Serge Pissard

List of Publications by Year
in descending order

17 papers	95 citations	1937685 4 h-index	1372567 10 g-index
17 all docs	17 docs citations	17 times ranked	208 citing authors

#	Article	IF	Citations
1	Hb Angers: A new Î±2â€globin variant [Î±2 (140)(HC2) TyrÂâ†'ÂSer; HBA2: C.422 A>C] with increased oxygen affinity leading to erythrocytosis. International Journal of Laboratory Hematology, 2021, 43, e114-e117.	1.3	0
2	Le dÃ©ficit en G6PD. Journal De Pediatrie Et De Puericulture, 2021, 34, 244-250.	0.0	0
3	A new gene associated with a Î²-thalassemia phenotype: the observation of variants in SUPT5H. Blood, 2020, 136, 1789-1793.	1.4	13
4	Red Blood Cells Free Î± Hemoglobin Pool: A Biomarker to Monitor Imbalanced Î±/Non Î±-Globin Chain Synthesis in Î²-Thalassemia Intermedia: The ALPHAPOOL Study. Hemoglobin, 2019, 43, 344-344.	0.8	0
5	Molecular Approach to Prenatal Diagnosis of Hemoglobinopathies and Thalassemias. Hemoglobin, 2019, 43, 335-335.	0.8	0
6	Short in-Frame Insertions/Deletions in the Coding Sequence of the Î±-Globin Gene. Consequences of the 3D Structure and Resulting Phenotypes: Hb Choisy as an Example. Hemoglobin, 2018, 42, 287-293.	0.8	1
7	Inherited or acquired modifiers of iron status may dramatically affect the phenotype in dehydrated hereditary stomatocytosis. European Journal of Haematology, 2018, 101, 566-569.	2.2	9
8	Evidence for a gene conversion in a Hb Arya Carrier [Î± codon 47 Asp>Asn, Hb A1(or Hb A2):c.142 G>A]. International Journal of Laboratory Hematology, 2017, 39, e55-e59.	1.3	2
9	Maternal consumption of quinine-containing sodas may induce G6PD crises in breastfed children. European Journal of Pediatrics, 2017, 176, 1415-1418.	2.7	9
10	Hb Olivet (HBA1: C.40Gâ€‰>â€‰A; p.Ala14Thr), a Novel Silent Hemoglobin Variant in Two Families of Distinct Origin. Hemoglobin, 2016, 40, 349-352.	0.8	1
11	Hb Savaria [Î±49(CE7)Serâ†'Arg;HBA2: c.150Câ€‰>â€‰A]: A New Case and Complete Description. Hemoglobin, 2016, 40, 267-269.	0.8	2
12	Haematological determinants of cardiac involvement in adults with sickle cell disease. European Heart Journal, 2016, 37, 1158-1167.	2.2	45
13	Improvements in phenotype studies of hemoglobin disorders brought by advances in reversedâ€phase chromatography of globin chains. International Journal of Laboratory Hematology, 2015, 37, 279-286.	1.3	5
14	A Second Observation of the Rare Frameshift Mutation in the Î²-Globin Gene: Codon 46 (+A) (Hbb:c.138_139insA). Hemoglobin, 2011, 35, 157-161.	0.8	2
15	Analysis of Rare deletional Thalassemia Using Custom CGH Array DNA Chip. Blood, 2010, 116, 4278-4278.	1.4	2
16	HFE Genotyping by Amplification Refractory Mutation Systemâ€"Denaturing HPLC. Clinical Chemistry, 2002, 48, 769-772.	3.2	3
17	HFE genotyping by amplification refractory mutation system-denaturing HPLC. Clinical Chemistry, 2002, 48, 769-72.	3.2	1