Bruna Ziegler Ziegler

List of Publications by Year
in descending order

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23 papers	259 citations	933447 10 h-index	940533 16 g-index
24 all docs	24 docs citations	24 times ranked	385 citing authors

#	Article	IF	Citations
1	Exercise programme in patients with cystic fibrosis: A randomized controlled trial. Respiratory Medicine, 2014, 108, 1134-1140.	2.9	33
2	Capacidade submÃ¡xima de exercÃcio em pacientes adolescentes e adultos com fibrose cÃstica. Jornal Brasileiro De Pneumologia, 2007, 33, 263-269.	0.7	29
3	PrevalÃªncia de hipertensÃ£o pulmonar avaliada por ecocardiografia Doppler em uma populaÃ§Ã£o de pacientes adolescentes e adultos com fibrose cÃstica. Jornal Brasileiro De Pneumologia, 2008, 34, 83-90.	0.7	25
4	Adherence to Airway Clearance Therapies by Adult Cystic Fibrosis Patients. Respiratory Care, 2013, 58, 279-285.	1.6	23
5	Glucose Intolerance in Patients With Cystic Fibrosis: Sex-Based Differences in Clinical Score, Pulmonary Function, Radiograph Score, and 6-Minute Walk Test. Respiratory Care, 2011, 56, 290-297.	1.6	19
6	Preditores da dessaturaÃ§Ã£o do oxigÃªnio no teste da caminhada de seis minutos em pacientes com fibrose cÃstica. Jornal Brasileiro De Pneumologia, 2009, 35, 957-965.	0.7	16
7	Repeatability of the 6-minute walk test in adolescents and adults with cystic fibrosis. Respiratory Care, 2010, 55, 1020-5.	1.6	16
8	Peripheral muscle strength is associated with lung function and functional capacity in patients with cystic fibrosis. Physiotherapy Research International, 2019, 24, e1771.	1.5	13
9	Doppler echocardiogram, oxygen saturation and submaximum capacity of exercise in patients with cystic fibrosis. Journal of Cystic Fibrosis, 2007, 6, 277-283.	0.7	12
10	Relationship between nutritional status and maximum inspiratory and expiratory pressures in cystic fibrosis. Respiratory Care, 2008, 53, 442-9.	1.6	12
11	Variability of the perception of dyspnea in healthy subjects assessed through inspiratory resistive loading. Jornal Brasileiro De Pneumologia, 2015, 41, 143-150.	0.7	8
12	Respiratory therapy: a problem among children and adolescents with cystic fibrosis. Jornal Brasileiro De Pneumologia, 2016, 42, 29-34.	0.7	8
13	Association between lung function, physical activity level and postural evaluation variables in adultÂpatients with cystic fibrosis. Clinical Respiratory Journal, 2018, 12, 1510-1517.	1.6	8
14	PadrÃµes ventilatÃ³rios na espirometria em pacientes adolescentes e adultos com fibrose cÃstica. Jornal Brasileiro De Pneumologia, 2009, 35, 854-859.	0.7	7
15	Pulmonary hypertension as estimated by Doppler echocardiography in adolescent and adult patients with cystic fibrosis and their relationship with clinical, lung function and sleep findings. Clinical Respiratory Journal, 2018, 12, 754-761.	1.6	7
16	Respiratory physical therapy techniques recommended for patients with cystic fibrosis treated in specialized centers. Brazilian Journal of Physical Therapy, 2020, 24, 532-538.	2.5	7
17	Functional capacity, pulmonary function, and quality of life in hematopoietic stem cell transplantation survivors. Supportive Care in Cancer, 2021, 29, 4015-4021.	2.2	5
18	Repeatability of the Evaluation of Perception of Dyspnea in Normal Subjects Assessed Through Inspiratory Resistive Loads. Open Respiratory Medicine Journal, 2014, 8, 41-47.	0.4	4

#	Article	IF	Citations
19	Clinical Outcomes and Prognostic Factors in a Cohort of Adults With Cystic Fibrosis: A 7-Year Follow-Up Study. Respiratory Care, 2016, 61, 192-199.	1.6	3
20	Repeatability of the 6-min walk test in non-cystic fibrosis bronchiectasis. Scientific Reports, 2020, 10, 19162.	3.3	1
21	Dyspnea perception during the inspiratory resistive loads test in obese subjects waiting bariatric surgery. Scientific Reports, 2020, 10, 8023.	3.3	1
22	Predictive factors for premature birth and respiratory exacerbation in pregnancies of women with cystic fibrosis. Jornal De Pediatria, 2021, , .	2.0	0
23	Physical activity and quality of life of children and adolescents with cystic fibrosis: a cross-sectional study. Fisioterapia Em Movimento, 0, 33, .	0.1	0