

Elisabetta Buscarini

List of Publications by Year in descending order

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Version: 2024-02-01

35
papers

3,361
citations

361413

20
h-index

414414

32
g-index

37
all docs

37
docs citations

37
times ranked

3292
citing authors

#	ARTICLE	IF	CITATIONS
1	Dysmetabolism, Diabetes and Clinical Outcomes in Patients Cured of Chronic Hepatitis C: A Real-Life Cohort Study. <i>Hepatology Communications</i> , 2022, 6, 867-877.	4.3	6
2	Hepatic Vascular Malformations in Hereditary Hemorrhagic Telangiectasia. , 2022, , 49-68.		0
3	Changes in digestive cancer diagnosis during the SARS-CoV-2 pandemic in Italy: A nationwide survey. <i>Digestive and Liver Disease</i> , 2021, 53, 682-688.	0.9	30
4	Second-generation thrombopoietin receptor agonists: New players in the management of cirrhotic patients undergoing therapeutic endoscopy?. <i>Digestive and Liver Disease</i> , 2021, 53, 1362-1363.	0.9	1
5	Needle-based confocal endomicroscopy in the discrimination of mucinous from non-mucinous pancreatic cystic lesions. <i>World Journal of Gastrointestinal Endoscopy</i> , 2021, 13, 555-564.	1.2	5
6	Dietary iron intake and anemia: food frequency questionnaire in patients with hereditary hemorrhagic telangiectasia. <i>Orphanet Journal of Rare Diseases</i> , 2020, 15, 295.	2.7	2
7	GI symptoms as early signs of COVID-19 in hospitalised Italian patients. <i>Gut</i> , 2020, 69, 1547-1548.	12.1	50
8	Multicentric Italian survey on daily practice for autoimmune pancreatitis: Clinical data, diagnosis, treatment, and evolution toward pancreatic insufficiency. <i>United European Gastroenterology Journal</i> , 2020, 8, 705-715.	3.8	25
9	High rates of 30-day mortality in patients with cirrhosis and COVID-19. <i>Journal of Hepatology</i> , 2020, 73, 1063-1071.	3.7	279
10	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.	2.7	28
11	Hereditary hemorrhagic telangiectasia and liver involvement. <i>Clinics and Research in Hepatology and Gastroenterology</i> , 2020, 44, 426-432.	1.5	5
12	Laboratory markers included in the Corona Score can identify false negative results on COVID-19 RT-PCR in the emergency room. <i>Biochimica Medica</i> , 2020, 30, 357-359.	2.7	7
13	Hospital admission for digestive diseases: Gastroenterology units offer a more effective and efficient care. <i>Digestive and Liver Disease</i> , 2019, 51, 43-46.	0.9	3
14	Prevention of serious infections in hereditary hemorrhagic telangiectasia: roles for prophylactic antibiotics, the pulmonary capillaries-but not vaccination. <i>Haematologica</i> , 2019, 104, e85-e86.	3.5	11
15	Safety of thalidomide and bevacizumab in patients with hereditary hemorrhagic telangiectasia. <i>Orphanet Journal of Rare Diseases</i> , 2019, 14, 28.	2.7	75
16	Liver involvement in hereditary hemorrhagic telangiectasia. <i>Abdominal Radiology</i> , 2018, 43, 1920-1930.	2.1	26
17	European Reference Network For Rare Vascular Diseases (VASCERN) Outcome Measures For Hereditary Haemorrhagic Telangiectasia (HHT). <i>Orphanet Journal of Rare Diseases</i> , 2018, 13, 136.	2.7	74
18	Long-term outcomes of patients with pulmonary arteriovenous malformations considered for lung transplantation, compared with similarly hypoxaemic cohorts. <i>BMJ Open Respiratory Research</i> , 2017, 4, e000198.	3.0	9

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19	Response to Bevacizumab for the treatment of Rendu-Osler disease”A note of caution. Liver International, 2017, 37, 928-928.	3.9	2
20	Evidence for the Presence of Non-Celiac Gluten Sensitivity in Patients with Functional Gastrointestinal Symptoms: Results from a Multicenter Randomized Double-Blind Placebo-Controlled Gluten Challenge. Nutrients, 2016, 8, 84.	4.1	155
21	Hereditary hemorrhagic telangiectasia: to transplant or not to transplant?. Liver International, 2016, 36, 1741-1744.	3.9	12
22	Hospital care services for digestive diseases in Italy: The first quantitative assessment. Digestive and Liver Disease, 2014, 46, 652-657.	0.9	6
23	White Paper of Italian Gastroenterology: Delivery of services for digestive diseases in Italy: Weaknesses and strengths. Digestive and Liver Disease, 2014, 46, 579-589.	0.9	40
24	Natural History and Outcome of Hepatic Vascular Malformations in a Large Cohort of Patients with Hereditary Hemorrhagic Teleangiectasia. Digestive Diseases and Sciences, 2011, 56, 2166-2178.	2.3	106
25	Endoscopic ultrasonography findings in autoimmune pancreatitis. World Journal of Gastroenterology, 2011, 17, 2080.	3.3	40
26	Interobserver Agreement in Diagnosing Liver Involvement in Hereditary Hemorrhagic Telangiectasia by Doppler Ultrasound. Ultrasound in Medicine and Biology, 2008, 34, 718-725.	1.5	21
27	Genotype-phenotype correlations in hereditary hemorrhagic telangiectasia: Data from the French-Italian HHT network. Genetics in Medicine, 2007, 9, 14-22.	2.4	196
28	Persistent elastic band in esophagus: An innocent bystander?. Digestive Endoscopy, 2006, 18, 239-239.	2.3	0
29	Radiofrequency thermal ablation with expandable needle of focal liver malignancies: complication report. European Radiology, 2004, 14, 31-37.	4.5	82
30	High prevalence of hepatic focal nodular hyperplasia in subjects with hereditary hemorrhagic telangiectasia. Ultrasound in Medicine and Biology, 2004, 30, 1089-1097.	1.5	117
31	EUS for suspected choledocholithiasis: Do benefits outweigh costs? A prospective, controlled study. Gastrointestinal Endoscopy, 2003, 57, 510-518.	1.0	141
32	Percutaneous radiofrequency ablation of small hepatocellular carcinoma: long-term results. European Radiology, 2001, 11, 914-921.	4.5	305
33	Diagnostic criteria for hereditary hemorrhagic telangiectasia (Rendu-Osler-Weber syndrome). American Journal of Medical Genetics Part A, 2000, 91, 66-67.	2.4	1,391
34	Diagnostic criteria for hereditary hemorrhagic telangiectasia (Rendu-Osler-Weber syndrome). , 2000, 91, 66.		3
35	Hepatic vascular malformations in hereditary hemorrhagic telangiectasia: Doppler sonographic screening in a large family. Journal of Hepatology, 1997, 26, 111-118.	3.7	106