Carla Olivieri

List of Publications by Year
in descending order

41 papers	1,416 citations	361413 20 h-index	330143 37 g-index
41 all docs	41 docs citations	41 times ranked	1386 citing authors

#	Article	IF	Citations
1	Genotype-phenotype correlations in hereditary hemorrhagic telangiectasia: Data from the French-Italian HHT network. Genetics in Medicine, 2007, 9, 14-22.	2.4	196
2	Overlapping spectra of SMAD4 mutations in juvenile polyposis (JP) and JPâ€“HHT syndrome. American Journal of Medical Genetics, Part A, 2010, 152A, 333-339.	1.2	128
3	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
4	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
5	Doppler Ultrasonographic Grading of Hepatic Vascular Malformations in Hereditary Hemorrhagic Telangiectasia - Results of Extensive Screening. Ultraschall in Der Medizin, 2004, 25, 348-355.	1.5	98
6	Contrast echocardiography for pulmonary arteriovenous malformations screening: does any bubble matter?. European Journal of Echocardiography, 2009, 10, 513-518.	2.3	79
7	Analysis of ENG and ACVRL1 genes in 137 HHT Italian families identifies 76 different mutations (24 novel). Comparison with other European studies. Journal of Human Genetics, 2007, 52, 820-829.	2.3	63
8	Efficacy and safety of thalidomide for the treatment of severe recurrent epistaxis in hereditary haemorrhagic telangiectasia: results of a non-randomised, single-centre, phase 2 study. Lancet Haematology,the, 2015, 2, e465-e473.	4.6	57
9	Echocardiographic screening discloses increased values of pulmonary artery systolic pressure in 9 of 68 unselected patients affected with hereditary hemorrhagic telangiectasia. Genetics in Medicine, 2006, 8, 183-190.	2.4	52
10	Liver involvement in hereditary haemorrhagic telangiectasia or Rendu-Osler-Weber disease. Digestive and Liver Disease, 2005, 37, 635-645.	0.9	51
11	Identification of 13 new mutations in the ACVRL1 gene in a group of 52 unselected Italian patients affected by hereditary haemorrhagic telangiectasia. Journal of Medical Genetics, 2002, 39, 39e-39.	3.2	49
12	Endoscopic evaluation of gastrointestinal tract in patients with hereditary hemorrhagic telangiectasia and correlation with their genotypes. Genetics in Medicine, 2014, 16, 3-10.	2.4	47
13	Familial partial monosomy 7 and myelodysplasia. Cancer Genetics and Cytogenetics, 2001, 124, 147-151.	1.0	38
14	Hereditary hemorrhagic telangiectasia: evidence for regional founder effects of ACVRL1 mutations in French and Italian patients. European Journal of Human Genetics, 2008, 16, 742-749.	2.8	35
15	Epidemiology and clinical aspects of Werner's syndrome in North Sardinia: description of a cluster. European Journal of Dermatology, 2007, 17, 213-6.	0.6	34
16	Isochromosome (7)(q10) in Shwachman Syndrome Without MDS/AML and Role of Chromosome 7 Anomalies in Myeloproliferative Disorders. Cancer Genetics and Cytogenetics, 2000, 121, 167-171.	1.0	32
17	Interstitial deletion of chromosomeÂ9, int del(9)(9q22.31-q31.2), including the genes causing multiple basal cell nevus syndrome and Robinow/brachydactylyÂ1 syndrome. European Journal of Pediatrics, 2003, 162, 100-103.	2.7	31
18	Shwachmanâ€Ðiamond syndrome with clonal interstitial deletion of the long arm of chromosome 20 in bone marrow: haematological features, prognosis and genomic instability. British Journal of Haematology, 2019, 184, 974-981.	2.5	24

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19	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
20	HHT diagnosis by Mid-infrared spectroscopy and artificial neural network analysis. Orphanet Journal of Rare Diseases, 2013, 8, 94.	2.7	20
21	Early Onset of Gastric Carcinoma and Constitutional Deletion of 18p. Cancer Genetics and Cytogenetics, 1999, 113, 96-99.	1.0	18
22	Correlation of Severity of Epistaxis with Nasal Telangiectasias in Hereditary Hemorrhagic Telangiectasia (HHT) Patients. American Journal of Rhinology and Allergy, 2009, 23, 52-58.	2.0	18
23	Argon plasma coagulation is an effective treatment for hereditary hemorrhagic telangiectasia patients with severe nosebleeds. Acta Oto-Laryngologica, 2013, 133, 174-180.	0.9	17
24	Bioinformatic Analysis of Pathogenic Missense Mutations of Activin Receptor Like Kinase 1 Ectodomain. PLoS ONE, 2011, 6, e26431.	2.5	14
25	Treatment of Epistaxis in Hereditary Hemorrhagic Telangiectasia Patients by Argon Plasma Coagulation with Local Anesthesia. American Journal of Rhinology & Allergy, 2006, 20, 421-425.	2.2	11
26	Meiotic origin of trisomy in neoplasms: evidence in a case of erythroleukaemia. Leukemia, 2001, 15, 971-975.	7.2	9
27	Functional analysis of a novel ENG variant in a patient with hereditary hemorrhagic telangiectasia (HHT) identifies a new Sp1 binding-site. Gene, 2018, 647, 85-92.	2.2	9
28	Endoscopic surgical treatment of epistaxis in hereditary haemorrhagic telangiectasia: our experience. Acta Otorhinolaryngologica Italica, 2021, 41, 59-68.	1.5	6
29	Fluoresceinâ€guided intraoperative endoscopy in patients with hereditary hemorrhagic telangiectasia: first impressions. International Forum of Allergy and Rhinology, 2017, 7, 300-303.	2.8	5
30	Characterization of a mutation in the zona pellucida module of Endoglin that causes Hereditary Hemorrhagic Telangiectasia. Gene, 2019, 696, 33-39.	2.2	5
31	Vascular abnormalities in the fingers of patients affected with hereditary hemorrhagic telangiectasia (HHT) as assessed by color doppler sonography. American Journal of Medical Genetics, Part A, 2005, 135A, 106-109.	1.2	4
32	Different forms of pulmonary hypertension in a family with clinical and genetic evidence for hereditary hemorrhagic teleangectasia type 2. Pulmonary Circulation, 2018, 8, 1-4.	1.7	4
33	BMPR2Âmutations and response to inhaled or parenteral prostanoids: a case series. Pulmonary Circulation, 2021, 11, 1-5.	1.7	4
34	Increase of circulating endothelial cells in patients with Hereditary Hemorrhagic Telangiectasia. International Journal of Hematology, 2015, 101, 23-31.	1.6	3
35	Nasal Endoscopy in the Clinical Diagnosis of Hereditary Hemorrhagic Telangiectasia. Journal of Pediatrics, 2021, 238, 74-79.e2.	1.8	3
36	Hereditary hemorrhagic telangiectasia: First demonstration of a founder effect in Italy; the ACVRL1 c.289_294del variant originated in the country of Bergamo 200â€‰years ago. Molecular Genetics & Genomic Medicine, 0, , .	1.2	3

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37	Hereditary haemorrhagic telangiectasia in North African and sub-Saharan patients. South African Medical Journal, 2014, 104, 256.	0.6	2
38	Correspondence. Cancer Genetics and Cytogenetics, 1999, 110, 140-142.	1.0	1
39	Hereditary Hemorrhagic Telangiectasia: Breakpoint Characterization of a Novel Large Deletion in ACVRL1 Suggests the Causing Mechanism. Molecular Syndromology, 2013, 4, 119-124.	0.8	1
40	Efficacy of Thalidomide in the Treatment of Severe Recurrent Epistaxis in Hereditary Hemorrhagic Telangiectasia (HHT): Preliminary Results of an Ongoing Study. Blood, 2012, 120, 629-629.	1.4	1
41	Immunohistochemical analysis of a merkeloma observed in a patient affected by hereditary haemorrhagic telangiectasia. BMJ Case Reports, 2010, 2010, bcr0920092251-bcr0920092251.	0.5	0