Curriculum Vitae



Charles James Shaw-Smith

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Currently tide

Personal details

Current Post Consultant in Clinical Genetics, Department of

Clinical Genetics, Heavitree Hospital, Exeter

Work address Royal Devon and Exeter NHS Trust, Heavitree

Hospital, Gladstone Road, Exeter EX2 1ED

Telephone 01392 405737 (work)

E mail charles.shaw-smith@nhs.net

Career Summary

 Consultant Clinical Geneticist with 15 years' experience at Consultant level

- Main areas of clinical expertise: chromosomal disorders; congenital malformations; renal genetics; connective tissues disorders such as Marfan syndrome, Ehlers-Danlos syndrome, osteogenesis imperfecta; cancer genetics; all common disorders in Clinical Genetics
- Extensive research background with 75 peer-reviewed research papers including reviews, plus contributions to text books, and a singleauthored text book

Medico-legal work

Training

- Module 1 of the Bond Solon course "Excellence in Report Writing" completed (June, 2013)
- Module 2 of the Bond Solon course "Courtroom Skills" Completed (April, 2015)

Instructions received/undertaken in the following areas:

- Chromosome disorders in children
- Congenital malformation syndromes
- Connective tissue disorders: Ehlers-Danlos syndrome, Marfan syndrome, osteogenesis imperfecta
- Single gene disorders

I have completed around 50 case reports since 2014. These have been in the areas of child protection and clinical negligence, with the majority for the complainant.

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Qualifications

1986 MA (Cantab)

1989 BM BCh (Oxon)

1992 MRCP (UK)

2000 PhD (University of London)

Previous Posts

1st September 2006 to 31st August, 2010

Wellcome Trust Intermediate Clinical Fellow, Sanger Institute, Cambridge and Institute of Child Health, London. Honorary Consultant in Clinical Genetics, Addenbrooke's Hospital, Cambridge

1st June 2004 to 31st August, 2006

University Lecturer and Honorary Consultant, Department of Medical Genetics Addenbrooke's Hospital, Cambridge

Other professional roles

- Rare Disease Lead for SouthWest Genomic Medicine Centre, Genomics England 100 000 Genomes Project, 2014-19
- Member, UKGTN Steering Group: Working Party on the evaluation of microarray technology for chromosomal disorders, Chair: Dr Hilary Burton.
- Patron and Member, Tracheo-Oesophageal Fistula Support (TOFS) [UK support group for patients born with oesophageal atresia/trachea-oesophageal fistula].

Membership of Clinical Societies

Member, British Society of Human Genetics Patron and Member, Tracheo-Oesophageal Fistula Support (TOFS) [UK support group for patients born with oesophageal atresia/trachea-oesophageal fistula].