Ralph Epaud

List of Publications by Year
in descending order

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87 papers	3,542 citations	136950 32 h-index	149698 56 g-index
98 all docs	98 docs citations	98 times ranked	3998 citing authors

#	Article	IF	Citations
1	Pulmonary alveolar proteinosis. European Respiratory Review, 2011, 20, 98-107.	7.1	212
2	European protocols for the diagnosis and initial treatment of interstitial lung disease in children. Thorax, 2015, 70, 1078-1084.	5.6	192
3	Benign hereditary chorea: phenotype, prognosis, therapeutic outcome and long term follow-up in a large series with new mutations in theTITF1/NKX2-1gene. Journal of Neurology, Neurosurgery and Psychiatry, 2012, 83, 956-962.	1.9	172
4	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
5	Chronic and acute anemia and extracranial internal carotid stenosis are risk factors for silent cerebral infarcts in sickle cell anemia. Blood, 2015, 125, 1653-1661.	1.4	144
6	Rapid Improvement after Starting Elexacaftorâ€“Tezacaftorâ€“Ivacaftor in Patients with Cystic Fibrosis and Advanced Pulmonary Disease. American Journal of Respiratory and Critical Care Medicine, 2021, 204, 64-73.	5.6	139
7	Bacterial Killing Is Enhanced by Expression of Lysozyme in the Lungs of Transgenic Mice. Journal of Immunology, 2000, 165, 5760-5766.	0.8	112
8	Interstitial lung diseases in children. Orphanet Journal of Rare Diseases, 2010, 5, 22.	2.7	112
9	NKX2-1mutations leading to surfactant protein promoter dysregulation cause interstitial lung disease in â€œBrain-Lung-Thyroid Syndromeâ€. Human Mutation, 2010, 31, E1146-E1162.	2.5	108
10	Characteristics of disorders associated with genetic mutations of surfactant protein C. Archives of Disease in Childhood, 2010, 95, 449-454.	1.9	103
11	New surfactant protein C gene mutations associated with diffuse lung disease. Journal of Medical Genetics, 2009, 46, 490-494.	3.2	100
12	New insights into pediatric idiopathic pulmonary hemosiderosis: the French RespiRareÂ® cohort. Orphanet Journal of Rare Diseases, 2013, 8, 161.	2.7	95
13	Molecular and cellular characteristics of ABCA3 mutations associated with diffuse parenchymal lung diseases in children. Human Molecular Genetics, 2012, 21, 765-775.	2.9	85
14	Surfactant Protein B Inhibits Endotoxin-Induced Lung Inflammation. American Journal of Respiratory Cell and Molecular Biology, 2003, 28, 373-378.	2.9	79
15	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
16	Long-term treatment follow-up of children with sickle cell disease monitored with abnormal transcranial Doppler velocities. Blood, 2016, 127, 1814-1822.	1.4	79
17	Effectiveness of Chest Physiotherapy in Infants Hospitalized with Acute Bronchiolitis: A Multicenter, Randomized, Controlled Trial. PLoS Medicine, 2010, 7, e1000345.	8.4	71
18	Incidence of paediatric pneumococcal meningitis and emergence of new serotypes: a time-series analysis of a 16-year French national survey. Lancet Infectious Diseases, The, 2018, 18, 983-991.	9.1	69

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19	Combined pulmonary fibrosis and emphysema syndrome associated with ABCA3 mutations. European Respiratory Journal, 2014, 43, 638-641.	6.7	68
20	Familial interstitial disease with I73T mutation: A midâ€•and longâ€•term study. Pediatric Pulmonology, 2009, 44, 167-175.	2.0	67
21	Impact of COVIDâ€•19 social distancing on viral infection in France: A delayed outbreak of RSV. Pediatric Pulmonology, 2021, 56, 3669-3673.	2.0	64
22	Surfactant protein C gene (SFTPC) mutationâ€•associated lung disease: Highâ€•resolution computed tomography (HRCT) findings and its relation to histological analysis. Pediatric Pulmonology, 2010, 45, 1021-1029.	2.0	58
23	Knockout of Insulin-Like Growth Factor-1 Receptor Impairs Distal Lung Morphogenesis. PLoS ONE, 2012, 7, e48071.	2.5	56
24	Heterogeneity of lung disease associated with NK2 homeobox 1 mutations. Respiratory Medicine, 2017, 129, 16-23.	2.9	54
25	Lung alveolar epithelium and interstitial lung disease. International Journal of Biochemistry and Cell Biology, 2009, 41, 1643-1651.	2.8	50
26	Lung sarcoidosis in children: update on disease expression and management. Thorax, 2015, 70, 537-542.	5.6	49
27	Assessment of blood enterovirus PCR testing in paediatric populations with fever without source, sepsis-like disease, or suspected meningitis: a prospective, multicentre, observational cohort study. Lancet Infectious Diseases, The, 2018, 18, 1385-1396.	9.1	43
28	Effect of Nebulized Hypertonic Saline Treatment in Emergency Departments on the Hospitalization Rate for Acute Bronchiolitis. JAMA Pediatrics, 2017, 171, e171333.	6.2	41
29	A randomised trial of high-flow nasal cannula in infants with moderate bronchiolitis. European Respiratory Journal, 2020, 56, 1901926.	6.7	40
30	Therapeutic strategies for idiopathic chylothorax. Journal of Pediatric Surgery, 2008, 43, 461-465.	1.6	37
31	Efficacy of Blebs Detection for Preventive Surgery in Childrenâ€™s Idiopathic Spontaneous Pneumothorax. World Journal of Surgery, 2010, 34, 185-189.	1.6	36
32	Combined Relay Therapy With Oral Cefixime and Clavulanate for Febrile Urinary Tract Infection Caused by Extended-Spectrum Î²-lactamaseâ€“producing Escherichia coli. Pediatric Infectious Disease Journal, 2013, 32, 96-97.	2.0	33
33	Management of children with interstitial lung diseases: the difficult issue of acute exacerbations. European Respiratory Journal, 2016, 48, 1559-1563.	6.7	33
34	Epidemiology and Clinical Presentation of Children Hospitalized with SARS-CoV-2 Infection in Suburbs of Paris. Journal of Clinical Medicine, 2020, 9, 2227.	2.4	33
35	EMLA Cream and Nitrous Oxide to Alleviate Pain Induced by Palivizumab (Synagis) Intramuscular Injections in Infants and Young Children. Pediatrics, 2008, 121, e1591-e1598.	2.1	32
36	Diffuse parenchymal lung disease caused by surfactant deficiency: dramatic improvement by azithromycin. BMJ Case Reports, 2013, 2013, bcr2013009988-bcr2013009988.	0.5	32

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37	Pulmonary hemosiderosis in children with Down syndrome: a national experience. Orphanet Journal of Rare Diseases, 2018, 13, 60.	2.7	32
38	Deficiency in type 1 insulin-like growth factor receptor in mice protects against oxygen-induced lung injury. Respiratory Research, 2005, 6, 31.	3.6	30
39	Research in progress: put the orphanage out of business: TableÂ1. Thorax, 2013, 68, 971-973.	5.6	28
40	Conservative use of chest-tube insertion in children with pleural effusion. Pediatric Surgery International, 2006, 22, 357-362.	1.4	26
41	Acute cervical lymphadenitis and infections of the retropharyngeal and parapharyngeal spaces in children. BMC Ear, Nose and Throat Disorders, 2014, 14, 8.	2.6	24
42	Biological impact of Î± genes, Î² haplotypes, and G6PD activity in sickle cell anemia at baseline and with hydroxyurea. Blood Advances, 2018, 2, 626-637.	5.2	24
43	Pilot experience of multidisciplinary team discussion dedicated to inherited pulmonary fibrosis. Orphanet Journal of Rare Diseases, 2019, 14, 280.	2.7	24
44	Clinical predictors of radiographic abnormalities among infants with bronchiolitis in a paediatric emergency department. BMC Pediatrics, 2014, 14, 143.	1.7	22
45	Sarcoidosis in children: HRCT findings and correlation with pulmonary function tests. Pediatric Pulmonology, 2014, 49, 1223-1233.	2.0	22
46	Tezacaftor/ivacaftor in people with cystic fibrosis who stopped lumacaftor/ivacaftor due to respiratory adverse events. Journal of Cystic Fibrosis, 2021, 20, 228-233.	0.7	21
47	Multiplex Ligation-Dependent Probe Amplification Improves the Detection Rate of <i>NKX2.1</i> Mutations in Patients Affected by Brain-Lung-Thyroid Syndrome. Hormone Research in Paediatrics, 2012, 77, 146-151.	1.8	20
48	Cladribine improves lung cysts and pulmonary function in a child with histiocytosis. European Respiratory Journal, 2015, 45, 831-833.	6.7	20
49	A recurrent deep-intronic splicing CF mutation emphasizes the importance of mRNA studies in clinical practice. Journal of Cystic Fibrosis, 2011, 10, 479-482.	0.7	19
50	Serotype 3 Pneumococcal Pleural Empyema in an Immunocompetent Child after 13-valent Pneumococcal Conjugate Vaccine. Pediatric Infectious Disease Journal, 2014, 33, 545-546.	2.0	17
51	Delayed acute bronchiolitis in infants hospitalized for COVIDâ€19. Pediatric Pulmonology, 2020, 55, 2211-2212.	2.0	16
52	Retrospective French nationwide survey of childhood aggressive vascular anomalies of bone, 1988-2009. Orphanet Journal of Rare Diseases, 2010, 5, 3.	2.7	15
53	New use for an old drug: COXâ€independent antiâ€inflammatory effects of sulindac in models of cystic fibrosis. British Journal of Pharmacology, 2016, 173, 1728-1741.	5.4	15
54	Targeting p16^{INK4a} Promotes Lipofibroblasts and Alveolar Regeneration after Early-Life Injury. American Journal of Respiratory and Critical Care Medicine, 2020, 202, 1088-1104.	5.6	15

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55	Mild cystic fibrosis revealed by persistent hyponatremia during the French 2003 heat wave, associated with the S1455X C-terminus CFTR mutation. Clinical Genetics, 2005, 68, 552-553.	2.0	14
56	Chronic eosinophilic pneumonia in a 13-year-old child. European Journal of Pediatrics, 2008, 167, 1203-1207.	2.7	14
57	Healthâ€related quality of life in infants and children with interstitial lung disease. Pediatric Pulmonology, 2019, 54, 828-836.	2.0	13
58	Childâ€“Adult Transition in Sarcoidosis: A Series of 52 Patients. Journal of Clinical Medicine, 2020, 9, 2097.	2.4	13
59	Impact of Chest Radiography for Children with Lower Respiratory Tract Infection: A Propensity Score Approach. PLoS ONE, 2014, 9, e96189.	2.5	12
60	Deciphering the mechanism of Q145H SFTPC mutation unmasks a splicing defect and explains the severity of the phenotype. European Journal of Human Genetics, 2017, 25, 779-782.	2.8	12
61	Clinical presentation of interstitial lung disease in children. Paediatric Respiratory Reviews, 2004, 5, 98-100.	1.8	11
62	Survival of an infant with homozygous surfactant protein C (SFTPC) mutation. Pediatric Pulmonology, 2014, 49, E112-5.	2.0	11
63	Brief report: International perspectives on the pediatric COVIDâ€19 experience. Pediatric Pulmonology, 2020, 55, 1598-1600.	2.0	10
64	Exposure to inorganic particles in paediatric sarcoidosis: the PEDIASARC study. Thorax, 2021, , thoraxjnl-2021-217870.	5.6	10
65	Scopulariopsis brevicaulis abscess in a child treated for myeloblastic leukaemia. Lancet Infectious Diseases, The, 2011, 11, 416.	9.1	9
66	Nonâ€invasive CT screening for pulmonary arteriovenous malformations in children with confirmed hereditary hemorrhagic telangiectasia: Results from two pediatric centers. Pediatric Pulmonology, 2017, 52, 642-649.	2.0	8
67	Unusual long survival despite severe lung disease of a child with biallelic loss of function mutations in ABCA-3. Respiratory Medicine Case Reports, 2018, 23, 173-175.	0.4	8
68	NKX2.1 (TTF1) germline mutation associated with pulmonary fibrosis and lung cancer. ERJ Open Research, 2021, 7, 00356-2021.	2.6	8
69	Pulmonary and hepatic nodular lesions precede the diagnosis of Crohnâ€™s disease in an 8â€yearâ€old girl: a case study and review of the literature. Acta Paediatrica, International Journal of Paediatrics, 2012, 101, e86-9.	1.5	7
70	Protein-losing gastropathy associated with cytomegalovirus in two sisters â€“ Case reports and review of the literature. Archives De Pediatrie, 2019, 26, 232-235.	1.0	7
71	Effectiveness of palivizumab in children with childhood interstitial lung disease: The French experience. Pediatric Pulmonology, 2016, 51, 688-695.	2.0	6
72	Identification of Clinical and Laboratory Parameters Associated with the Development of Acute Chest Syndrome during Vaso-Occlusive Episodes in Children with Sickle Cell Disease: A Preliminary Step before Assessing Specific and Early Treatment Strategies. Journal of Clinical Medicine, 2019, 8, 1839.	2.4	6

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73	Serum Immunoglobulin Levels in Children with Sickle Cell Disease: A Large Prospective Study. Journal of Clinical Medicine, 2019, 8, 1688.	2.4	5
74	BAL Fluid Surfactant Protein C Level Is Related to Parenchymal Lung Disease in Children With Sarcoidosis. Chest, 2011, 140, 1104-1105.	0.8	3
75	Ultrasonography and Computed Tomographic Manifestations of Abdominal Sarcoidosis in Children. Journal of Pediatric Gastroenterology and Nutrition, 2016, 63, 195-199.	1.8	3
76	Clinical course and cost assessment of infants with a first episode of acute bronchiolitis presenting to the emergency department: Data from the GUERANDE clinical trial. Pediatric Pulmonology, 2021, 56, 3802-3812.	2.0	3
77	An update on paediatric respiratory diseases. European Respiratory Review, 2018, 27, 180013.	7.1	1
78	Methylprednisolone pulse treatment improves ProSPâ€C trafficking in twins with SFTPC mutation: An isoform story?. British Journal of Clinical Pharmacology, 2021, 87, 2361-2373.	2.4	1
79	Prise enÂcharge desÂpneumonies chezÂl'enfant. Journal De Pediatrie Et De Puericulture, 2006, 19, 145-148.	0.0	0
80	High Resolution Computed Tomography in Children with SFTPC Gene Mutation and Its Relationship to Histological Findings from the Open Lung Biopsies.. , 2009, , .		0
81	Dramatic Improvement by Macrolides in Surfactant Deficiency with ABCA3 Mutation.. , 2009, , .		0
82	Identification and Characterization of Two New TTF-1 Mutations Associated with Pediatric Interstitial Lung Diseases.. , 2009, , .		0
83	Azithromycin In Interstitial Lung Disease Associated With Surfactant Metabolism Disorders. , 2010, , .		0
84	New Mutations Of ABCA3 Associated With Neonatal Respiratory Distress And Diffuse Lung Disease. , 2010, , .		0
85	Interstitial lung disease reveals 48,XXYY syndrome in a child. Acta Paediatrica, International Journal of Paediatrics, 2020, 109, 1060-1061.	1.5	0
86	BiothÃ©rapies dans lâ€™asthme sÃ©vÃ¨re de lâ€™enfant et de lâ€™adolescent. Revue Des Maladies Respiratoires Actualites, 2020, 12, 2S415-2S422.	0.0	0
87	Impact of a rare respiratory diseases reference centre set-up on primary ciliary dyskinesia care pathway. European Respiratory Journal, 2022, 59, 2102413.	6.7	0