

## Dr David Withers PhD

Wellcome Trust Research Fellow

School of Immunity and Infection

### Contact details

Telephone [+44 \(0\)7534 288956](tel:+44%207534%20288956) (tel:+44 7534 288956)

Email [d.withers@bham.ac.uk](mailto:d.withers@bham.ac.uk) (mailto:d.withers@bham.ac.uk)

School of Immunity and Infection  
College of Medical and Dental Sciences  
University of Birmingham  
Edgbaston  
Birmingham  
B15 2TT  
UK



### About

David Withers is a Wellcome Trust funded Research Fellow

His work is focused on CD4 T cell responses and the secondary lymphoid tissue within which these responses are initiated.

### Qualifications

- PhD Immunology 2004
- BSc (Hons) Microbiology and Virology 2000

### Biography

David Withers qualified with a BSc (Hons) in Virology and Microbiology from the University of Warwick in 2000. He went on to study for a PhD in Immunology at the Institute for Animal Health in conjunction with the University of Bristol. After obtaining his PhD, David continued his studies in the laboratory of Peter Lipsky at NIAMS, NIH, Bethesda (2004-2006). He then returned to the UK to study with Peter Lane at the University of Birmingham, cementing his interest in secondary lymphoid tissue development/structure and how this controlled CD4 T cell responses. In 2011 he was awarded a Wellcome Trust Career Development Fellowship to continue his research interests and establish his own research group

### Teaching

#### Teaching Programmes

- [MBChB IIH \(2nd year\) \(/undergraduate/courses/med/medicine.aspx\)](#)
- [BMedSc3 ExplImm \(/undergraduate/courses/med/medical-sci.aspx\)](#)

### Postgraduate supervision

David is interested in supervising doctoral research students in the following areas:

- Generation and survival of memory CD4 T cells
- Secondary lymphoid tissue generation, organisation and maintenance

If you are interesting in studying any of these subject areas please contact David on the contact details above, or for any general doctoral research enquiries, please email: [dr@contacts.bham.ac.uk](mailto:dr@contacts.bham.ac.uk) (mailto:dr@contacts.bham.ac.uk) or call or call +44 (0)121 414 5005.

For a full list of available Doctoral Research opportunities, please visit our Doctoral Research programme listings.

### Research

#### RESEARCH THEMES

T cell responses, Immunological memory, Secondary lymphoid tissue.

#### RESEARCH ACTIVITY

David's research is focused on understanding the signals involved in the development and maintenance of T cell responses, in particular, the development of memory CD4 T cells which are essential for developing immunological memory. Immunological memory underpins our strategies of vaccination. Understanding how memory CD4 T cells are generated and maintained is crucial for improving upon our ability to both vaccinate and modulate unwanted self reactivity. CD4 T cell responses are initiated within secondary lymphoid tissues such as lymph nodes. Understanding how these structures are formed and also maintained is also important for a full understanding of the response.

#### Key areas of research:

##### *Memory CD4 T cell survival and function*

To improve our ability to modulate memory CD4 T cells, greater understanding of the signals involved is required. Since any given T cell clone is present in very low numbers within the host, it is imperative to track low numbers of antigen-specific T cells within our experimental approaches. Using a combination of techniques including

### **Lymph node structure and function**

The development of lymph nodes is dependent upon lymphoid tissue inducer cells, a specialised innate cell type that initiates the formation of these structures during embryonic development. Given that lymph node development is restricted to a window of embryonic development, it is somewhat surprising that these cells persist with adult lymph nodes, suggesting further functions within this tissue. Utilising different genetically modified mice, these cells can be removed from developed lymph nodes and the impact this has on lymph node function assessed.

### **Publications**

Withers, D.R., Gaspal, F.M., Bekiaris, V., McConnell, F.M., Kim, M.-Y., Anderson, G., and Lane, P.J.L. (2011). Role of OX40 and CD30 in murine CD4 effector and memory function. *Immunol Rev* In press

Withers, D.R. (2011). Lymphoid tissue inducer cells. *Curr Biol* 21, R381-382.

Li, Y., Innocentin, S., Withers, D.R., Roberts, N.A., Gallagher, A.R., Grigorieva, E.F., Wilhelm, C., and Veldhoen, M. (2011). Exogenous Stimuli Maintain Intraepithelial Lymphocytes via Aryl Hydrocarbon Receptor Activation. *Cell*.

Gaspal, F.M., Withers, D., Saini, M., Bekiaris, V., McConnell, F.M., White, A., Yagita, H., Walker, L.S.K., Anderson, G., and Lane, P.J.L. (2011). Abrogation of CD30 and OX40 signals prevents autoimmune disease in FoxP3 deficient mice. *J Exp Med* 208 1579-1584, 1579-1584.

Kim, S., Han, S., Withers, D.R., Gaspal, F., Bae, J., Baik, S., Shin, H.C., Kim, K.S., Bekiaris, V., Anderson, G., et al. (2011). CD117(+) CD3(-) CD56(-) OX40L(high) cells express IL-22 and display an LT $\alpha$ i phenotype in human secondary lymphoid tissues. *Eur J Immunol* 41, 1563-1572.

Withers, D.R., Jaensson, E., Gaspal, F., McConnell, F.M., Eksteen, B., Anderson, G., Agace, W.W., and Lane, P.J. (2009). The survival of memory CD4+ T cells within the gut lamina propria requires OX40 and CD30 signals. *J Immunol* 183, 5079-5084.

