

# Lucia De Martino

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/589838/publications.pdf>

Version: 2024-02-01

15  
papers

493  
citations

759233

12  
h-index

996975

15  
g-index

16  
all docs

16  
docs citations

16  
times ranked

695  
citing authors

| #  | ARTICLE   | IF  | CITATIONS |
|----|---|-----|-----------|
| 1  | Lung Ultrasound Score Predicts Surfactant Need in Extremely Preterm Neonates. <i>Pediatrics</i> , 2018, 142, .  | 2.1 | 173       |
| 2  | Comparative Evaluation of Therapy with <i>L</i> -Thyroxine versus No Treatment in Children with Idiopathic and Mild Subclinical Hypothyroidism. <i>Hormone Research in Paediatrics</i> , 2012, 77, 376-381.         | 1.8 | 63        |
| 3  | Autoimmune Polyendocrinopathy Candidiasis Ectodermal Dystrophy: Insights into Genotype-Phenotype Correlation. <i>International Journal of Endocrinology</i> , 2012, 2012, 1-9.                                      | 1.5 | 42        |
| 4  | APECED: A Paradigm of Complex Interactions between Genetic Background and Susceptibility Factors. <i>Frontiers in Immunology</i> , 2013, 4, 331.  | 4.8 | 32        |
| 5  | Porcine vs bovine surfactant therapy for preterm neonates with RDS: systematic review with biological plausibility and pragmatic meta-analysis of respiratory outcomes. <i>Respiratory Research</i> , 2019, 20, 28. | 3.6 | 32        |
| 6  | Novel Findings into AIRE Genetics and Functioning: Clinical Implications. <i>Frontiers in Pediatrics</i> , 2016, 4, 86.   | 1.9 | 25        |
| 7  | Autoimmune polyendocrinopathy-candidiasis-ectodermal dystrophy from the pediatric perspective. <i>Journal of Endocrinological Investigation</i> , 2013, 36, 903-12.   | 3.3 | 20        |
| 8  | Genetic Basis of Altered Central Tolerance and Autoimmune Diseases: A Lesson from AIRE Mutations. <i>International Reviews of Immunology</i> , 2012, 31, 344-362.   | 3.3 | 18        |
| 9  | Symptomatic malignant spinal cord compression in children: a single-center experience. <i>Italian Journal of Pediatrics</i> , 2019, 45, 80.   | 2.6 | 18        |
| 10 | Noonan-like syndrome with loose anagen hair associated with growth hormone insensitivity and atypical neurological manifestations. <i>American Journal of Medical Genetics, Part A</i> , 2012, 158A, 856-860.       | 1.2 | 16        |
| 11 | Long-term effects of growth hormone (GH) replacement therapy on hematopoiesis in a large cohort of children with GH deficiency. <i>Endocrine</i> , 2016, 53, 192-198.   | 2.3 | 15        |
| 12 | Germline mutations and new copy number variants among 40 pediatric cancer patients suspected for genetic predisposition. <i>Clinical Genetics</i> , 2019, 96, 359-365.  | 2.0 | 14        |
| 13 | Pediatric lung adenocarcinoma presenting with brain metastasis: a case report. <i>Journal of Medical Case Reports</i> , 2018, 12, 243.  | 0.8 | 11        |
| 14 | Investigation of chronic diarrhoea in infancy. <i>Early Human Development</i> , 2013, 89, 893-897.  | 1.8 | 10        |
| 15 | Acute adrenal insufficiency in a neonate with bilateral adrenal hemorrhage and combined prothrombotic risk factors. <i>Journal of Endocrinological Investigation</i> , 2012, 35, 449-449.                           | 3.3 | 4         |