## Armando Magrelli

## List of Publications by Citations

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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
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h-index
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1,393
ext. papers
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avg, IF
L-index

#	Paper	IF	Citations
28	Quantitative technologies establish a novel microRNA profile of chronic lymphocytic leukemia. <i>Blood</i> , <b>2007</b> , 109, 4944-51	2.2	422
27	White collar-1, a central regulator of blue light responses in Neurospora, is a zinc finger protein <i>EMBO Journal</i> , <b>1996</b> , 15, 1650-1657	13	350
26	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> , <b>2004</b> , 101, 10189-94	11.5	63
25	Identification of key regions and genes important in the pathogenesis of sezary syndrome by combining genomic and expression microarrays. <i>Cancer Research</i> , <b>2009</b> , 69, 8438-46	10.1	62
24	Altered microRNA Expression Patterns in Hepatoblastoma Patients. <i>Translational Oncology</i> , <b>2009</b> , 2, 157-63	4.9	59
23	Oxidative stress activation of miR-125b is part of the molecular switch for Hailey-Hailey disease manifestation. <i>Experimental Dermatology</i> , <b>2011</b> , 20, 932-7	4	50
22	In utero exposure to di-(2-ethylhexyl) phthalate affects liver morphology and metabolism in post-natal CD-1 mice. <i>Reproductive Toxicology</i> , <b>2010</b> , 29, 427-32	3.4	40
21	Splicing of the rolA transcript of Agrobacterium rhizogenes in Arabidopsis. <i>Science</i> , <b>1994</b> , 266, 1986-8	33.3	33
20	MicroRNA profiling of multiple osteochondromas: identification of disease-specific and normal cartilage signatures. <i>Clinical Genetics</i> , <b>2010</b> , 78, 507-16	4	29
19	Complex multipathways alterations and oxidative stress are associated with Hailey-Hailey disease. <i>British Journal of Dermatology</i> , <b>2010</b> , 162, 518-26	4	28
18	The artificial gene Jazz, a transcriptional regulator of utrophin, corrects the dystrophic pathology in mdx mice. <i>Human Molecular Genetics</i> , <b>2010</b> , 19, 752-60	5.6	28
17	European regulatorsTviews on a wearable-derived performance measurement of ambulation for Duchenne muscular dystrophy regulatory trials. <i>Neuromuscular Disorders</i> , <b>2019</b> , 29, 514-516	2.9	27
16	Pathways Implicated in Tadalafil Amelioration of Duchenne Muscular Dystrophy. <i>Journal of Cellular Physiology</i> , <b>2016</b> , 231, 224-32	7	19
15	The role of microRNAs in the biology of rare diseases. <i>International Journal of Molecular Sciences</i> , <b>2011</b> , 12, 6733-42	6.3	11
14	Nonclinical data supporting orphan medicinal product designations: lessons from rare neurological conditions. <i>Drug Discovery Today</i> , <b>2018</b> , 23, 26-48	8.8	10
13	Demonstrating significant benefit of orphan medicines: analysis of 15 years of experience in Europe. <i>Drug Discovery Today</i> , <b>2018</b> , 23, 90-100	8.8	10
12	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> , <b>2019</b> , 14, 154	4.2	8

## LIST OF PUBLICATIONS

11	Haemophilia, state of the art and new therapeutic opportunities, a regulatory perspective. <i>British Journal of Clinical Pharmacology</i> , <b>2021</b> , 87, 4183-4196	3.8	7
10	Genomic Analysis of GPR37 and Related Orphan G-Protein Coupled Receptor Genes Highly Expressed in the Mammalian Brain. <i>Current Genomics</i> , <b>2001</b> , 2, 253-260	2.6	5
9	Defining orphan conditions in the context of the European orphan regulation: challenges and evolution. <i>Nature Reviews Drug Discovery</i> , <b>2019</b> , 18, 479-480	64.1	5
8	Nonclinical data supporting orphan medicinal product designations in the area of rare infectious diseases. <i>Drug Discovery Today</i> , <b>2020</b> , 25, 274-291	8.8	4
7	Predictive medicine and biomarkers: the case of rare diseases. <i>Personalized Medicine</i> , <b>2012</b> , 9, 143-146	2.2	2
6	Reply to the article "Management of status epilepticus in adults. Position paper of the Italian League against Epilepsy". <i>Epilepsy and Behavior</i> , <b>2020</b> , 107, 106866	3.2	1
5	The Italian National Centre for Rare Diseases: where research and public health translate into action. <i>Blood Transfusion</i> , <b>2014</b> , 12 Suppl 3, s591-605	3.6	1
4	Regulatory Standards in Orphan Medicinal Product Designation in the EU. <i>Frontiers in Medicine</i> , <b>2021</b> , 8, 698534	4.9	1
3	Defining Satisfactory Methods of Treatment in Rare Diseases When Evaluating Significant Benefit-The EU Regulator's Perspective. <i>Frontiers in Medicine</i> , <b>2021</b> , 8, 744625	4.9	О
2	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> , <b>2020</b> , 110, 10716	58 <sup>.2</sup>	
1	Orphan Medicinal Products for the Treatment of Pancreatic Cancer: Lessons Learned From Two Decades of Orphan Designation <i>Frontiers in Oncology</i> , <b>2021</b> , 11, 809035	5.3	