JosÃ© D Ribeiro

List of Publications by Year
in descending order

183	2,838	185998 28	276539 41
papers	citations	h-index	g-index

206	206	206	3726
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Rhinoviruses as critical agents in severe bronchiolitis in infants. Jornal De Pediatria, 2022, 98, 362-368.	0.9	4
2	The Sun also rises. Jornal Brasileiro De Pneumologia, 2022, 47, e20210473.	0.4	0
3	Can continuous glucose monitoring predict cystic fibrosis-related diabetes and worse clinical outcome?. Jornal Brasileiro De Pneumologia, 2022, 48, e20210307.	0.4	5
4	Challenges in Diagnosing Primary Ciliary Dyskinesia in a Brazilian Tertiary Hospital. Genes, 2022, 13, 1252.	1.0	3
5	Are there differences in the physical activity level and functional capacity among children and adolescents with and without asthma?. Jornal De Pediatria, 2021, 97, 295-301.	0.9	5
6	Reliability of impulse oscillometry parameters in healthy children and in children with cystic fibrosis. International Journal of Clinical Practice, 2021, 75, e13715.	0.8	4
7	Applicability of lung ultrasound in COVID-19 diagnosis and evaluation of the disease progression: A systematic review. Pulmonology, 2021, 27, 529-562.	1.0	41
8	Risk factors for recurrent wheezing in preterm infants who received prophylaxis with palivizumab. Jornal Brasileiro De Pneumologia, 2021, 47, e20210157.	0.4	1
9	Volumetric capnography versus spirometry for the evaluation of pulmonary function in cystic fibrosis and allergic asthma. Jornal De Pediatria, 2020, 96, 255-264.	0.9	6
10	A negative screening of rare genetic variants in the ADIPOQ and STATH genes in cystic fibrosis. Pulmonology, 2020, 26, 138-144.	1.0	2
11	Safety, Tolerability, and Effects of Sodium Bicarbonate Inhalation in Cystic Fibrosis. Clinical Drug Investigation, 2020, 40, 105-117.	1.1	20
12	The Use of Ultrasound as a Tool to Evaluate Pulmonary Disease in Cystic Fibrosis. Respiratory Care, 2020, 65, 293-303.	0.8	12
13	Distal intestinal obstruction syndrome: a diagnostic and therapeutic challenge in cystic fibrosis. Jornal De Pediatria, 2020, 96, 732-740.	0.9	8
14	Differences between patients who achieved asthma control and those who remain uncontrolled after standardized severe asthma care strategy. Journal of Asthma, 2020, , 1-13.	0.9	4
15	Lung function in obese children and adolescents without respiratory disease: a systematic review. BMC Pulmonary Medicine, 2020, 20, 281.	0.8	10
16	Prevalence of constipation in cystic fibrosis patients: a systematic review of observational studies. Jornal De Pediatria, 2020, 96, 686-692.	0.9	4
17	P103 Lymphocyte responses to Mycobacterium tuberculosis and Mycobacterium bovis are similar between BCG-vaccinated patients with cystic fibrosis and healthy controls. Journal of Cystic Fibrosis, 2020, 19, S84.	0.3	0
18	Volumetric capnography versus spirometry for the evaluation of pulmonary function in cystic fibrosis and allergic asthma. Jornal De Pediatria (VersÃ£o Em PortuguÃªs), 2020, 96, 255-263.	0.2	0

#	Article	IF	Citations
19	Lymphocyte responses to Mycobacterium tuberculosis and Mycobacterium bovis are similar between BCG-vaccinated patients with cystic fibrosis and healthy controls. Journal of Cystic Fibrosis, 2020, 19, 575-579.	0.3	1
20	Multichannel intraluminal impedance-pH and psychometric properties in gastroesophageal reflux: systematic review. Jornal De Pediatria, 2020, 96, 673-685.	0.9	4
21	Assessment of asthma control among different measures and evaluation of functional exercise capacity in children and adolescents with asthma. Jornal Brasileiro De Pneumologia, 2020, 46, e20190102-e20190102.	0.4	4
22	MON-LB114 CGM in Cystic Fibrosis Patients to Predict Cystic Fibrosis-Related Diabetes Onset. Journal of the Endocrine Society, 2020, 4, .	0.1	0
23	Recurrent wheezing in preterm infants: Prevalence and risk factors. Jornal De Pediatria, 2019, 95, 720-727.	0.9	11
24	Chloride and sodium ion concentrations in saliva and sweat as a method to diagnose cystic fibrosis. Jornal De Pediatria, 2019, 95, 443-450.	0.9	22
25	Correlation between parameters of volumetric capnography and spirometry during a submaximal exercise protocol on a treadmill in patients with cystic fibrosis and healthy controls. Pulmonology, 2019, 25, 21-31.	1.0	2
26	Chloride and sodium ion concentrations in saliva and sweat as a method to diagnose cystic fibrosis. Jornal De Pediatria (VersÃ£o Em PortuguÃªs), 2019, 95, 443-450.	0.2	1
27	Lung disease and vitamin D levels in cystic fibrosis infants and preschoolers. Pediatric Pulmonology, 2019, 54, 563-574.	1.0	10
28	Analysis of motor and respiratory function in Duchenne muscular dystrophy patients. Respiratory Physiology and Neurobiology, 2019, 262, 1-11.	0.7	6
29	Secretory IgA-mediated immune response in saliva and early detection of Pseudomonas aeruginosa in the lower airways of pediatric cystic fibrosis patients. Medical Microbiology and Immunology, 2019, 208, 205-213.	2.6	13
30	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. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2019, 1865, 1323-1331.	1.8	28
31	Severe pulmonary disease in an adult primary ciliary dyskinesia population in Brazil. Scientific Reports, 2019, 9, 8693.	1.6	15
32	Microbiological contamination of nebulizers used by cystic fibrosis patients: an underestimated problem. Jornal Brasileiro De Pneumologia, 2019, 45, e20170351.	0.4	25
33	P173 Improvement of the diagnosis of P. aeruginosa infection in cystic fibrosis using real-time PCR: a pilot analysis. Journal of Cystic Fibrosis, 2019, 18, S106.	0.3	0
34	Evaluation of respiratory dynamics by volumetric capnography during submaximal exercise protocol of six minutes on treadmill in cystic fibrosis patients. Jornal De Pediatria (VersÃ£o Em PortuguÃªs), 2019, 95, 76-86.	0.2	0
35	Novel, rare and common pathogenic variants in the CFTR gene screened by high-throughput sequencing technology and predicted by in silico tools. Scientific Reports, 2019, 9, 6234.	1.6	33
36	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. Nutrition, 2019, 66, 78-86.	1.1	2

#	Article	IF	Citations
37	Recurrent wheezing in preterm infants: Prevalence and risk factors. Jornal De Pediatria (VersÃ£o Em) Tj ETQq1 1 0.784314 rgBT /Over	0.2	0
38	Evaluation of respiratory dynamics by volumetric capnography during submaximal exercise protocol of six minutes on treadmill in cystic fibrosis patients. Jornal De Pediatria, 2019, 95, 76-86.	0.9	0
39	Lung ultrasound assessment of response to antibiotic therapy in cystic fibrosis exacerbations: a study of two cases. Jornal Brasileiro De Pneumologia, 2019, 45, e20190128.	0.4	5
40	Pseudomonas aeruginosa colonization in the upper and lower airways of a child with cystic fibrosis: a fatherâ€™s meticulous approach to successful eradication. Jornal Brasileiro De Pneumologia, 2019, 45, e20190191.	0.4	1
41	Are there differences in the level of physical activity and functional capacity between healthy and asthmatic children and adolescents?. , 2019, , .		0
42	Clinical characteristics, lung function, quality of life, exercise capacity and inflammatory biomarkers in induced sputum and serum in children and adolescents with uncontrolled severe asthma. , 2019, , .		0
43	Interaction among variants in the SLC gene family (SLC6A14, SLC26A9,) Tj ETQq1 1 0.784314 rgBT /Overlock 10 T Pediatric Pulmonology, 2018, 53, 888-900.	1.0	15
44	Association between oxygenation and ventilation indices with the time on invasive mechanical ventilation in infants. Pulmonology, 2018, 24, 241-249.	1.0	3
45	Continuous glucose monitoring to evaluate glycaemic abnormalities in cystic fibrosis. Archives of Disease in Childhood, 2018, 103, 592-596.	1.0	9
46	Association between single nucleotide polymorphisms in TLR4 , TLR2 , TLR9 , VDR , NOS2 and CCL5 genes with acute viral bronchiolitis. Gene, 2018, 645, 7-17.	1.0	24
47	Correlation between acoustic rhinometry, computed rhinomanometry and cone-beam computed tomography in mouth breathers with transverse maxillary deficiency. Brazilian Journal of Otorhinolaryngology, 2018, 84, 40-50.	0.4	1
48	Cystic fibrosis transmembrane regulator haplotypes in households of patients with cystic fibrosis. Gene, 2018, 641, 137-143.	1.0	1
49	Impulse oscillometry and obesity in children. Jornal De Pediatria, 2018, 94, 419-424.	0.9	10
50	Respiratory syncytial virus in Brazilian infants â€“ Ten years, two cohorts. Journal of Clinical Virology, 2018, 98, 33-36.	1.6	6
51	Pancreatic Insufficiency in Cystic Fibrosis. Pancreas, 2018, 47, 99-109.	0.5	5
52	Evaluation of continuous constant current and continuous pulsed current in sweat induction for cystic fibrosis diagnosis. BMC Pulmonary Medicine, 2018, 18, 153.	0.8	1
53	Impulse oscillometry and obesity in children. Jornal De Pediatria (VersÃ£o Em PortuguÃªs), 2018, 94, 419-424.	0.2	0
54	Prevalence and clinical outcomes of nontuberculous mycobacteria in a Brazilian cystic fibrosis reference center. Pathogens and Disease, 2018, 76, .	0.8	9

#	Article	IF	Citations
55	Impulse Oscillometry System and Anthropometric Variables of Preschoolers, Children and Adolescents Systematic Review. Current Pediatric Reviews, 2018, 13, 126-135.	0.4	8
56	Use of Phase III Slope of Volumetric Capnography in Outpatient Clinical Practice: A Descriptive Analysis. Current Respiratory Medicine Reviews, 2018, 13, 152-158.	0.1	2
57	The correlation between age and sweat chloride levels in sweat tests. Revista Portuguesa De Pneumologia, 2017, 23, 227-230.	0.7	5
58	Alternative Indexes to Estimate the Functional Capacity From the 6-Minute Walk Test in Children and Adolescents With Cystic Fibrosis. Respiratory Care, 2017, 62, 324-332.	0.8	10
59	Hypertonic Saline as a Useful Tool for Sputum Induction and Pathogen Detection in Cystic Fibrosis. Lung, 2017, 195, 431-439.	1.4	9
60	Spirometry and volumetric capnography in lung function assessment of obese and normal-weight individuals without asthma. Jornal De Pediatria, 2017, 93, 398-405.	0.9	9
61	Physical performance, quality of life and sexual satisfaction evaluation in adults with cystic fibrosis: An unexplored correlation. Revista Portuguesa De Pneumologia, 2017, 23, 179-192.	0.7	4
62	Physical activity and asthma control level in children and adolescents. Respirology, 2017, 22, 1643-1648.	1.3	18
63	13 Saliva for newborn screening for CF. Journal of Cystic Fibrosis, 2017, 16, S66.	0.3	0
64	Burkholderia cepacia complex in cystic fibrosis in a Brazilian reference center. Medical Microbiology and Immunology, 2017, 206, 447-461.	2.6	3
65	Variants in the interleukin 8 gene and the response to inhaled bronchodilators in cystic fibrosis. Jornal De Pediatria, 2017, 93, 639-648.	0.9	7
66	WS10.3 Chloride in saliva and sweat in age-matched individuals with and without CF. Journal of Cystic Fibrosis, 2017, 16, S18.	0.3	0
67	Association of clinical severity of cystic fibrosis with variants in the SLC gene family (SLC6A14,) Tj ETQq1 1 0.784314 rgBT /Overlock 10	1.0	34
68	Impulse oscillometry, spirometry, and passive smoking in healthy children and adolescents. Revista Portuguesa De Pneumologia, 2017, 23, 311-316.	0.7	3
69	Secretory IgA response against Pseudomonas aeruginosa in the upper airways and the link with chronic lung infection in cystic fibrosis. Pathogens and Disease, 2017, 75, .	0.8	13
70	Personalized or Precision Medicine? The Example of Cystic Fibrosis. Frontiers in Pharmacology, 2017, 8, 390.	1.6	56
71	Thirty Years of Sweat Chloride Testing at One Referral Center. Frontiers in Pediatrics, 2017, 5, 222.	0.9	10
72	Sweat test and cystic fibrosis: overview of test performance at public and private centers in the state of SÃ£o Paulo, Brazil. Jornal Brasileiro De Pneumologia, 2017, 43, 121-128.	0.4	18

#	Article	IF	Citations
73	Brazilian guidelines for the diagnosis and treatment of cystic fibrosis. Jornal Brasileiro De Pneumologia, 2017, 43, 219-245.	0.4	73
74	Skin Biomarkers for Cystic Fibrosis: A Potential Non-Invasive Approach for Patient Screening. Frontiers in Pediatrics, 2017, 5, 290.	0.9	12
75	VIVER COM FIBROSE CÃSTICA: A VISÃƒO PESSOAL DO ADOLESCENTE BRASILEIRO. Psicologia Em Estudo, 2016, 21, 211.	0.2	1
76	IL8 gene as modifier of cystic fibrosis: unraveling the factors which influence clinical variability. Human Genetics, 2016, 135, 881-894.	1.8	22
77	AssociaÃ§Ã£o dos parÃ¢metros de crescimento e nutricionais com funÃ§Ã£o pulmonar na fibrose cÃstica: revisÃ£o da literatura. Revista Paulista De Pediatria, 2016, 34, 503-509.	0.4	16
78	Association between clinical variables related to asthma in schoolchildren born with very low birth weight with and without bronchopulmonary dysplasia. Revista Paulista De Pediatria (English Edition), 2016, 34, 271-280.	0.3	4
79	103 Low prevalence of mycobacteria among Brazilian CF patients: possible explanations. Journal of Cystic Fibrosis, 2016, 15, S77.	0.3	0
80	48 The humoral immune response against Pseudomonas aeruginosa correlates with decreased lung function in a cohort of Brazilian CF patients. Journal of Cystic Fibrosis, 2016, 15, S63-S64.	0.3	0
81	Classification of CFTR mutation classes. Lancet Respiratory Medicine,the, 2016, 4, e37-e38.	5.2	115
82	110 Pitfalls in the diagnosis of Aspergillus disease in a cohort of Brazilian CF patients. Journal of Cystic Fibrosis, 2016, 15, S79.	0.3	0
83	Insulin therapy in patients with cystic fibrosis in the pre-diabetes stage: a systematic review. Revista Paulista De Pediatria (English Edition), 2016, 34, 367-373.	0.3	10
84	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?. Diagnostic Pathology, 2016, 11, 103.	0.9	4
85	AssociaÃ§Ã£o entre variÃ¡veis clÃnicas relacionadas Ã asma em escolares nascidos com muito baixo peso com e sem displasia broncopulmonar. Revista Paulista De Pediatria, 2016, 34, 271-280.	0.4	9
86	Association of growth and nutritional parameters with pulmonary function in cystic fibrosis: a literature review. Revista Paulista De Pediatria (English Edition), 2016, 34, 503-509.	0.3	16
87	Insulinoterapia em pacientes com fibrose cÃstica na fase de prÃ©â€diabetes: uma revisÃ£o sistemÃ¡tica. Revista Paulista De Pediatria, 2016, 34, 367-373.	0.4	11
88	Severe acute viral bronchiolitis: A genetic entity. , 2016, , .		0
89	Influence of single nucleotide polymorphisms in post-bronchiolitis wheezing. , 2016, , .		0
90	Chronic obstructive pulmonary diseases in children. Jornal De Pediatria (VersÃ£o Em PortuguÃªs), 2015, 91, S11-S25.	0.2	0

#	Article	IF	Citations
91	Evaluation of quality of life according to asthma control and asthma severity in children and adolescents. Jornal Brasileiro De Pneumologia, 2015, 41, 502-508.	0.4	33
92	Nasal Potential Difference in Cystic Fibrosis considering SevereCFTRMutations. Disease Markers, 2015, 2015, 1-11.	0.6	7
93	Ventilatory Efficiency in Children and Adolescents: A Systematic Review. Disease Markers, 2015, 2015, 1-10.	0.6	8
94	Demographic, clinical, and laboratory parameters of cystic fibrosis during the last two decades: a comparative analysis. BMC Pulmonary Medicine, 2015, 15, 3.	0.8	14
95	63 Pseudomonas aeruginosa pulmonary infection in CF patients in a Brazilian reference center: Antibody response monitoring. Journal of Cystic Fibrosis, 2015, 14, S73.	0.3	0
96	Chronic obstructive pulmonary diseases in children. Jornal De Pediatria, 2015, 91, S11-S25.	0.9	9
97	Ventilatory abnormalities in patients with cystic fibrosis undergoing the submaximal treadmill exercise test. BMC Pulmonary Medicine, 2015, 15, 63.	0.8	8
98	Personalized Drug Therapy in Cystic Fibrosis: From Fiction to Reality. Current Drug Targets, 2015, 16, 1007-1017.	1.0	25
99	OscilaÃ§Ã£o oral de alta frequÃªncia e fibrose cÃstica: comparaÃ§Ã£o entre instrumentais. ConScientiae SaÃºde, 2015, 14, 283-290.	0.1	1
100	Epidemiological aspects of and risk factors for wheezing in the first year of life. Jornal Brasileiro De Pneumologia, 2014, 40, 617-625.	0.4	11
101	Pulsed direct and constant direct currents in the pilocarpine iontophoresis sweat chloride test. BMC Pulmonary Medicine, 2014, 14, 198.	0.8	8
102	The relationship between physical functional capacity and lung function in obese children and adolescents. BMC Pulmonary Medicine, 2014, 14, 199.	0.8	15
103	Assessment of IgG antibodies to Pseudomonas aeruginosa in patients with cystic fibrosis by an enzyme-linked immunosorbent assay (ELISA). Diagnostic Pathology, 2014, 9, 158.	0.9	22
104	SLC23A2-05 (rs4987219) and KRAS-LCS6 (rs61764370) polymorphisms in patients with squamous cell carcinoma of the head and neck. Oncology Letters, 2014, 7, 1803-1811.	0.8	8
105	CFTR genotype and clinical outcomes of adult patients carried as cystic fibrosis disease. Gene, 2014, 540, 183-190.	1.0	37
106	Asthma: Gln27Glu and Arg16Gly polymorphisms of the beta2-adrenergic receptor gene as risk factors. Allergy, Asthma and Clinical Immunology, 2014, 10, 8.	0.9	38
107	Effect of exercise test on pulmonary function of obese adolescents. Jornal De Pediatria, 2014, 90, 242-249.	0.9	27
108	Polymorphisms in the glutathione pathway modulate cystic fibrosis severity: a cross-sectional study. BMC Medical Genetics, 2014, 15, 27.	2.1	28

#	Article	IF	Citations
109	Effect of exercise test on pulmonary function of obese adolescents. Jornal De Pediatria (VersÃ£o Em) Tj ETQq1 1 0.784314 rgBT /Over	0.2	3
110	32 Correlation between structural and functional lung injury in children and adolescents with cystic fibrosis. Journal of Cystic Fibrosis, 2014, 13, S53.	0.3	0
111	TNF-alpha polymorphisms as a potential modifier gene in the cystic fibrosis. International Journal of Molecular Epidemiology and Genetics, 2014, 5, 87-99.	0.4	7
112	Saliva as a potential tool for cystic fibrosis diagnosis. Diagnostic Pathology, 2013, 8, 46.	0.9	45
113	Rectal forceps biopsy procedure in cystic fibrosis: technical aspects and patients perspective for clinical trials feasibility. BMC Gastroenterology, 2013, 13, 91.	0.8	24
114	Severe lower respiratory tract infection in infants and toddlers from a non-affluent population: viral etiology and co-detection as risk factors. BMC Infectious Diseases, 2013, 13, 41.	1.3	60
115	Genetic interaction of GSH metabolic pathway genes in cystic fibrosis. BMC Medical Genetics, 2013, 14, 60.	2.1	24
116	Obesity increases eosinophil activity in asthmatic children and adolescents. BMC Pulmonary Medicine, 2013, 13, 39.	0.8	62
117	Epidemiological and genetic characteristics associated with the severity of acute viral bronchiolitis by respiratory syncytial virus. Jornal De Pediatria (VersÃ£o Em PortuguÃªs), 2013, 89, 531-543.	0.2	1
118	Epidemiological and genetic characteristics associated with the severity of acute viral bronchiolitis by respiratory syncytial virus. Jornal De Pediatria, 2013, 89, 531-543.	0.9	55
119	Manual Therapy for Childhood Respiratory Disease: A Systematic Review. Journal of Manipulative and Physiological Therapeutics, 2013, 36, 57-65.	0.4	13
120	Neonatal and Pediatric Manual Hyperinflation: Influence of Oxygen Flow on Ventilation Parameters. Respiratory Care, 2013, 58, 2127-2133.	0.8	2
121	APC germline mutations in families with familial adenomatous polyposis. Oncology Reports, 2013, 30, 2081-2088.	1.2	16
122	Walk test and school performance in mouth-breathing children. Brazilian Journal of Otorhinolaryngology, 2013, 79, 212-218.	0.4	13
123	Cystic fibrosis transmembrane conductance regulator mutations at a referral center for cystic fibrosis. Jornal Brasileiro De Pneumologia, 2013, 39, 555-561.	0.4	6
124	Pseudomonas aeruginosa infection in patients with cystic fibrosis: scientific evidence regarding clinical impact, diagnosis, and treatment. Jornal Brasileiro De Pneumologia, 2013, 39, 495-512.	0.4	49
125	Screening for F508del as a first step in the molecular diagnosis of cystic fibrosis. Jornal Brasileiro De Pneumologia, 2013, 39, 306-316.	0.4	11
126	Measurements of CFTR-Mediated Clâ¯ Secretion in Human Rectal Biopsies Constitute a Robust Biomarker for Cystic Fibrosis Diagnosis and Prognosis. PLoS ONE, 2012, 7, e47708.	1.1	52

#	Article	IF	Citations
127	Effect of Omalizumab as Add-On Therapy on Asthma-Related Quality of Life in Severe Allergic Asthma: A Brazilian Study (QUALITX). Journal of Asthma, 2012, 49, 288-293.	0.9	40
128	Does Experience Influence the Performance of Neonatal and Pediatric Manual Hyperinflation?. Respiratory Care, 2012, 57, 1908-1913.	0.8	4
129	Polymorphisms in ADRB2 gene can modulate the response to bronchodilators and the severity of cystic fibrosis. BMC Pulmonary Medicine, 2012, 12, 50.	0.8	31
130	The ACE gene D/I polymorphism as a modulator of severity of cystic fibrosis. BMC Pulmonary Medicine, 2012, 12, 41.	0.8	26
131	Evaluation of oral functions of the stomatognathic system according to the levels of asthma severity. Jornal Da Sociedade Brasileira De Fonoaudiologia, 2012, 24, 119-124.	0.4	10
132	Pneumonia adquirida na comunidade e derrame pleural parapneumÃ´nico relacionados a Mycoplasma pneumoniae em crianÃ§as e adolescentes. Jornal Brasileiro De Pneumologia, 2012, 38, 226-236.	0.4	4
133	Pneumonia lipoide em lactente de 40 dias de vida. Jornal Brasileiro De Pneumologia, 2012, 38, 535-537.	0.4	2
134	Influence of thoracic spine postural disorders on cardiorespiratory parameters in children and adolescents with cystic fibrosis. Jornal De Pediatria, 2012, 88, 310-6.	0.9	11
135	Volumetric capnography as a tool to detect early peripheric lung obstruction in cystic fibrosis patients. Jornal De Pediatria, 2012, 88, 509-17.	0.9	20
136	Fatores que afetam a ventilaÃ§Ã£o com o reanimador manual autoinflÃ¡vel: uma revisÃ£o sistemÃ¡tica. Revista Paulista De Pediatria, 2011, 29, 645-655.	0.4	2
137	AvaliaÃ§Ã£o da qualidade de vida de pacientes com fibrose cÃstica por meio do Cystic Fibrosis Questionnaire. Jornal Brasileiro De Pneumologia, 2011, 37, 184-192.	0.4	18
138	Desempenho funcional de pacientes com fibrose cÃstica e indivÃduos saudÃ¡veis no teste de caminhada de seis minutos. Jornal Brasileiro De Pneumologia, 2011, 37, 735-744.	0.4	22
139	RespiraÃ§Ã£o bucal e anterioriza Ã§Ã£o da cabeÃ§a: efeitos na biomecÃ¢nica respiratÃ³ria e na capacidade de exercÃcio em crianÃ§as. Jornal Brasileiro De Pneumologia, 2011, 37, 471-479.	0.4	44
140	RelaÃ§Ã£o entre Ãndice de oxigenaÃ§Ã£o e ventilaÃ§Ã£o com o tempo em ventilaÃ§Ã£o mecÃ¢nica de pacientes em terapia intensiva pediÃ¡trica. Revista Paulista De Pediatria, 2011, 29, 348-351.	0.4	3
141	Exercise capacity, respiratory mechanics and posture in mouth breathers. Brazilian Journal of Otorhinolaryngology, 2011, 77, 656-662.	0.4	23
142	Volumetric capnography to detect ventilation inhomogeneity in children and adolescents with controlled persistent asthma. Jornal De Pediatria, 2011, 87, 163-168.	0.9	23
143	Assessment of the body posture of mouth-breathing children and adolescents. Jornal De Pediatria, 2011, 87, 357-363.	0.9	26
144	Volumetric Capnography for the Evaluation of Pulmonary Disease in Adult Patients with Cystic Fibrosis and Noncystic Fibrosis Bronchiectasis. Lung, 2010, 188, 263-268.	1.4	28

#	Article	IF	Citations
145	Obesidade e asma: associaÃ§Ã£o ou coincidÃªncia?. Jornal De Pediatria, 2010, 86, 6-14.	0.9	9
146	Glutathione S-transferase mu 1 (GSTM1) and theta 1 (GSTT1) genetic polymorphisms and atopic asthma in children from Southeastern Brazil. Genetics and Molecular Biology, 2010, 33, 438-441.	0.6	19
147	Antibiotic therapy and Effects of Respiratory Physiotherapy Techniques Cystic Fibrosis Patients Treated for Acute Lung Exacerbation: an Experimental Study. Archivos De Bronconeumologia, 2010, 46, 310-316.	0.4	1
148	AvaliaÃ§Ã£o espiromÃ©trica e da hiper-responsividade brÃ´nquica de crianÃ§as e adolescentes com asma atÃ³pica persistente moderada submetidos a nataÃ§Ã£o. Jornal De Pediatria, 2010, 86, 384-390.	0.9	26
149	Obesity and asthma: association or coincidence?. Jornal De Pediatria, 2010, 86, 6-14.	0.9	14
150	Asthma and swimming: weighing the benefits and the risks. Jornal De Pediatria, 2010, 86, 384-390.	0.9	7
151	AssociaÃ§Ã£o entre os polimorfismos dos genes MBL2, TGF-Î²1 e CD14 com a gravidade da doenÃ§a pulmonar na fibrose cÃstica. Jornal Brasileiro De Pneumologia, 2009, 35, 334-342.	0.4	24
152	ComparaÃ§Ã£o dos efeitos de duas tÃ©cnicas fisioterapÃªuticas respiratÃ³rias em parÃ¢metros cardiorrespiratÃ³rios de lactentes com bronquiolite viral aguda. Jornal Brasileiro De Pneumologia, 2009, 35, 860-867.	0.4	31
153	DeposiÃ§Ã£o pulmonar de tobramicina inalatÃ³ria antes e apÃ³s fisioterapia respiratÃ³ria e uso de salbutamol inalatÃ³rio em pacientes com fibrose cÃstica colonizados por Pseudomonas aeruginosa. Jornal Brasileiro De Pneumologia, 2009, 35, 35-43.	0.4	6
154	Genetic associations with asthma and virus-induced wheezing: a systematic review. Jornal Brasileiro De Pneumologia, 2009, 35, 1220-1226.	0.4	5
155	Risk factors for gastroesophageal reflux disease in very low birth weight infants with bronchopulmonary dysplasia. Jornal De Pediatria, 2008, 84, 154-159.	0.9	16
156	AspiraÃ§Ã£o de corpo estranho em crianÃ§as: aspectos clÃnicos, radiolÃ³gicos e tratamento broncoscÃ³pico. Jornal Brasileiro De Pneumologia, 2008, 34, 74-82.	0.4	36
157	Association of TGF-β1, CD14, IL-4, IL-4R and ADAM33 gene polymorphisms with asthma severity in children and adolescents. Jornal De Pediatria, 2008, 84, 203-210.	0.9	33
158	Relationship between physiologic deadspace/tidal volume ratio and gas exchange in infants with acute bronchiolitis on invasive mechanical ventilation*. Pediatric Critical Care Medicine, 2007, PAP, 372-7.	0.2	21
159	Nasal and paranasal sinus endoscopy, computed tomography and microbiology of upper airways and the correlations with genotype and severity of cystic fibrosis. International Journal of Pediatric Otorhinolaryngology, 2007, 71, 41-50.	0.4	49
160	Oral magnesium supplementation in asthmatic children: a double-blind randomized placebo-controlled trial. European Journal of Clinical Nutrition, 2007, 61, 54-60.	1.3	60
161	Respiratory syncytial virus (RSV) in infants hospitalized for acute lower respiratory tract disease: incidence and associated risks. Brazilian Journal of Infectious Diseases, 2006, 10, 357-61.	0.3	26
162	Antileucotrienos no tratamento da asma e rinite alÃ©rgica. Jornal De Pediatria, 2006, 82, S213.	0.9	1

#	Article	IF	Citations
163	Antileukotrienes in the treatment of asthma and allergic rhinitis. Jornal De Pediatria, 2006, 82, 213-213.	0.9	19
164	Effect of expiratory flow increase technique on pulmonary function of infants on mechanical ventilation. Physiotherapy Research International, 2005, 10, 213-221.	0.7	17
165	Risk Factors for Bronchopulmonary Dysplasia in very Low Birth Weight Newborns Treated with Mechanical Ventilation in the First Week of Life. Journal of Tropical Pediatrics, 2005, 51, 334-340.	0.7	41
166	AerossÃ³is e espaÃ§adores na crise aguda de asma: evoluÃ§Ã£o e hora de mudar a rotina. Jornal De Pediatria, 2005, 81, 274-276.	0.9	1
167	AssociaÃ§Ã£o entre deficiÃªncia de alfa-1-antitripsina e a gravidade da fibrose cÃstica. Jornal De Pediatria, 2005, 81, 485-490.	0.9	2
168	Utilidade de um escore e de variÃ¡veis indicativas de drenagem pleural em crianÃ§as com derrame pleural parapneumÃ´nico. Jornal Brasileiro De Pneumologia, 2005, 31, 205-211.	0.4	3
169	Aerosols and spacers for acute asthma in children. Evolution and time to change the routine. Jornal De Pediatria, 2005, 81, 274-276.	0.9	0
170	AssociaÃ§Ã£o entre Ãndice de ventilaÃ§Ã£o e tempo de ventilaÃ§Ã£o mecÃ¢nica em lactentes com bronquiolite viral aguda. Jornal De Pediatria, 2005, 81, 466-470.	0.9	3
171	Cystic fibrosis at a Brazilian center of excellence: clinical and laboratory characteristics of 104 patients and their association with genotype and disease severity. Jornal De Pediatria, 2004, 80, 371-379.	0.9	38
172	Fibrose cÃstica em um centro de referÃªncia no Brasil: caracterÃsticas clÃnicas e laboratoriais de 104 pacientes e sua associaÃ§Ã£o com o genÃ³tipo e a gravidade da doenÃ§a. Jornal De Pediatria, 2004, 80, .	0.9	2
173	AnÃ¡lise crÃtica dos escores de avaliaÃ§Ã£o de gravidade da fibrose cÃstica: estado da arte. Jornal Brasileiro De Pneumologia, 2004, 30, 286-298.	0.4	30
174	Cystic fibrosis at a Brazilian center of excellence: clinical and laboratory characteristics of 104 patients and their association with genotype and disease severity. Jornal De Pediatria, 2004, 80, 371-9.	0.9	10
175	Linear growth in asthmatic children. Jornal De Pneumologia, 2003, 29, 36-42.	0.1	8
176	Fatores maternos e neonatais na incidÃªncia de displasia broncopulmonar em recÃ©m-nascidos de muito baixo peso. Jornal De Pediatria, 2003, 79, 550-556.	0.9	16
177	Maternal and neonatal factors affecting the incidence of bronchopulmonary dysplasia in very low birth weight newborns. Jornal De Pediatria, 2003, 79, 550-6.	0.9	2
178	Are immunoglobulin E levels associated with early wheezing? A prospective study in Brazilian infants. European Respiratory Journal, 2002, 20, 640-645.	3.1	35
179	Eosinophilic lung diseases. Paediatric Respiratory Reviews, 2002, 3, 278-284.	1.2	17
180	ControvÃ©rsias na fibrose cÃstica: do pediatra ao especialista. Jornal De Pediatria, 2002, 78, 171.	0.9	10

#	Article	IF	Citations
181	Abernethy malformation: One of the etiologies of hepatopulmonary syndrome. Pediatric Pulmonology, 2002, 34, 391-394.	1.0	101
182	Theophylline therapy inhibits neutrophil and mononuclear cell chemotaxis from chronic asthmatic children.. British Journal of Clinical Pharmacology, 1991, 32, 557-561.	1.1	33
183	Oral health in patients with cystic fibrosis. Brazilian Journal of Oral Sciences, 0, 17, e18160.	0.1	0