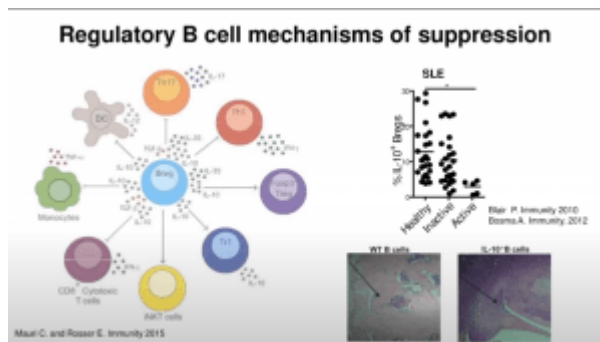
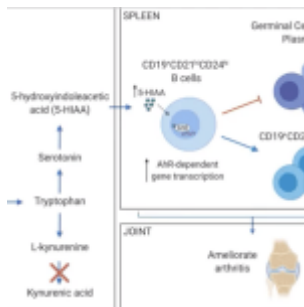


IUIS-Immunopaedia-Frontiers webinar: Getting to the “bottom” of arthritis



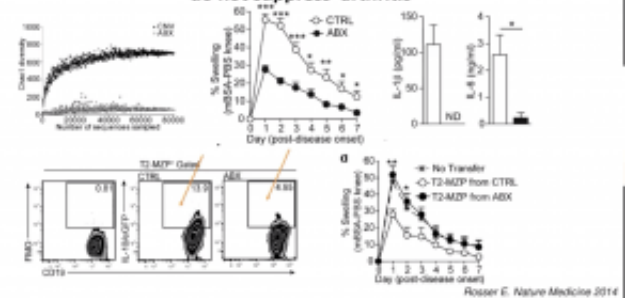
Regulatory B cells (Bregs) are immunosuppressive cells that contribute to the reduction of excessive inflammatory responses. Researchers have demonstrated that interaction between the gut microbiota and

the innate immune system in the gut-associated lymphoid tissue (GALT) promotes the differentiation of immature B cells into Bregs. Mice depleted of endogenous bacteria following administration of broad-spectrum antibiotics do not develop

arthritis or Bregs, suggesting an intricate relationship between the microbiota, inflammation, and B-reg differentiation. These findings suggest a potential link between microbiota and Bregs, which could occur either (1)

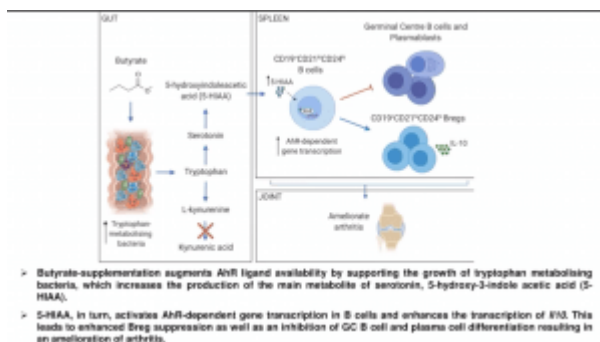
due to inflammatory signals induced by interaction with microbiota or (2) direct induced by microbial products by inflammatory. In a recent IUIS-Immunopaedia-Frontiers in Immunology webinar, Claudia Mauri (University College London, UK) discussed how microbial products induce Bregs and their

Bregs from broad spectrum ABX-treated mice produce less IL-10 and do not suppress arthritis



relevance for arthritis pathology.

Murine models of arthritis have demonstrated that pathology is associated with a reduction of commensal bacteria and short-chain fatty acids such as butyrate compared to naïve mice, a similar finding also observed in humans. Using a murine model, Claudia Mauri demonstrated that suppression of arthritis pathology by butyrate supplementation is B-reg dependent and requires induction of B-regs via aryl-hydrocarbon receptor (AhR). She also showed that butyrate-supplementation prevents plasmablast and germinal centre differentiation. Is this process direct or indirect? Microbial analysis butyrate supplemented mice demonstrated an increase in bacteria that metabolise tryptophan, an amino acid involved in and leads to the production of serotonin, indole and kynurenine, and also activate AhR. Analysis of metabolic products in mice supplemented with butyrate demonstrated a significant increase in serotonin-derived metabolite 5-Hydroxyindole-3-acetic acid (5-HIAA). Further, she showed data that 5-HIAA activates B-regs via activation of AhR and suppressed arthritis pathology.



In summary, results presented by Claudia Mauri demonstrated a role of microbial metabolites in the suppression of arthritis pathology, thus providing evidence for the link between arthritis and the gut microbiota.

Summary by Cheleka AM Mpande

Brief B-reg Introduction

Regulatory B cells (Bregs) are immunosuppressive cells that contribute to the maintenance of immunological tolerance (Mauri and Bosma, 2012). B-regs suppress a variety of immune pathologies including autoimmune diseases through the production of interleukin (IL)-10, IL-35, and transforming growth factor beta 1 (TGFb1). Toll-like receptor (TLR) agonists, including lipopolysaccharide (TLR4) and CpG oligodeoxynucleotides (TLR9), in combination with low grade levels of inflammatory cytokines, for example IFNa and/or IL-1b and IL-6, induce IL-10-producing B-reg differentiation (Lampropoulou et al., 2008; Menon et al., 2016; Rosser et al., 2014).