

Jos D Ribeiro

List of Publications by Year in Descending Order

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Version: 2024-04-20

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

165
papers

2,046
citations

24
h-index

35
g-index

206
ext. papers

2,470
ext. citations

2.5
avg, IF

4.94
L-index

#	Paper	IF	Citations
165	Can continuous glucose monitoring predict cystic fibrosis-related diabetes and worse clinical outcome?. <i>Jornal Brasileiro De Pneumologia</i> , 2022 , 48, e20210307	1.1	
164	Applicability of lung ultrasound in COVID-19 diagnosis and evaluation of the disease progression: A systematic review. <i>Pulmonology</i> , 2021 , 27, 529-562	3.7	14
163	Are there differences in the physical activity level and functional capacity among children and adolescents with and without asthma?. <i>Jornal De Pediatria</i> , 2021 , 97, 295-301	2.6	1
162	Reliability of impulse oscillometry parameters in healthy children and in children with cystic fibrosis. <i>International Journal of Clinical Practice</i> , 2021 , 75, e13715	2.9	1
161	Risk factors for recurrent wheezing in preterm infants who received prophylaxis with palivizumab. <i>Jornal Brasileiro De Pneumologia</i> , 2021 , 47, e20210157	1.1	0
160	Distal intestinal obstruction syndrome: a diagnostic and therapeutic challenge in cystic fibrosis. <i>Jornal De Pediatria (Versão Em Português)</i> , 2020 , 96, 732-740	0.2	
159	Multichannel intraluminal impedance-pH and psychometric properties in gastroesophageal reflux: systematic review. <i>Jornal De Pediatria (Versão Em Português)</i> , 2020 , 96, 673-685	0.2	
158	Prevalence of constipation in cystic fibrosis patients: a systematic review of observational studies. <i>Jornal De Pediatria</i> , 2020 , 96, 686-692	2.6	2
157	Volumetric capnography versus spirometry for the evaluation of pulmonary function in cystic fibrosis and allergic asthma. <i>Jornal De Pediatria (Versão Em Português)</i> , 2020 , 96, 255-263	0.2	
156	Lymphocyte responses to Mycobacterium tuberculosis and Mycobacterium bovis are similar between BCG-vaccinated patients with cystic fibrosis and healthy controls. <i>Journal of Cystic Fibrosis</i> , 2020 , 19, 575-579	4.1	1
155	Multichannel intraluminal impedance-pH and psychometric properties in gastroesophageal reflux: systematic review. <i>Jornal De Pediatria</i> , 2020 , 96, 673-685	2.6	0
154	Assessment of asthma control among different measures and evaluation of functional exercise capacity in children and adolescents with asthma. <i>Jornal Brasileiro De Pneumologia</i> , 2020 , 46, e20190102	1.1	3
153	MON-LB114 CGM in Cystic Fibrosis Patients to Predict Cystic Fibrosis-Related Diabetes Onset. <i>Journal of the Endocrine Society</i> , 2020 , 4,	0.4	78
152	Safety, Tolerability, and Effects of Sodium Bicarbonate Inhalation in Cystic Fibrosis. <i>Clinical Drug Investigation</i> , 2020 , 40, 105-117	3.2	14
151	The Use of Ultrasound as a Tool to Evaluate Pulmonary Disease in Cystic Fibrosis. <i>Respiratory Care</i> , 2020 , 65, 293-303	2.1	4
150	Distal intestinal obstruction syndrome: a diagnostic and therapeutic challenge in cystic fibrosis. <i>Jornal De Pediatria</i> , 2020 , 96, 732-740	2.6	2
149	Differences between patients who achieved asthma control and those who remain uncontrolled after standardized severe asthma care strategy. <i>Journal of Asthma</i> , 2020 , 1-13	1.9	0

148	Prevalence of constipation in cystic fibrosis patients: a systematic review of observational studies. <i>Jornal De Pediatria (Versão Em Português)</i> , 2020 , 96, 686-692	0.2	
147	Lung function in obese children and adolescents without respiratory disease: a systematic review. <i>BMC Pulmonary Medicine</i> , 2020 , 20, 281	3.5	4
146	Volumetric capnography versus spirometry for the evaluation of pulmonary function in cystic fibrosis and allergic asthma. <i>Jornal De Pediatria</i> , 2020 , 96, 255-264	2.6	1
145	A negative screening of rare genetic variants in the ADIPOQ and STATH genes in cystic fibrosis. <i>Pulmonology</i> , 2020 , 26, 138-144	3.7	0
144	Chloride and sodium ion concentrations in saliva and sweat as a method to diagnose cystic fibrosis. <i>Jornal De Pediatria (Versão Em Português)</i> , 2019 , 95, 443-450	0.2	1
143	Lung disease and vitamin D levels in cystic fibrosis infants and preschoolers. <i>Pediatric Pulmonology</i> , 2019 , 54, 563-574	3.5	4
142	Analysis of motor and respiratory function in Duchenne muscular dystrophy patients. <i>Respiratory Physiology and Neurobiology</i> , 2019 , 262, 1-11	2.8	4
141	Secretory IgA-mediated immune response in saliva and early detection of <i>Pseudomonas aeruginosa</i> in the lower airways of pediatric cystic fibrosis patients. <i>Medical Microbiology and Immunology</i> , 2019 , 208, 205-213	4	6
140	Extent of rescue of F508del-CFTR function by VX-809 and VX-770 in human nasal epithelial cells correlates with SNP rs7512462 in SLC26A9 gene in F508del/F508del Cystic Fibrosis patients. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2019 , 1865, 1323-1331	6.9	17
139	Severe pulmonary disease in an adult primary ciliary dyskinesia population in Brazil. <i>Scientific Reports</i> , 2019 , 9, 8693	4.9	7
138	Microbiological contamination of nebulizers used by cystic fibrosis patients: an underestimated problem. <i>Jornal Brasileiro De Pneumologia</i> , 2019 , 45, e20170351	1.1	13
137	Evaluation of respiratory dynamics by volumetric capnography during submaximal exercise protocol of six minutes on treadmill in cystic fibrosis patients. <i>Jornal De Pediatria (Versão Em Português)</i> , 2019 , 95, 76-86	0.2	
136	Novel, rare and common pathogenic variants in the CFTR gene screened by high-throughput sequencing technology and predicted by in silico tools. <i>Scientific Reports</i> , 2019 , 9, 6234	4.9	15
135	Concordance between whole- and half-body scans to evaluate body composition in dual-energy X-ray absorptiometry in children and adolescents with different nutritional and pubertal conditions. <i>Nutrition</i> , 2019 , 66, 78-86	4.8	1
134	Recurrent wheezing in preterm infants: Prevalence and risk factors. <i>Jornal De Pediatria</i> , 2019 , 95, 720-726	2.6	5
133	Chloride and sodium ion concentrations in saliva and sweat as a method to diagnose cystic fibrosis. <i>Jornal De Pediatria</i> , 2019 , 95, 443-450	2.6	9
132	Correlation between parameters of volumetric capnography and spirometry during a submaximal exercise protocol on a treadmill in patients with cystic fibrosis and healthy controls. <i>Pulmonology</i> , 2019 , 25, 21-31	3.7	2
131	Lung ultrasound assessment of response to antibiotic therapy in cystic fibrosis exacerbations: a study of two cases. <i>Jornal Brasileiro De Pneumologia</i> , 2019 , 45, e20190128	1.1	1

130	Pseudomonas aeruginosa colonization in the upper and lower airways of a child with cystic fibrosis: a father's meticulous approach to successful eradication. <i>Jornal Brasileiro De Pneumologia</i> , 2019 , 45, e20190191	1.1	
129	Recurrent wheezing in preterm infants: Prevalence and risk factors. <i>Jornal De Pediatria (Versão Em Português)</i> , 2019 , 95, 720-727	0.2	
128	Evaluation of respiratory dynamics by volumetric capnography during submaximal exercise protocol of six minutes on treadmill in cystic fibrosis patients. <i>Jornal De Pediatria</i> , 2019 , 95, 76-86	2.6	
127	Interaction among variants in the SLC gene family (SLC6A14, SLC26A9, SLC11A1, and SLC9A3) and CFTR mutations with clinical markers of cystic fibrosis. <i>Pediatric Pulmonology</i> , 2018 , 53, 888-900	3.5	7
126	Association between oxygenation and ventilation indices with the time on invasive mechanical ventilation in infants. <i>Pulmonology</i> , 2018 , 24, 241-241	3.7	0
125	Continuous glucose monitoring to evaluate glycaemic abnormalities in cystic fibrosis. <i>Archives of Disease in Childhood</i> , 2018 , 103, 592-596	2.2	3
124	Association between single nucleotide polymorphisms in TLR4, TLR2, TLR9, VDR, NOS2 and CCL5 genes with acute viral bronchiolitis. <i>Gene</i> , 2018 , 645, 7-17	3.8	17
123	Cystic fibrosis transmembrane regulator haplotypes in households of patients with cystic fibrosis. <i>Gene</i> , 2018 , 641, 137-143	3.8	1
122	Impulse oscillometry and obesity in children. <i>Jornal De Pediatria</i> , 2018 , 94, 419-424	2.6	5
121	Impulse oscillometry and obesity in children. <i>Jornal De Pediatria (Versão Em Português)</i> , 2018 , 94, 419-424	0.2	
120	Prevalence and clinical outcomes of nontuberculous mycobacteria in a Brazilian cystic fibrosis reference center. <i>Pathogens and Disease</i> , 2018 , 76,	4.2	6
119	Use of Phase III Slope of Volumetric Capnography in Outpatient Clinical Practice: A Descriptive Analysis. <i>Current Respiratory Medicine Reviews</i> , 2018 , 13, 152-158	0.3	2
118	Respiratory syncytial virus in Brazilian infants - Ten years, two cohorts. <i>Journal of Clinical Virology</i> , 2018 , 98, 33-36	14.5	5
117	Pancreatic Insufficiency in Cystic Fibrosis: Influence of Inflammatory Response Genes. <i>Pancreas</i> , 2018 , 47, 99-109	2.6	3
116	Evaluation of continuous constant current and continuous pulsed current in sweat induction for cystic fibrosis diagnosis. <i>BMC Pulmonary Medicine</i> , 2018 , 18, 153	3.5	0
115	The correlation between age and sweat chloride levels in sweat tests. <i>Revista Portuguesa De Pneumologia</i> , 2017 , 23, 227-230		4
114	Alternative Indexes to Estimate the Functional Capacity From the 6-Minute Walk Test in Children and Adolescents With Cystic Fibrosis. <i>Respiratory Care</i> , 2017 , 62, 324-332	2.1	8
113	Hypertonic Saline as a Useful Tool for Sputum Induction and Pathogen Detection in Cystic Fibrosis. <i>Lung</i> , 2017 , 195, 431-439	2.9	4

112	Spirometry and volumetric capnography in lung function assessment of obese and normal-weight individuals without asthma. <i>Jornal De Pediatria</i> , 2017 , 93, 398-405	2.6	5
111	Physical performance, quality of life and sexual satisfaction evaluation in adults with cystic fibrosis: An unexplored correlation. <i>Revista Portuguesa De Pneumologia</i> , 2017 , 23, 179-192		3
110	Physical activity and asthma control level in children and adolescents. <i>Respirology</i> , 2017 , 22, 1643-1648	3.6	15
109	Burkholderia cepacia complex in cystic fibrosis in a Brazilian reference center. <i>Medical Microbiology and Immunology</i> , 2017 , 206, 447-461	4	2
108	Sweat test and cystic fibrosis: overview of test performance at public and private centers in the state of S ^ˆ ˆ Paulo, Brazil. <i>Jornal Brasileiro De Pneumologia</i> , 2017 , 43, 121-128	1.1	12
107	Brazilian guidelines for the diagnosis and treatment of cystic fibrosis. <i>Jornal Brasileiro De Pneumologia</i> , 2017 , 43, 219-245	1.1	39
106	Skin Biomarkers for Cystic Fibrosis: A Potential Non-Invasive Approach for Patient Screening. <i>Frontiers in Pediatrics</i> , 2017 , 5, 290	3.4	9
105	Variants in the interleukin 8 gene and the response to inhaled bronchodilators in cystic fibrosis. <i>Jornal De Pediatria</i> , 2017 , 93, 639-648	2.6	7
104	Association of clinical severity of cystic fibrosis with variants in the SLC gene family (SLC6A14, SLC26A9, SLC11A1 and SLC9A3). <i>Gene</i> , 2017 , 629, 117-126	3.8	22
103	Spirometry and volumetric capnography in lung function assessment of obese and normal-weight individuals without asthma. <i>Jornal De Pediatria (Versˆo Em Portuguˆs)</i> , 2017 , 93, 398-405	0.2	
102	Impulse oscillometry, spirometry, and passive smoking in healthy children and adolescents. <i>Revista Portuguesa De Pneumologia</i> , 2017 , 23, 311-316		2
101	Secretory IgA response against Pseudomonas aeruginosa in the upper airways and the link with chronic lung infection in cystic fibrosis. <i>Pathogens and Disease</i> , 2017 , 75,	4.2	8
100	Variants in the interleukin 8 gene and the response to inhaled bronchodilators in cystic fibrosis. <i>Jornal De Pediatria (Versˆo Em Portuguˆs)</i> , 2017 , 93, 639-648	0.2	
99	Personalized or Precision Medicine? The Example of Cystic Fibrosis. <i>Frontiers in Pharmacology</i> , 2017 , 8, 390	5.6	39
98	Thirty Years of Sweat Chloride Testing at One Referral Center. <i>Frontiers in Pediatrics</i> , 2017 , 5, 222	3.4	8
97	Impulse Oscillometry System and Anthropometric Variables of Preschoolers, Children and Adolescents Systematic Review. <i>Current Pediatric Reviews</i> , 2017 , 13, 126-135	2.8	7
96	Correlation between acoustic rhinometry, computed rhinomanometry and cone-beam computed tomography in mouth breathers with transverse maxillary deficiency. <i>Brazilian Journal of Otorhinolaryngology</i> , 2016 , 84, 40-40	1.6	1
95	Classification of CFTR mutation classes. <i>Lancet Respiratory Medicine</i> , 2016 , 4, e37-e38	35.1	76

94	Insulin therapy in patients with cystic fibrosis in the pre-diabetes stage: a systematic review. <i>Revista Paulista De Pediatria (English Edition)</i> , 2016 , 34, 367-73		6
93	Quality of sweat test (ST) based on the proportion of sweat sodium (Na) and sweat chloride (Cl) as diagnostic parameter of cystic fibrosis: are we on the right way?. <i>Diagnostic Pathology</i> , 2016 , 11, 103	3	3
92	Associação entre variáveis clínicas relacionadas à asma em escolares nascidos com muito baixo peso com e sem displasia broncopulmonar. <i>Revista Paulista De Pediatria</i> , 2016 , 34, 271-280	1.2	2
91	Association of growth and nutritional parameters with pulmonary function in cystic fibrosis: a literature review. <i>Revista Paulista De Pediatria (English Edition)</i> , 2016 , 34, 503-509		9
90	Insulinoterapia em pacientes com fibrose cística na fase de pré-diabetes: uma revisão sistemática. <i>Revista Paulista De Pediatria</i> , 2016 , 34, 367-373	1.2	9
89	VIVER COM FIBROSE CÍSTICA: A VISÃO PESSOAL DO ADOLESCENTE BRASILEIRO. <i>Psicologia Em Estudo</i> , 2016 , 21, 211	0.7	1
88	IL8 gene as modifier of cystic fibrosis: unraveling the factors which influence clinical variability. <i>Human Genetics</i> , 2016 , 135, 881-94	6.3	19
87	Association of growth and nutritional parameters with pulmonary function in cystic fibrosis: a literature review. <i>Revista Paulista De Pediatria</i> , 2016 , 34, 503-509	1.2	10
86	Association between clinical variables related to asthma in schoolchildren born with very low birth weight with and without bronchopulmonary dysplasia. <i>Revista Paulista De Pediatria (English Edition)</i> , 2016 , 34, 271-80		
85	Chronic obstructive pulmonary diseases in children. <i>Jornal De Pediatria</i> , 2015 , 91, S11-25	2.6	7
84	Ventilatory abnormalities in patients with cystic fibrosis undergoing the submaximal treadmill exercise test. <i>BMC Pulmonary Medicine</i> , 2015 , 15, 63	3.5	5
83	Chronic obstructive pulmonary diseases in children. <i>Jornal De Pediatria (Versão Em Português)</i> , 2015 , 91, S11-S25	0.2	
82	Evaluation of quality of life according to asthma control and asthma severity in children and adolescents. <i>Jornal Brasileiro De Pneumologia</i> , 2015 , 41, 502-8	1.1	24
81	Nasal potential difference in cystic fibrosis considering severe CFTR mutations. <i>Disease Markers</i> , 2015 , 2015, 306825	3.2	7
80	Ventilatory Efficiency in Children and Adolescents: A Systematic Review. <i>Disease Markers</i> , 2015 , 2015, 546891	3.2	4
79	Demographic, clinical, and laboratory parameters of cystic fibrosis during the last two decades: a comparative analysis. <i>BMC Pulmonary Medicine</i> , 2015 , 15, 3	3.5	12
78	Personalized Drug Therapy in Cystic Fibrosis: From Fiction to Reality. <i>Current Drug Targets</i> , 2015 , 16, 1007-17	3	19
77	CFTR genotype and clinical outcomes of adult patients carried as cystic fibrosis disease. <i>Gene</i> , 2014 , 540, 183-90	3.8	32

76	Asthma: Gln27Glu and Arg16Gly polymorphisms of the beta2-adrenergic receptor gene as risk factors. <i>Allergy, Asthma and Clinical Immunology</i> , 2014 , 10, 8	3.2	34
75	Effect of exercise test on pulmonary function of obese adolescents. <i>Jornal De Pediatria</i> , 2014 , 90, 242-9	2.6	18
74	Polymorphisms in the glutathione pathway modulate cystic fibrosis severity: a cross-sectional study. <i>BMC Medical Genetics</i> , 2014 , 15, 27	2.1	25
73	Effect of exercise test on pulmonary function of obese adolescents. <i>Jornal De Pediatria (Versão Em Português)</i> , 2014 , 90, 242-249	0.2	3
72	Epidemiological aspects of and risk factors for wheezing in the first year of life. <i>Jornal Brasileiro De Pneumologia</i> , 2014 , 40, 617-25	1.1	9
71	Pulsed direct and constant direct currents in the pilocarpine iontophoresis sweat chloride test. <i>BMC Pulmonary Medicine</i> , 2014 , 14, 198	3.5	6
70	The relationship between physical functional capacity and lung function in obese children and adolescents. <i>BMC Pulmonary Medicine</i> , 2014 , 14, 199	3.5	8
69	Assessment of IgG antibodies to <i>Pseudomonas aeruginosa</i> in patients with cystic fibrosis by an enzyme-linked immunosorbent assay (ELISA). <i>Diagnostic Pathology</i> , 2014 , 9, 158	3	15
68	SLC23A2-05 (rs4987219) and KRAS-LCS6 (rs61764370) polymorphisms in patients with squamous cell carcinoma of the head and neck. <i>Oncology Letters</i> , 2014 , 7, 1803-1811	2.6	4
67	TNF-alpha polymorphisms as a potential modifier gene in the cystic fibrosis. <i>International Journal of Molecular Epidemiology and Genetics</i> , 2014 , 5, 87-99	0.9	6
66	Saliva as a potential tool for cystic fibrosis diagnosis. <i>Diagnostic Pathology</i> , 2013 , 8, 46	3	39
65	Rectal forceps biopsy procedure in cystic fibrosis: technical aspects and patients perspective for clinical trials feasibility. <i>BMC Gastroenterology</i> , 2013 , 13, 91	3	22
64	Severe lower respiratory tract infection in infants and toddlers from a non-affluent population: viral etiology and co-detection as risk factors. <i>BMC Infectious Diseases</i> , 2013 , 13, 41	4	45
63	Genetic interaction of GSH metabolic pathway genes in cystic fibrosis. <i>BMC Medical Genetics</i> , 2013 , 14, 60	2.1	23
62	Obesity increases eosinophil activity in asthmatic children and adolescents. <i>BMC Pulmonary Medicine</i> , 2013 , 13, 39	3.5	50
61	Epidemiological and genetic characteristics associated with the severity of acute viral bronchiolitis by respiratory syncytial virus. <i>Jornal De Pediatria (Versão Em Português)</i> , 2013 , 89, 531-543	0.2	1
60	Epidemiological and genetic characteristics associated with the severity of acute viral bronchiolitis by respiratory syncytial virus. <i>Jornal De Pediatria</i> , 2013 , 89, 531-43	2.6	38
59	Manual therapy for childhood respiratory disease: a systematic review. <i>Journal of Manipulative and Physiological Therapeutics</i> , 2013 , 36, 57-65	1.3	7

58	Neonatal and pediatric manual hyperinflation: influence of oxygen flow on ventilation parameters. <i>Respiratory Care</i> , 2013 , 58, 2127-33	2.1	1
57	APC germline mutations in families with familial adenomatous polyposis. <i>Oncology Reports</i> , 2013 , 30, 2081-8	3.5	14
56	Walk test and school performance in mouth-breathing children. <i>Brazilian Journal of Otorhinolaryngology</i> , 2013 , 79, 212-8	1.6	7
55	Cystic fibrosis transmembrane conductance regulator mutations at a referral center for cystic fibrosis. <i>Jornal Brasileiro De Pneumologia</i> , 2013 , 39, 555-61	1.1	4
54	<i>Pseudomonas aeruginosa</i> infection in patients with cystic fibrosis: scientific evidence regarding clinical impact, diagnosis, and treatment. <i>Jornal Brasileiro De Pneumologia</i> , 2013 , 39, 495-512	1.1	39
53	Screening for F508del as a first step in the molecular diagnosis of cystic fibrosis. <i>Jornal Brasileiro De Pneumologia</i> , 2013 , 39, 306-16	1.1	8
52	Effect of omalizumab as add-on therapy on asthma-related quality of life in severe allergic asthma: a Brazilian study (QUALITX). <i>Journal of Asthma</i> , 2012 , 49, 288-93	1.9	30
51	Does experience influence the performance of neonatal and pediatric manual hyperinflation?. <i>Respiratory Care</i> , 2012 , 57, 1908-13	2.1	1
50	Polymorphisms in ADRB2 gene can modulate the response to bronchodilators and the severity of cystic fibrosis. <i>BMC Pulmonary Medicine</i> , 2012 , 12, 50	3.5	29
49	The ACE gene D/I polymorphism as a modulator of severity of cystic fibrosis. <i>BMC Pulmonary Medicine</i> , 2012 , 12, 41	3.5	21
48	Evaluation of oral functions of the stomatognathic system according to the levels of asthma severity. <i>Jornal Da Sociedade Brasileira De Fonoaudiologia</i> , 2012 , 24, 119-24		8
47	Measurements of CFTR-mediated Cl ⁻ secretion in human rectal biopsies constitute a robust biomarker for Cystic Fibrosis diagnosis and prognosis. <i>PLoS ONE</i> , 2012 , 7, e47708	3.7	42
46	<i>Mycoplasma pneumoniae</i> -related community-acquired pneumonia and parapneumonic pleural effusion in children and adolescents. <i>Jornal Brasileiro De Pneumologia</i> , 2012 , 38, 226-36	1.1	4
45	Lipoid pneumonia in a 40-day-old infant. <i>Jornal Brasileiro De Pneumologia</i> , 2012 , 38, 535-7	1.1	2
44	Influence of thoracic spine postural disorders on cardiorespiratory parameters in children and adolescents with cystic fibrosis. <i>Jornal De Pediatria</i> , 2012 , 88, 310-6	2.6	10
43	Volumetric capnography as a tool to detect early peripheral lung obstruction in cystic fibrosis patients. <i>Jornal De Pediatria</i> , 2012 , 88, 509-17	2.6	18
42	Fatores que afetam a ventilação com o reanimador manual autoinflável: uma revisão sistemática. <i>Revista Paulista De Pediatria</i> , 2011 , 29, 645-655	1.2	2
41	Quality of life assessment in patients with cystic fibrosis by means of the Cystic Fibrosis Questionnaire. <i>Jornal Brasileiro De Pneumologia</i> , 2011 , 37, 184-92	1.1	15

40	Functional performance on the six-minute walk test in patients with cystic fibrosis. <i>Jornal Brasileiro De Pneumologia</i> , 2011 , 37, 735-44	1.1	14
39	Mouth breathing and forward head posture: effects on respiratory biomechanics and exercise capacity in children. <i>Jornal Brasileiro De Pneumologia</i> , 2011 , 37, 471-9	1.1	26
38	Rela ^ç o entre ^o \dot{V}_E e ventila ^ç o mec ^â nica de pacientes em terapia intensiva pedi ^á tica. <i>Revista Paulista De Pediatria</i> , 2011 , 29, 348-351	1.2	2
37	Exercise capacity, respiratory mechanics and posture in mouth breathers. <i>Brazilian Journal of Otorhinolaryngology</i> , 2011 , 77, 656-62	1.6	18
36	Volumetric capnography to detect ventilation inhomogeneity in children and adolescents with controlled persistent asthma. <i>Jornal De Pediatria</i> , 2011 , 87, 163-8	2.6	20
35	Assessment of the body posture of mouth-breathing children and adolescents. <i>Jornal De Pediatria</i> , 2011 , 87, 357-63	2.6	17
34	Obesidade e asma: associa ^ç o ou coincid ^ê ncia?. <i>Jornal De Pediatria</i> , 2010 , 86, 6-14	2.6	2
33	Glutathione S-transferase mu 1 (GSTM1) and theta 1 (GSTT1) genetic polymorphisms and atopic asthma in children from Southeastern Brazil. <i>Genetics and Molecular Biology</i> , 2010 , 33, 438-41	2	16
32	Antibiotic therapy and Effects of Respiratory Physiotherapy Techniques Cystic Fibrosis Patients Treated for Acute Lung Exacerbation: an Experimental Study. <i>Archivos De Bronconeumologia</i> , 2010 , 46, 310-316	0.7	0
31	Volumetric capnography for the evaluation of pulmonary disease in adult patients with cystic fibrosis and noncystic fibrosis bronchiectasis. <i>Lung</i> , 2010 , 188, 263-8	2.9	24
30	Avalia ^ç o espirom ^é trica e da hiper-responsividade br ^ô nquica de crian ^ç as e adolescentes com asma at ^é rica persistente moderada submetidos a nata ^ç o. <i>Jornal De Pediatria</i> , 2010 , 86, 384-390	2.6	22
29	Obesity and asthma: association or coincidence?. <i>Jornal De Pediatria</i> , 2010 , 86, 6-14	2.6	5
28	Asthma and swimming: weighing the benefits and the risks. <i>Jornal De Pediatria</i> , 2010 , 86, 351-2	2.6	4
27	Comparison of the effects that two different respiratory physical therapy techniques have on cardiorespiratory parameters in infants with acute viral bronchiolitis. <i>Jornal Brasileiro De Pneumologia</i> , 2009 , 35, 860-7	1.1	18
26	Association of MBL2, TGF-beta1 and CD14 gene polymorphisms with lung disease severity in cystic fibrosis. <i>Jornal Brasileiro De Pneumologia</i> , 2009 , 35, 334-42	1.1	20
25	Pulmonary deposition of inhaled tobramycin prior to and after respiratory therapy and use of inhaled albuterol in cystic fibrosis patients colonized with <i>Pseudomonas aeruginosa</i> . <i>Jornal Brasileiro De Pneumologia</i> , 2009 , 35, 35-43	1.1	4
24	Genetic associations with asthma and virus-induced wheezing: a systematic review. <i>Jornal Brasileiro De Pneumologia</i> , 2009 , 35, 1220-6	1.1	4
23	Risk factors for gastroesophageal reflux disease in very low birth weight infants with bronchopulmonary dysplasia. <i>Jornal De Pediatria</i> , 2008 , 84, 154-9	2.6	14

22	Foreign body aspiration in children: clinical aspects, radiological aspects and bronchoscopic treatment. <i>Jornal Brasileiro De Pneumologia</i> , 2008 , 34, 74-82	1.1	28
21	Association of TGF-beta1, CD14, IL-4, IL-4R and ADAM33 gene polymorphisms with asthma severity in children and adolescents. <i>Jornal De Pediatria</i> , 2008 , 84, 203-10	2.6	28
20	Oral magnesium supplementation in asthmatic children: a double-blind randomized placebo-controlled trial. <i>European Journal of Clinical Nutrition</i> , 2007 , 61, 54-60	5.2	46
19	Relationship between physiologic deadspace/tidal volume ratio and gas exchange in infants with acute bronchiolitis on invasive mechanical ventilation. <i>Pediatric Critical Care Medicine</i> , 2007 , 8, 372-7	3	11
18	Nasal and paranasal sinus endoscopy, computed tomography and microbiology of upper airways and the correlations with genotype and severity of cystic fibrosis. <i>International Journal of Pediatric Otorhinolaryngology</i> , 2007 , 71, 41-50	1.7	43
17	Respiratory syncytial virus (RSV) in infants hospitalized for acute lower respiratory tract disease: incidence and associated risks. <i>Brazilian Journal of Infectious Diseases</i> , 2006 , 10, 357-61	2.8	21
16	Antileucotrienos no tratamento da asma e rinite alérgica. <i>Jornal De Pediatria</i> , 2006 , 82, S213	2.6	1
15	Antileukotrienes in the treatment of asthma and allergic rhinitis. <i>Jornal De Pediatria</i> , 2006 , 82, S213-21	2.6	15
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12	Utilidade de um escore e de variáveis indicativas de drenagem pleural em crianças com derrame pleural parapneumônico. <i>Jornal Brasileiro De Pneumologia</i> , 2005 , 31, 205-211	1.1	2
11	Cystic fibrosis at a Brazilian center of excellence: clinical and laboratory characteristics of 104 patients and their association with genotype and disease severity. <i>Jornal De Pediatria</i> , 2004 , 80, 371-379	2.6	24
10	Fibrose cística em um centro de referência no Brasil: características clínicas e laboratoriais de 104 pacientes e sua associação com o genótipo e a gravidade da doença. <i>Jornal De Pediatria</i> , 2004 , 80,	2.6	2
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