

Patrizia Suppressa

List of Publications by Year in descending order

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Version: 2024-02-01

17
papers

457
citations

1170033

9
h-index

1113639

15
g-index

17
all docs

17
docs citations

17
times ranked

599
citing authors

| # | ARTICLE | IF | CITATIONS |
|----|--|-----|-----------|
| 1 | Safety of reduced or absent antithrombotic therapy after left atrial appendage closure in patients affected by hereditary hemorrhagic telangiectasia and atrial fibrillation. <i>Minerva Cardiology and Angiology</i> , 2022, 70, . | 0.4 | 5 |
| 2 | Abdominal obesity negatively influences key metrics of reverse cholesterol transport. <i>Biochimica Et Biophysica Acta - Molecular and Cell Biology of Lipids</i> , 2022, 1867, 159087. | 1.2 | 7 |
| 3 | Monocyte-to-HDL Ratio (MHR) Predicts Vitamin D Deficiency in Healthy and Metabolic Women: A Cross-Sectional Study in 1048 Subjects. <i>Nutrients</i> , 2022, 14, 347. | 1.7 | 9 |
| 4 | AST to Platelet Ratio Index (APRI) is an easy-to-use predictor score for cardiovascular risk in metabolic subjects. <i>Scientific Reports</i> , 2021, 11, 14834. | 1.6 | 24 |
| 5 | Minimal portosystemic encephalopathy: A new nosological entity in patients with hereditary haemorrhagic telangiectasia. <i>European Journal of Internal Medicine</i> , 2021, 90, 43-48. | 1.0 | 0 |
| 6 | 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> , 2021, 16, 378. | 1.2 | 4 |
| 7 | 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> , 2020, 15, 165. | 1.2 | 28 |
| 8 | Gender, BMI and fasting hyperglycaemia influence Monocyte to-HDL ratio (MHR) index in metabolic subjects. <i>PLoS ONE</i> , 2020, 15, e0231927. | 1.1 | 14 |
| 9 | Safety of thalidomide and bevacizumab in patients with hereditary hemorrhagic telangiectasia. <i>Orphanet Journal of Rare Diseases</i> , 2019, 14, 28. | 1.2 | 75 |
| 10 | Efficacy of Lomitapide in the Treatment of Familial Homozygous Hypercholesterolemia: Results of a Real-World Clinical Experience in Italy. <i>Advances in Therapy</i> , 2017, 34, 1200-1210. | 1.3 | 56 |
| 11 | A case of sarcoidosis with isolated hepatosplenic onset and development of inflammatory bowel disease during recovery stage. <i>Autoimmunity Highlights</i> , 2017, 8, 6. | 3.9 | 0 |
| 12 | Hepatic angiodynamic profile in paediatric patients with hereditary haemorrhagic telangiectasia type 1 and type 2. <i>Vasa - European Journal of Vascular Medicine</i> , 2017, 46, 195-202. | 0.6 | 4 |
| 13 | Liver involvement in hereditary hemorrhagic telangiectasia: can breath test unmask impaired hepatic first-pass effect?. <i>Internal and Emergency Medicine</i> , 2012, 7, 323-329. | 1.0 | 9 |
| 14 | Low dose intravenous bevacizumab for the treatment of anaemia in hereditary haemorrhagic telangiectasia. <i>British Journal of Haematology</i> , 2011, 152, 365-365. | 1.2 | 19 |
| 15 | 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> , 2008, 48, 811-820. | 1.8 | 71 |
| 16 | DHPLC-based mutation analysis of ENG and ALK-1 genes in HHT Italian population. <i>Human Mutation</i> , 2006, 27, 213-214. | 1.1 | 38 |
| 17 | Health-related quality of life in a rare disease: Hereditary hemorrhagic telangiectasia (HHT) or Rendu-Osler-Weber Disease. <i>Quality of Life Research</i> , 2004, 13, 1715-1723. | 1.5 | 94 |