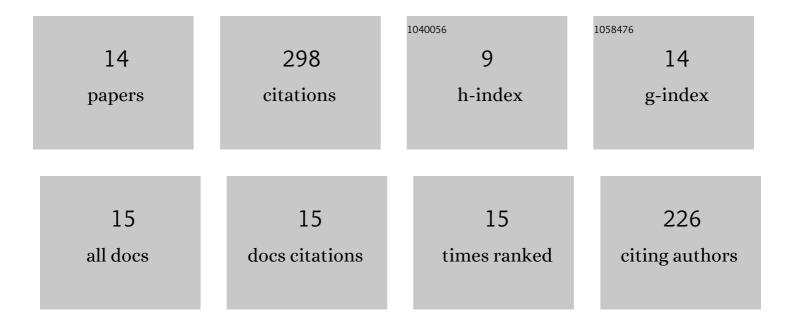
Véronique Baudin-Creuza

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Unstable and Thalassemic α Chain Hemoglobin Variants: A Cause of Hb H Disease and Thalassemia Intermedia. Hemoglobin, 2008, 32, 327-349.	0.8	117
2	Impaired binding of AHSP to α chain variants: Hb Groene Hart illustrates a mechanism leading to unstable hemoglobins with α thalassemic like syndrome. Blood Cells, Molecules, and Diseases, 2006, 37, 173-179.	1.4	37
3	Transfer of Human α- to β-Hemoglobin via Its Chaperone Protein. Journal of Biological Chemistry, 2004, 279, 36530-36533.	3.4	33
4	The α-hemoglobin stabilizing protein and expression of unstable α-Hb variants. Clinical Biochemistry, 2009, 42, 1818-1823.	1.9	21
5	α-Hemoglobin Stabilizing Protein (AHSP), a Kinetic Scheme of the Action of a Human Mutant, AHSPV56G. Journal of Biological Chemistry, 2010, 285, 17986-17992.	3.4	21
6	Role of α-Globin H Helix in the Building of Tetrameric Human Hemoglobin: Interaction with α-Hemoglobin Stabilizing Protein (AHSP) and Heme Molecule. PLoS ONE, 2014, 9, e111395.	2.5	17
7	Evaluation of the free αâ€hemoglobin pool in red blood cells: A new test providing a scale of βâ€thalassemia severity. American Journal of Hematology, 2011, 86, 199-202.	4.1	14
8	Dynamics of α-Hb chain binding to its chaperone AHSP depends on heme coordination and redox state. Biochimica Et Biophysica Acta - General Subjects, 2014, 1840, 277-287.	2.4	10
9	Red blood cells free αâ€haemoglobin pool: a biomarker to monitor the βâ€thalassemia intermedia variability. The <scp>ALPHAPOOL</scp> study. British Journal of Haematology, 2017, 179, 142-153.	2.5	10
10	Elevated soluble αâ€hemoglobin pool in sickle cell anemia. American Journal of Hematology, 2017, 92, E593-E595.	4.1	6
11	αâ€Haemoglobin pool measurement: a useful biomarker for evaluation of βâ€ŧhalassaemia intermedia? – response to Huang and Li. British Journal of Haematology, 2018, 183, 671-673.	2.5	1
12	Hydroxycarbamide decreases the free alphaâ€hemoglobin pool in red blood cells of adult patients with sickle cell anemia. American Journal of Hematology, 2020, 95, E302-E305.	4.1	1
13	Alpha haemoglobinâ€stabilising protein concentration in the red blood cells of patients with sickle cell anaemia with and without hydroxycarbamide treatment. British Journal of Haematology, 2021, , .	2.5	1
14	Detection and follow-up of a soluble alpha-haemoglobin pool in the red cells of stored blood units. Blood Transfusion, 2021, , .	0.4	0