Personal Information

Name: Biliana Lozanoska-Ochser

Education

2003 – 2007	PhD in Immunology, King's College London, UK
2001 - 2003	MSc Medical Immunology, University College London, UK
1998 – 2000	MSc Human Metabolism, University of Aberdeen, UK
1995 – 1998	$\textbf{BSc Biomedical Sciences,} \ \textit{University of Westminster, London, UK}$

Employment	
2014 - Present	Research Scientist, Sapienza University of Rome, IT
	Project: The immune response in Duchenne Muscular Dystrophy
2012 – 2013	Post-Doctoral Fellow, California Institute for Biomedical Research, San Diego, US Project: M1 and M2 macrophages in inflammation.
2009 – 2011	Post-Doctoral Scientist, Yale University School of Medicine, New Haven, US Department of Pulmonary Medicine

Post-Doctoral Fellow, Kings College London, UK 2007 - 2009

Department of Immunobiology

Project: Pancreatic islet endothelial cells and plasmacytoid dendritic cells in Type 1 Diabetes.

Key responsibilities in addition to research:

- Under-graduate and Post-graduate teaching.
- Project supervisor of postgraduate and PhD students.

Project: The role of macrophages in lung inflammation.

- Writing manuscripts and paper reviews for publication.
- Oral presentations at international conferences and departmental seminars.
- Organizing and participating in journal clubs.

Research Skills

Broad range of practical skills in human and mouse immunology including:

- Isolation (cell sort or magnetic beads), culture and in-vitro study of mouse and human cells including: macrophages derived from lung tissue and BAL fluid; peripheral blood monocytes; dendritic cells (myeloid and plasmacytoid); endothelial cells; T cells (effector, memory and regulatory T cells); cells derived from mouse spleen and lymph nodes.
- Extensive experience working with mouse models of lung inflammation and Type 1 Diabetes.
- Multicolour Flow cytometry (surface and intracellular staining).
- Functional assays: T cell proliferation (by CFSE or thymidine incorporation), Cytokine production by ELISPOT, ELISA, LUMINEX or CBA assay.
- Trans-migration and adhesion assays.
- Molecular biology techniques including RT-qPCR.
- Confocal microscopy and immunohistochemistry.
- Assay development and validation.

Awards & Achievements

- Small Project Grant (awarded 2018), Dutch Parent Project.
- Oliver Bird grant (awarded 2009) for my written proposal to study monocyte subsets in Rheumatoid Arthritis.
- JDRF (Juvenile Diabetes Research Foundation) Fellowship Award (2007-2009).

Research Publications

Peer-reviewed research articles

Fiore P, Benedetti A, Sandonà M, Madaro L, De Bardi M, Saccone V, Puri PL, Gargioli C, Lozanoska-Ochser B and Bouché M. Lack of PKCθ promotes regenerative ability of muscle stem cells in chronic muscle injury. January 2020, Submitted to International Journal of Molecular Sciences.

Rizzo G, Di Maggio R, Benedetti A, Morroni J, Bouche M, and **Lozanoska-Ochser B**. Splenic Ly6C^{hi} monocytes are critical players in dystrophic muscle injury and repair. *JCI Insight* 2019; Epub ahead of print.

Bouchè M, Lozanoska-Ochser B, Proietti D, Madaro L. Do neurogenic and cancer-induced muscle atrophy follow common or divergent paths? *Eur J Transl Myol*. 2018, 28 (4): 393-400.

Lozanoska-Ochser B, Benedetti A, Rizzo G, Marrocco V, Di Maggio R, Fiore P, Bouche M. Targeting early PKC0-dependent T-cell infiltration of dystrophic muscle reduces disease severity in a mouse model of muscular dystrophy. *J Pathol. 2018, 244:323-333*

Marrocco V, Fiore P, Benedetti A, Pisu S, Rizzuto E, Musarò A, Madaro L, **Lozanoska-Ochser B**, Bouché M. Pharmacological Inhibition of PKCθ Counteracts Muscle Disease in a Mouse Model of Duchenne Muscular Dystrophy. *EBioMedicine*. 2017, 16:150-161.

Valeria Marrocco, Piera Fiore, Luca Madaro, Annunziata Crupi, Biliana Lozanoska-Ochser and Marina Bouché. Targeting PKC theta in skeletal muscle and muscle diseases: good or bad? Biochemical Society Transactions. 2014, 42:1550-155

Lozanoska-Ochser B and Peakman M. Level of MHC class I expression on endothelium in non-obese diabetic mice influences CD8 T cell adhesion and migration. *Clinical and Experimental Immunology*. 2009, 157:119-127.

Lozanoska-Ochser B, Klein NJ, Huang GC, Alvarez RA, Peakman M. Expression of CD86 on human islet endothelial cells facilitates T cell adhesion and migration. *Journal of Immunology*. 2008, 181:6109-16.

Lozanoska-Ochser B, Barone F, Pitzalis C, Peakman M. Atorvastatin fails to prevent the development of autoimmune diabetes despite inhibition of pathogenic beta-cell-specific CD8 T-cells. *Diabetes*.2006, 55:1004-10.

Thrower SL, James L, Hall W, Green KM, Arif S, Allen JS, Van-Krinks C, **Lozanoska-Ochser B**, Marquesini L, Brown S, Wong FS, Dayan CM, Peakman M. Proinsulin peptide immunotherapy in type 1 diabetes: report of a first-in-man Phase I safety study. *Clinical and Experimental Immunology*, 2009, 155:156-65.

Allen JS, Pang K, Skowera A, Ellis R, Rackham C, **Lozanoska-Ochser B**, Tree T, Leslie RD, Tremble JM, Dayan CM, Peakman M. Plasmacytoid dendritic cells are proportionally expanded at diagnosis of type 1 diabetes and enhance islet autoantigen presentation to T-cells through immune complex capture. *Diabetes*.2009, 58:138-45.

Zanone MM, Favaro E, Doublier S, Lozanoska-Ochser B, Deregibus MC, Greening J, Huang GC, Klein N, Cavallo Perin P, Peakman M, Camussi G. Expression of nephrin by human pancreatic islet endothelial cells. *Diabetologia*. 2005, 48:1789-97.

Favaro E, Bottelli A, Lozanoska-Ochser B, Ferioli E, Huang GC, Klein N, Chiaravalli A, Perin PC, Camussi G, Peakman M, Conaldi PG, ZanoneMM. Primary and immortalised human pancreatic islet endothelial cells: phenotypic and immunological characterisation. *Diabetologia*.2005, 48:2552-62.

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