Graham Serjeant

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/9423678/publications.pdf

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		1040056	1199594
13	285	9	12
papers	citations	h-index	g-index
13	13	13	180
all docs	docs citations	times ranked	citing authors

#	Article	lF	CITATIONS
1	Post-natal decline of fetal haemoglobin in homozygous sickle cell disease: relationship to parental Hb F levels. British Journal of Haematology, 1982, 52, 455-463.	2.5	48
2	The haematology of homozygous sickle cell disease after the age of 40 years. British Journal of Haematology, 1991, 77, 382-385.	2.5	44
3	Determinants of haemoglobin level in steady-state homozygous sickle cell disease. British Journal of Haematology, 1996, 92, 143-149.	2.5	40
4	The Mechanisms of Low Birth Weight in Infants of Mothers With Homozygous Sickle Cell Disease. Pediatrics, 2007, 120, e686-e693.	2.1	36
5	Haematological indices in normal negro children: a Jamaican cohort from birth to five years. International Journal of Laboratory Hematology, 1980, 2, 169-178.	0.2	31
6	The clinical significance of serum transferrin receptor levels in sickle cell disease. British Journal of Haematology, 1993, 84, 301-304.	2.5	25
7	Systemic <i>Salmonella</i> infections in sickle cell anaemia. Annals of Tropical Paediatrics, 1989, 9, 169-172.	1.0	18
8	Alpha thalassaemia and the haematology of normal Jamaican children. International Journal of Laboratory Hematology, 1985, 7, 289-295.	0.2	14
9	Haemoglobin Variant Screening in Jamaica: Meeting Student's Request. British Journal of Haematology, 2016, 172, 634-636.	2.5	10
10	Intravenous oxpentifylline and the painful crisis of sickle cell disease. Clinical Hemorheology and Microcirculation, 2016, 10, 35-42.	1.7	8
11	Pregnancy outcome in homozygous sickle cell disease: observations from the Jamaican Birth Cohort. BJOG: an International Journal of Obstetrics and Gynaecology, 2021, 128, 1703-1710.	2.3	7
12	Sickle Cell-Hemoglobin Caribbean - A Benign Syndrome. Hemoglobin, 1982, 6, 403-405.	0.8	4
13	Stilboestrol and Stuttering Priapism in Homozygous Sickle-Cell Disease. Journal of Urology, 1986, 136, 543-544.	0.4	О