

## Sally Watt's publications

Diagnosis of cystic fibrosis following the NHS newborn blood spot screening programme.  
**Sally Watts** and Roberta Rizzo British Journal of Midwifery • August 2015 • Vol 23, No 8  
pp394-398

Ethical and cultural issues. Chapter 34. Athalie Melville and **Sally Watts**  
Genetics for Health Care Professionals in Cancer Care: From Principles to Practice  
Oxford University Press 2014

Marion McAllister, Gillian Scott, **Sally Watts**, Charlotte Eddy, Athalie Melville, Anita Bruce  
A new approach to service evaluation in clinical genetics services using a Patient Reported  
Outcome Measure: Measuring patient benefits. The Newsletter of the British Society for  
Human Genetics Issue 48 January 2013

Gareth J Hollands, Sophia C L Whitwell, Richard A Parker, Natalie J Prescott, Alastair  
Forbes, Jeremy Sanderson, Christopher G Mathew, Cathryn M Lewis, **Sally Watts**, Stephen  
Sutton, David Armstrong, Ann Louise Kinmonth, A Toby Prevost, Theresa M Marteau  
Effect of communicating DNA based risk assessments for Crohn's disease on smoking  
cessation: randomised controlled trial.  
BMJ2012;345:e4708doi:10.1136/bmj.e4708 (published 20 July 2012)

**Sally Watts**, Tootie Bueser and MLP Robert  
A multidisciplinary service for inherited cardiac disease in a regional clinic genetics service.  
British Journal of Cardiac Nursing 2009 Vol 4 pp321-325.

Fraser L., **Watts S.**, Cargill A., Sutton S and Hodgson S  
Study comparing two types of screening provision for people with von Hippel Lindau  
disease.  
Familial Cancer 2007 (1) pp 103-11.

Warren et al including collaborators in the UK MRC study of MRI screening for breast cancer  
in women at high risk (MARIBS) (**Watts S.** (local collaborator)).  
A test of performance of breast MRI interpretation in a multicentre screening study.  
MRI 2006 24(7) pp 917-29.

**Watts S.** and O'Shea J.  
Guy's Experience as the first Genetics Department to undergo Agenda for Change. Editorial  
Article British Society of Human Genetics  
Newsletter February 2004

Stermer T., Hodgson S., Kavalier F., **Watts S.** and Jones R.  
Patients' and professionals' opinions of services for people at an increased risk of colorectal  
cancer: an exploratory qualitative study.  
Familial Cancer 2004 3:49-53 2004

D.Ellis, J. Greenman, S.V. Hodgson, S. McCall, F. Laloo, J. Cameron , L. Izatt, G. Scott, C.  
Jacobs, **S. Watts**, W. Chorley, C. Perrett, K MacDermot, S. Mohammed,  
G Evans, C. Mathew.  
Low prevalence of germline BRCA1 mutations in early onset breast cancer without a family  
history.  
J. Med.Genet 2000 37 (8) 792-794

J. Greenman, S. Mohammed, D. Ellis, **S. Watts**, G.Scott, D. Barnes, E. Solomon, S. Hodgson, C. Mathew.  
Identification of missense and truncating mutations in the BRCA1 gene in sporadic and familial breast and ovarian cancer.  
Genes, Chromosomes and Cancer September 1997