signaling. <i>Journal of Experimental Medicine</i> , 2020, 217, .	8.5	130
7	Germline <i>SFTPA1</i> mutation in familial idiopathic interstitial pneumonia and lung cancer. <i>Human Molecular Genetics</i> , 2016, 25, 1457-1467.	2.9	119
8	Interstitial lung diseases in children. <i>Orphanet Journal of Rare Diseases</i> , 2010, 5, 22.	2.7	112
9	New insights into pediatric idiopathic pulmonary hemosiderosis: the French RespiRare® cohort. <i>Orphanet Journal of Rare Diseases</i> , 2013, 8, 161.	2.7	95
10	A national internet-linked based database for pediatric interstitial lung diseases: the French network. <i>Orphanet Journal of Rare Diseases</i> , 2012, 7, 40.	2.7	79
11	Atypical presentation of COVID-19 in young infants. <i>Lancet, The</i> , 2020, 395, 1481.	13.7	78
12	International management platform for children's interstitial lung disease (chILD-EU). <i>Thorax</i> , 2018, 73, 231-239.	5.6	64
13	Nonsteroidal Anti-Inflammatory Drug without Antibiotics for Acute Viral Infection Increases the Empyema Risk in Children: A Matched Case-Control Study. <i>Journal of Pediatrics</i> , 2016, 175, 47-53.e3.	1.8	58
14	Dramatic improvement after tocilizumab of severe COVID-19 in a child with sickle cell disease and acute chest syndrome. <i>American Journal of Hematology</i> , 2020, 95, E192-E194.	4.1	56
15	Management of suspected monogenic lung fibrosis in a specialised centre. <i>European Respiratory Review</i> , 2017, 26, 160122.	7.1	54
16	Heterogeneity of lung disease associated with NK2 homeobox 1 mutations. <i>Respiratory Medicine</i> , 2017, 129, 16-23.	2.9	54
17	Surfactant protein A: A key player in lung homeostasis. <i>International Journal of Biochemistry and Cell Biology</i> , 2016, 81, 151-155.	2.8	50
18	Lung sarcoidosis in children: update on disease expression and management. <i>Thorax</i> , 2015, 70, 537-542.	5.6	49

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19	Long-term effects of azithromycin in patients with cystic fibrosis. <i>Respiratory Medicine</i> , 2016, 117, 1-6.	2.9	42
20	Familial pulmonary fibrosis. <i>Revue Des Maladies Respiratoires</i> , 2015, 32, 413-434.	1.7	39
21	Efficacy of Blebs Detection for Preventive Surgery in Children's Idiopathic Spontaneous Pneumothorax. <i>World Journal of Surgery</i> , 2010, 34, 185-189.	1.6	36
22	Use of ruxolitinib in COPA syndrome manifesting as life-threatening alveolar haemorrhage. <i>Thorax</i> , 2020, 75, 92-95.	5.6	36
23	Management of children with interstitial lung diseases: the difficult issue of acute exacerbations. <i>European Respiratory Journal</i> , 2016, 48, 1559-1563.	6.7	33
24	Diffuse parenchymal lung disease caused by surfactant deficiency: dramatic improvement by azithromycin. <i>BMJ Case Reports</i> , 2013, 2013, bcr2013009988-bcr2013009988.	0.5	32
25	Pulmonary hemosiderosis in children with Down syndrome: a national experience. <i>Orphanet Journal of Rare Diseases</i> , 2018, 13, 60.	2.7	32
26	Eosinophilic pneumonias in children: A review of the epidemiology, diagnosis, and treatment. <i>Pediatric Pulmonology</i> , 2016, 51, 203-216.	2.0	31
27	Paediatric sarcoidosis. <i>Paediatric Respiratory Reviews</i> , 2019, 29, 53-59.	1.8	31
28	Genetic causes and clinical management of pediatric interstitial lung diseases. <i>Current Opinion in Pulmonary Medicine</i> , 2018, 24, 253-259.	2.6	30
29	Glucocorticoid receptor gene polymorphisms associated with progression of lung disease in young patients with cystic fibrosis. <i>Respiratory Research</i> , 2007, 8, 88.	3.6	28
30	The Wide Spectrum of COVID-19 Clinical Presentation in Children. <i>Journal of Clinical Medicine</i> , 2020, 9, 2950.	2.4	28
31	Familial vs. sporadic sarcoidosis: BTNL2 polymorphisms, clinical presentations, and outcomes in a French cohort. <i>Orphanet Journal of Rare Diseases</i> , 2016, 11, 165.	2.7	27
32	Whole exome sequencing in three families segregating a pediatric case of sarcoidosis. <i>BMC Medical Genomics</i> , 2018, 11, 23.	1.5	26
33	Interstitial lung diseases in children. <i>Presse Medicale</i> , 2020, 49, 103909.	1.9	26
34	Interstitial lung disease: Physiopathology in the context of lung growth. <i>Paediatric Respiratory Reviews</i> , 2011, 12, 216-222.	1.8	25
35	Pilot experience of multidisciplinary team discussion dedicated to inherited pulmonary fibrosis. <i>Orphanet Journal of Rare Diseases</i> , 2019, 14, 280.	2.7	24
36	Functional assessment and phenotypic heterogeneity of <i>SFTPA1</i> and <i>SFTPA2</i> mutations in interstitial lung diseases and lung cancer. <i>European Respiratory Journal</i> , 2020, 56, 2002806.	6.7	23

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37	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
38	Bi-allelic missense <i>ABCA3</i> mutations in a patient with childhood ILD who reached adulthood. ERJ Open Research, 2019, 5, 00066-2019.	2.6	22
39	Macrolides: New therapeutic perspectives in lung diseases. International Journal of Biochemistry and Cell Biology, 2011, 43, 1241-1246.	2.8	21
40	COPA syndrome, 5 years after: Where are we?. Joint Bone Spine, 2021, 88, 105070.	1.6	21
41	Biomarkers in Interstitial lung diseases. Paediatric Respiratory Reviews, 2015, 16, 219-224.	1.8	20
42	Idiopathic eosinophilic pneumonia in children: the French experience. Orphanet Journal of Rare Diseases, 2014, 9, 28.	2.7	19
43	Pulmonary Fibrosis in Children. Journal of Clinical Medicine, 2019, 8, 1312.	2.4	19
44	Chronic interstitial lung diseases in children: diagnosis approaches. Expert Review of Respiratory Medicine, 2018, 12, 1051-1060.	2.5	16
45	Chronic eosinophilic pneumonia in a 13-year-old child. European Journal of Pediatrics, 2008, 167, 1203-1207.	2.7	14
46	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
47	Health-related quality of life in infants and children with interstitial lung disease. Pediatric Pulmonology, 2019, 54, 828-836.	2.0	13
48	Child-Adult Transition in Sarcoidosis: A Series of 52 Patients. Journal of Clinical Medicine, 2020, 9, 2097.	2.4	13
49	Respiratory Distress, Congenital Hypothyroidism and Hypotonia in a Newborn. Respiration, 2016, 92, 188-191.	2.6	10
50	Exposure to inorganic particles in paediatric sarcoidosis: the PEDIASARC study. Thorax, 2021, , thoraxjnl-2021-217870.	5.6	10
51	Work of breathing in children with diffuse parenchymal lung disease. Respiratory Physiology and Neurobiology, 2015, 206, 45-52.	1.6	8
52	AB0007...Shared genetic predisposition in rheumatoid arthritis...interstitial lung disease and familial pulmonary fibrosis. , 2017, , .		7
53	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
54	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

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55	Respiratory Outcome in Children with Scimitar Syndrome. <i>Journal of Pediatrics</i> , 2013, 162, 275-279.e1.	1.8	5
56	Endobronchial avium mycobacteria infection in an immunocompetent child. <i>BMJ Case Reports</i> , 2013, 2013, bcr2013200776-bcr2013200776.	0.5	5
57	Usefulness of bronchoalveolar lavage in a French pediatric cohort with hypersensitivity pneumonitis. <i>Pediatric Pulmonology</i> , 2020, 55, 136-140.	2.0	5
58	A Nonsmoker Man in His 40s With a Diagnosis of Genetic-Related Idiopathic Pulmonary Fibrosis (Surfactant-Protein C Gene Mutation). <i>Chest</i> , 2019, 155, e91-e96.	0.8	4
59	An idiopathic congenital chylothorax: surgery or conservative management?. <i>BMJ Case Reports</i> , 2014, 2014, bcr2014204147-bcr2014204147.	0.5	3
60	Ultrasonography and Computed Tomographic Manifestations of Abdominal Sarcoidosis in Children. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2016, 63, 195-199.	1.8	3
61	Interstitial lung diseases in the neonatal period. , 2021, , 213-230.		2
62	Pulmonary sarcoid-like granulomatous disease in an 11-month-old girl. <i>BMJ Case Reports</i> , 2013, 2013, bcr2012008024-bcr2012008024.	0.5	2
63	Benefits and risks of bronchoalveolar lavage in severe asthma in children. <i>ERJ Open Research</i> , 2021, 7, 00332-2021.	2.6	2
64	Childhood Interstitial Lung Diseases (chILD) Recognition: When Epidemiology Increases a Rare Disease Incidence. <i>Archivos De Bronconeumologia</i> , 2022, , .	0.8	2
65	French national cohort of neuroendocrine cell hyperplasia of infancy (FRENCHI) study: diagnosis and initial management. <i>European Journal of Pediatrics</i> , 2022, 181, 3067-3073.	2.7	2
66	Inorganic exposome in pediatric sarcoidosis: The PEDIASARC study. , 2015, , .		1
67	GermlineSFTPA1mutation in familial idiopathic interstitial pneumonia and lung cancer. , 2016, , .		1
68	Contribution of mutations in genes of the surfactant system to idiopathic interstitial pneumonia (IIP). , 2018, , .		1
69	Down syndrome and pulmonary hemosiderosis: an under-recognized association. , 2018, , .		1
70	COPA syndrome restricted to life-threatening alveolar hemorrhages: clinical, pathological, molecular and biological characterization. , 2018, , .		1
71	Multidisciplinary team dedicated to suspected heritable pulmonary fibrosis. , 2018, , .		1
72	Azithromycin In Interstitial Lung Disease Associated With Surfactant Metabolism Disorders. , 2010, , .		0

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73	A National French Web Database For Children Rare Lung Diseases. , 2010, , .		0
74	Search For Interstitial Lung Disease Etiology In Children: A Step By Step Approach. , 2011, , .		0
75	Inborn Errors of Metabolism: The Achilles' Heel of the Respiratory System. Respiration, 2017, 94, 14-15.	2.6	0
76	OP0107â€¦HETEROZYGOUS MUTATIONS IN COPA ARE ASSOCIATED WITH ENHANCED TYPE I INTERFERON SIGNALLING. , 2019, , .		0
77	Interstitial lung diseases. , 2021, , 588-601.		0
78	Syndrome COPA, quoi de neuf cinq ans aprÃ©s? . Revue Du Rhumatisme (Edition Francaise), 2021, 88, 183-189.	0.0	0
79	Diffuse Parenchymal Lung Disease in Early Childhood. , 2022, , 229-243.		0
80	Genetic testing in idiopathic interstitial pneumonia. , 2015, , .		0
81	Phenotype heterogeneity in a familial â€œbrain lung thyroid syndromeâ€ related to a novel <i>NKX-2.1</i> mutation. , 2016, , .		0
82	SFTPA mutations in interstitial lung disease (ILD) and lung cancer. , 2017, , .		0
83	A genetic landscape of familial predisposition to sarcoidosis identified by whole exome sequencing. , 2017, , .		0
84	OP0284â€¦Muc5b promoter variant rs35705950 is a risk factor for rheumatoid arthritis â€œ interstitial lung disease. , 2018, , .		0
85	TERT/TERC mutations in a Greek cohort of suspected genetic pulmonary fibrosis patients. , 2018, , .		0
86	Lung disease caused by non-null ABCA3 mutations: long-term follow-up. , 2018, , .		0
87	Functional assessment of newly identified SFTPA1 and SFTPA2 mutations in patients with idiopathic interstitial pneumonia (IIP) and lung cancer. , 2018, , .		0
88	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
89	Health-related quality of life in children interstitial lung disease. , 2019, , .		0
90	Paediatric sarcoidosis. , 2022, , 285-294.		0