

Armando Magrelli

List of Publications by Year in Descending Order

Source: <https://exaly.com/author-pdf/4806707/armando-magrelli-publications-by-year.pdf>

Version: 2024-04-10

This document has been generated based on the publications and citations recorded by exaly.com. For the latest version of this publication list, visit the link given above.

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

| | | | |
|-------------------|-------------------------|----------------|----------------|
| 28 papers | 1,275 citations | 13 h-index | 29 g-index |
| 29 ext. papers | 1,393 ext. citations | 8.2 avg, IF | 3.3 L-index |

| # | Paper | IF | Citations |
|----|---|------|-----------|
| 28 | Haemophilia, state of the art and new therapeutic opportunities, a regulatory perspective. <i>British Journal of Clinical Pharmacology</i> , 2021 , 87, 4183-4196 | 3.8 | 7 |
| 27 | Regulatory Standards in Orphan Medicinal Product Designation in the EU. <i>Frontiers in Medicine</i> , 2021 , 8, 698534 | 4.9 | 1 |
| 26 | Defining Satisfactory Methods of Treatment in Rare Diseases When Evaluating Significant Benefit-The EU Regulator's Perspective. <i>Frontiers in Medicine</i> , 2021 , 8, 744625 | 4.9 | 0 |
| 25 | Orphan Medicinal Products for the Treatment of Pancreatic Cancer: Lessons Learned From Two Decades of Orphan Designation.. <i>Frontiers in Oncology</i> , 2021 , 11, 809035 | 5.3 | |
| 24 | Reply to the article "Management of status epilepticus in adults. Position paper of the Italian League against Epilepsy". <i>Epilepsy and Behavior</i> , 2020 , 107, 106866 | 3.2 | 1 |
| 23 | Nonclinical data supporting orphan medicinal product designations in the area of rare infectious diseases. <i>Drug Discovery Today</i> , 2020 , 25, 274-291 | 8.8 | 4 |
| 22 | Reply to the reply of the authors of the review article entitled "Management of status epilepticus in adults. Position paper of the Italian League against Epilepsy". <i>Epilepsy and Behavior</i> , 2020 , 110, 107168 ^{3,2} | | |
| 21 | Using a stated preference discrete choice experiment to assess societal value from the perspective of patients with rare diseases in Italy. <i>Orphanet Journal of Rare Diseases</i> , 2019 , 14, 154 | 4.2 | 8 |
| 20 | European regulators' views on a wearable-derived performance measurement of ambulation for Duchenne muscular dystrophy regulatory trials. <i>Neuromuscular Disorders</i> , 2019 , 29, 514-516 | 2.9 | 27 |
| 19 | Defining orphan conditions in the context of the European orphan regulation: challenges and evolution. <i>Nature Reviews Drug Discovery</i> , 2019 , 18, 479-480 | 64.1 | 5 |
| 18 | Nonclinical data supporting orphan medicinal product designations: lessons from rare neurological conditions. <i>Drug Discovery Today</i> , 2018 , 23, 26-48 | 8.8 | 10 |
| 17 | Demonstrating significant benefit of orphan medicines: analysis of 15 years of experience in Europe. <i>Drug Discovery Today</i> , 2018 , 23, 90-100 | 8.8 | 10 |
| 16 | Pathways Implicated in Tadalafil Amelioration of Duchenne Muscular Dystrophy. <i>Journal of Cellular Physiology</i> , 2016 , 231, 224-32 | 7 | 19 |
| 15 | The Italian National Centre for Rare Diseases: where research and public health translate into action. <i>Blood Transfusion</i> , 2014 , 12 Suppl 3, s591-605 | 3.6 | 1 |
| 14 | Predictive medicine and biomarkers: the case of rare diseases. <i>Personalized Medicine</i> , 2012 , 9, 143-146 | 2.2 | 2 |
| 13 | Oxidative stress activation of miR-125b is part of the molecular switch for Hailey-Hailey disease manifestation. <i>Experimental Dermatology</i> , 2011 , 20, 932-7 | 4 | 50 |
| 12 | The role of microRNAs in the biology of rare diseases. <i>International Journal of Molecular Sciences</i> , 2011 , 12, 6733-42 | 6.3 | 11 |

| | | | |
|----|--|------|-----|
| 11 | Complex multipathways alterations and oxidative stress are associated with Hailey-Hailey disease. <i>British Journal of Dermatology</i> , 2010 , 162, 518-26 | 4 | 28 |
| 10 | MicroRNA profiling of multiple osteochondromas: identification of disease-specific and normal cartilage signatures. <i>Clinical Genetics</i> , 2010 , 78, 507-16 | 4 | 29 |
| 9 | The artificial gene Jazz, a transcriptional regulator of utrophin, corrects the dystrophic pathology in mdx mice. <i>Human Molecular Genetics</i> , 2010 , 19, 752-60 | 5.6 | 28 |
| 8 | In utero exposure to di-(2-ethylhexyl) phthalate affects liver morphology and metabolism in post-natal CD-1 mice. <i>Reproductive Toxicology</i> , 2010 , 29, 427-32 | 3.4 | 40 |
| 7 | Identification of key regions and genes important in the pathogenesis of Sezary syndrome by combining genomic and expression microarrays. <i>Cancer Research</i> , 2009 , 69, 8438-46 | 10.1 | 62 |
| 6 | Altered microRNA Expression Patterns in Hepatoblastoma Patients. <i>Translational Oncology</i> , 2009 , 2, 157-63 | 4.9 | 59 |
| 5 | Quantitative technologies establish a novel microRNA profile of chronic lymphocytic leukemia. <i>Blood</i> , 2007 , 109, 4944-51 | 2.2 | 422 |
| 4 | Altered dopamine signaling and MPTP resistance in mice lacking the Parkinson's disease-associated GPR37/parkin-associated endothelin-like receptor. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2004 , 101, 10189-94 | 11.5 | 63 |
| 3 | Genomic Analysis of GPR37 and Related Orphan G-Protein Coupled Receptor Genes Highly Expressed in the Mammalian Brain. <i>Current Genomics</i> , 2001 , 2, 253-260 | 2.6 | 5 |
| 2 | White collar-1, a central regulator of blue light responses in <i>Neurospora</i> , is a zinc finger protein.. <i>EMBO Journal</i> , 1996 , 15, 1650-1657 | 13 | 350 |
| 1 | Splicing of the rolA transcript of <i>Agrobacterium rhizogenes</i> in <i>Arabidopsis</i> . <i>Science</i> , 1994 , 266, 1986-8 | 33.3 | 33 |