Dr Jo Howard's publications

Community Acquired Salmonella Bacteraemia in patients with Sickle Cell Disease 1969-2008. A Single Centre Study. Richards LH, Howard J, Klein JL. Scandinavian J Infection. 2011. 43 (2) 89-94

A prospective Diary Study of Stuttering Priapism in Adolescents and Young men with Sickle Cell Anaemia: Report of an International Randomized Controlled Trial: the Priapism in Sickle Cell Study (PISCES). Olujohungbe AB, Adeyoju A, Yarduminan A, Akinyanjy J, Morris J, Westerdale N, Akenova Y, Kehinde O, Anie K, Howard J, Brooks A, Davis V, Inati A. J Andrology. 2010 epub

Pregnancy outcomes in sickle cell disease: a retrospective cohort study from two cohort studies in the UK. Chase AR, Sohal M, Howard J, Laher R, McCarthy A, Layton DM, Oteng-Ntim E. Obstetric Medicine 2010

Silent cerebral infarcts occur despite regular blood transfusion therapy after first stroke in children with sickle cell disease. Hulbert ML, McKinstry RC, Lacey JL, Moran CJ, Panepinto JA, Thompson AA, Sarnaik SA, Woods GM, Casella JF, Inusa B, Howard J, Kirkham FJ, Anie KJ, Mullin JE, Ichord R, Noetzel M, Yan Y, Rodeghier M, Debaun M et al. Blood 2010

A qualitative study of perceptions of using prophylactic penicillin in patients with Sickle Cell Disease. Abedian M, Howard J, Rawle, H, Thomas VJ. Primary Health Care. 2010, 20, 7, 26-31)

Influence of a thalassaemia on hydroxycarbamide response in sickle cell disease. Vasavada N, Badiger S, Rees D, Height S, Howard J, Thein SL. BJ Haem 2008 143, 589-592

Longitudinal Analysis of Pulmonary Function in Adults with Sickle Cell Disease. American Journal of Haematology. Field JJ, Glassberg J, Gilmore A, Howard J, Patankar S, Yan Y, Davies SC, DeBaun MR, Strunk RC. Am J Hematol. 2008 Jul 83 (7): 574-6

Moving young people with sickle cell disease from paediatric to adult services. Howard J, Woodhead T, Musumadi L, Martell A, Inusa B. 2010 British Journal Hospital Medicine. 71 (6) 310-314

The role of hydroxycarbamide in the prevention of complications in patients with sickle cell disease. Wiles N, Howard J. Therapeutics and Clinical Risk Management. 2009, 5, 745-755

Pain management and quality of life in sickle cell disease. Howard J, Thomas VJ, Rawle HM. Expert Review of Pharmacoeconomics and Outcomes Research, 2009. 9; 4 ; 347- 352

The clinical care of adult patients with sickle cell disease. Olujohungbe A, Howard J. BJ Hospital Medicine, 2008. 69; 11; 616-619