

# Bruna Ziegler Ziegler

## List of Publications by Year in descending order

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Version: 2024-02-01

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  | Functional capacity, pulmonary function, and quality of life in hematopoietic stem cell transplantation survivors. <i>Supportive Care in Cancer</i> , 2021, 29, 4015-4021.   | 2.2 | 5         |
| 2  | Predictive factors for premature birth and respiratory exacerbation in pregnancies of women with cystic fibrosis. <i>Jornal De Pediatria</i> , 2021, , .   | 2.0 | 0         |
| 3  | Respiratory physical therapy techniques recommended for patients with cystic fibrosis treated in specialized centers. <i>Brazilian Journal of Physical Therapy</i> , 2020, 24, 532-538.  | 2.5 | 7         |
| 4  | Repeatability of the 6-min walk test in non-cystic fibrosis bronchiectasis. <i>Scientific Reports</i> , 2020, 10, 19162.   | 3.3 | 1         |
| 5  | Dyspnea perception during the inspiratory resistive loads test in obese subjects waiting bariatric surgery. <i>Scientific Reports</i> , 2020, 10, 8023.  | 3.3 | 1         |
| 6  | Peripheral muscle strength is associated with lung function and functional capacity in patients with cystic fibrosis. <i>Physiotherapy Research International</i> , 2019, 24, e1771.   | 1.5 | 13        |
| 7  | 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. <i>Clinical Respiratory Journal</i> , 2018, 12, 754-761. | 1.6 | 7         |
| 8  | Association between lung function, physical activity level and postural evaluation variables in adult patients with cystic fibrosis. <i>Clinical Respiratory Journal</i> , 2018, 12, 1510-1517.  | 1.6 | 8         |
| 9  | Respiratory therapy: a problem among children and adolescents with cystic fibrosis. <i>Jornal Brasileiro De Pneumologia</i> , 2016, 42, 29-34.   | 0.7 | 8         |
| 10 | Clinical Outcomes and Prognostic Factors in a Cohort of Adults With Cystic Fibrosis: A 7-Year Follow-Up Study. <i>Respiratory Care</i> , 2016, 61, 192-199.  | 1.6 | 3         |
| 11 | Variability of the perception of dyspnea in healthy subjects assessed through inspiratory resistive loading. <i>Jornal Brasileiro De Pneumologia</i> , 2015, 41, 143-150.  | 0.7 | 8         |
| 12 | Repeatability of the Evaluation of Perception of Dyspnea in Normal Subjects Assessed Through Inspiratory Resistive Loads. <i>Open Respiratory Medicine Journal</i> , 2014, 8, 41-47.   | 0.4 | 4         |
| 13 | Exercise programme in patients with cystic fibrosis: A randomized controlled trial. <i>Respiratory Medicine</i> , 2014, 108, 1134-1140.  | 2.9 | 33        |
| 14 | Adherence to Airway Clearance Therapies by Adult Cystic Fibrosis Patients. <i>Respiratory Care</i> , 2013, 58, 279-285.  | 1.6 | 23        |
| 15 | Glucose Intolerance in Patients With Cystic Fibrosis: Sex-Based Differences in Clinical Score, Pulmonary Function, Radiograph Score, and 6-Minute Walk Test. <i>Respiratory Care</i> , 2011, 56, 290-297.  | 1.6 | 19        |
| 16 | Repeatability of the 6-minute walk test in adolescents and adults with cystic fibrosis. <i>Respiratory Care</i> , 2010, 55, 1020-5.  | 1.6 | 16        |
| 17 | Padrões ventilatórios na espirometria em pacientes adolescentes e adultos com fibrose cística. <i>Jornal Brasileiro De Pneumologia</i> , 2009, 35, 854-859.  | 0.7 | 7         |
| 18 | Preditores da dessaturação do oxigênio no teste da caminhada de seis minutos em pacientes com fibrose cística. <i>Jornal Brasileiro De Pneumologia</i> , 2009, 35, 957-965.  | 0.7 | 16        |

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|----|---|-----|-----------|
| 19 | Prevalência de hipertensão pulmonar avaliada por ecocardiografia Doppler em uma população de pacientes adolescentes e adultos com fibrose cística. Jornal Brasileiro De Pneumologia, 2008, 34, 83-90. | 0.7 | 25        |
| 20 | Relationship between nutritional status and maximum inspiratory and expiratory pressures in cystic fibrosis. Respiratory Care, 2008, 53, 442-9.   | 1.6 | 12        |
| 21 | 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        |
| 22 | Capacidade submáxima de exercício em pacientes adolescentes e adultos com fibrose cística. Jornal Brasileiro De Pneumologia, 2007, 33, 263-269.   | 0.7 | 29        |
| 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         |