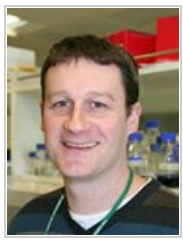
Dr Neil Morgan

Lecturer in Cardiovascular Genetics

Cardiovascular and Respiratory Sciences



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About

Neil Morgan is a Lecturer in Cardiovascular Genetics within the School of Clinical and Experimental Medicine.

He has published over 50 research papers in high impact scientific journals in the field of human genetics. His research has primarily involved the identification of novel genes for autosomal recessive inherited diseases.

Qualifications

- PhD in Molecular Genetics 2005
- BSc (Hons) Applied Biochemistry 1994

Biography

Neil Morgan qualified with a BSc (Hons) in Applied Biochemistry from Liverpool John Moores University in 1994. His first post was as Research Assistant at Guy's Hospital until 2000 followed by a brief spell as a Research Associate at Leicester University. He joined the Medical and Molecular Genetics group at the University of Birmingham where he went onto study for a PhD in Molecular Genetics. He has recently been appointed as a lecturer in the Centre for Cardiovascular Sciences.

Teaching

- **Medical Science BMedSc (/undergraduate/courses/med/medical-sci.aspx)**
- Biomedical Research - Molecular and Cellular Medicine MRes

Research

His current research is focussing on the molecular genetics of patients with platelet bleeding disorders and low platelet counts (thrombocytopenia). The identification of novel gene defects will provide clues to genes and proteins involved in normal platelet physiology and ultimately lead to devising new treatment strategies to minimise the risk of bleeding in such patients.

His past research has primarily involved identification of the novel genes for autosomal recessive inherited diseases. He has been extremely successful in his pursuit and key findings include:

- *TRAC* mutations causing a novel immunodeficiency disorder
- *SLC29A3* mutations causing familial Faisalabad histiocytosis/Rosai-Dorfman disease
- *BLOC1S3* mutations causing a novel form of Hermansky Pudlak disorder (HPS8)
- *CHRNG*, *RAPSN* and *DOK7* mutations causing multiple pterygia syndrome/foetal akinesia deformation sequence
- *PLA2G6* mutations causing a spectrum of childhood onset neurodegenerative disorders associated with brain iron accumulation

Publications

Morgan NV, Hartley JL, Setchell KDR,Simpson MA, Brown R, Tee L, Kirkham S,Pasha S, Trembath RC, Maher ER, Gissen P, Kelly DA (2013) A combination of mutations in *AKR1D1* and *SKIV2L* in a family with severe infantile liver disease. Orphanet Journal Rare Diseases; 8: 74.

Watson SP, Lowe GC, Lordkipanidze M, Morgan NV (2013) Genotyping and phenotyping of platelet function disorders. J Thromb Haemost. 11 Suppl 1:351-63.

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Nieminen P, Morgan NV, Fenwick AL, Parmanen S, Veistinen L, Mikkola ML, van der Spek PJ, Giraud A, Judd L, Arte S, Brueton LA, Wall SA, Mathijssen IMJ, Maher

ER, Wilkie AOM, Kreiborg S, Thesleff I (2011) Inactivation of IL11 Signaling Causes Craniosynostosis, Delayed Tooth Eruption and Supernumerary Teeth. Am J Hum Genet July 7. [Epub ahead of print]

Morgan NV, Goddard S, Cardno TS, McDonald D, Rahman F, Barge D, Ciupek A, Straatman-Iwanowska A ,Pasha S, Guckian M, Anderson G, Huissoon A, Cant A, Tate WP, Hambleton S, Maher ER (2011) Mutation in the TCRa subunit constant gene (*TRAC*) leads to a human immunodeficiency disorder characterized by a lack of TCRab^{+} T cells. J Clin Invest 121(2):695-702

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Morgan NV, Morris MR, Cangul H, Gleeson D, Straatman-Iwanowska A, Davies N, Keenan S, Pasha S, Rahman F, Gentle D,Vreeswijk MPG, Devilee P, Knowles MA, Ceylaner S, Trembath RC, Dalence C, Kismet E, Koseoglu V, Rossbach H-C, Gissen P, Tannanhill D, Maher ER (2010). Mutations in SLC29A3, encoding an equilibrative nucleoside transporter ENT3, cause a familial Histiocytosis syndrome (Faisalabad histiocytosis) and Familial Rosai-Dorfman disease. PLoS Genetics 6(2):e1000833