## Patrizia Suppressa

## List of Publications by Year in Descending Order

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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.

15	311	8	17
papers	citations	h-index	g-index
17	384 ext. citations	3.9	2.5
ext. papers		avg, IF	L-index

#	Paper	IF	Citations
15	Abdominal obesity negatively influences key metrics of reverse cholesterol transport. <i>Biochimica Et Biophysica Acta - Molecular and Cell Biology of Lipids</i> , <b>2021</b> , 1867, 159087	5	О
14	AST to Platelet Ratio Index (APRI) is an easy-to-use predictor score for cardiovascular risk in metabolic subjects. <i>Scientific Reports</i> , <b>2021</b> , 11, 14834	4.9	7
13	Minimal portosystemic encephalopathy: A new nosological entity in patients with hereditary haemorrhagic telangiectasia. <i>European Journal of Internal Medicine</i> , <b>2021</b> , 90, 43-48	3.9	
12	Characterization of epidemiological distribution and outcome of COVID-19 in patients with hereditary hemorrhagic telangiectasia: a nationwide retrospective multi-centre study during first wave in Italy. <i>Orphanet Journal of Rare Diseases</i> , <b>2021</b> , 16, 378	4.2	1
11	European Reference Network for Rare Vascular Diseases (VASCERN) position statement on cerebral screening in adults and children with hereditary haemorrhagic telangiectasia (HHT). <i>Orphanet Journal of Rare Diseases</i> , <b>2020</b> , 15, 165	4.2	14
10	Gender, BMI and fasting hyperglycaemia influence Monocyte to-HDL ratio (MHR) index in metabolic subjects. <i>PLoS ONE</i> , <b>2020</b> , 15, e0231927	3.7	4
9	Safety of thalidomide and bevacizumab in patients with hereditary hemorrhagic telangiectasia. <i>Orphanet Journal of Rare Diseases</i> , <b>2019</b> , 14, 28	4.2	45
8	Efficacy of Lomitapide in the Treatment of Familial Homozygous Hypercholesterolemia: Results of a Real-World Clinical Experience in Italy. <i>Advances in Therapy</i> , <b>2017</b> , 34, 1200-1210	4.1	39
7	A case of sarcoidosis with isolated hepatosplenic onset and development of inflammatory bowel disease during recovery stage. <i>Autoimmunity Highlights</i> , <b>2017</b> , 8, 6	3.7	
6	Hepatic angiodynamic profile in paediatric patients with hereditary haemorrhagic telangiectasia type 1 and type 2. <i>Vasa - European Journal of Vascular Medicine</i> , <b>2017</b> , 46, 195-202	1.9	4
5	Liver involvement in hereditary hemorrhagic telangiectasia: can breath test unmask impaired hepatic first-pass effect?. <i>Internal and Emergency Medicine</i> , <b>2012</b> , 7, 323-9	3.7	8
4	Low dose intravenous bevacizumab for the treatment of anaemia in hereditary haemorrhagic telangiectasia. <i>British Journal of Haematology</i> , <b>2011</b> , 152, 365	4.5	15
3	Liver involvement in a large cohort of patients with hereditary hemorrhagic telangiectasia: echo-color-Doppler vs multislice computed tomography study. <i>Journal of Hepatology</i> , <b>2008</b> , 48, 811-20	13.4	68
2	DHPLC-based mutation analysis of ENG and ALK-1 genes in HHT Italian population. <i>Human Mutation</i> , <b>2006</b> , 27, 213-4	4.7	32
1	Health-related quality of life in a rare disease: hereditary hemorrhagic telangiectasia (HHT) or Rendu-Osler-Weber disease. <i>Quality of Life Research</i> , <b>2004</b> , 13, 1715-23	3.7	74