

# universitäts klinikumbonn

# PRESS RELEASE

# New findings on the immune system

Bonn researchers decipher an underlying mechanism for the development of T follicular helper cells

Bonn, March 04 - T follicular helper cells (Tfh) are essential for strong antibody-mediated reactions of our immune system during infections and vaccinations. However, if they get out of control, this can cause diseases such as autoimmunity, allergies or cancer. Researchers from the University Hospital Bonn (UKB) and the Cluster of Excellence ImmunoSensation<sup>2</sup> at the University of Bonn investigated the underlying mechanisms of Tfh cell development in a mouse model and thus decoded their internal networking. They hope that this will lead to new strategies for the development of highly effective vaccines and new therapies to combat various diseases. The results have now been published in the renowned journal "Science Immunology".

T follicular helper cells (Tfh cells) are a specialized subgroup within the socalled CD4<sup>+</sup> T helper cells in the immune system. Their main task is to assist the B cells in the immune defense. They are essential for the generation of highly effective antibodies. Tfh cells therefore play a decisive role in protecting against and fighting infections. "Although Tfh cells were first described over 20 years ago, there is still no reliable protocol for their generation in cell culture," says co-first author Dr. Yinshui Chang, former postdoctoral researcher at the University of Bonn at the UKB, describing the motivation to take a closer look at the process in the mouse model.

The transforming growth factor TGF- $\beta$  is a cytokine. This is a group of proteins that initiates and regulates the growth and differentiation of cells. The Bonn team led by Prof. Dr. Dirk Baumjohann has now discovered that this signaling molecule induces strong protein expression of both the transcription factor Bcl6 and the chemokine receptor CXCR5, which are characteristic of Tfh cells. The latter plays an important role in the targeted migration of Tfh cells into the vicinity of B cells. "We were able to show that the Tfh cells induced by TGF- $\beta$  in cell culture are quite similar to the Tfh cells generated in a living organism. They provide crucial help for B cells," says co-first author Luisa Bach, doctoral student at the University of Bonn at the UKB.

### Transcription factor c-Maf controls the fate of T helper cells

Using a new method based on CRISPR gene scissors, the international team led by the Bonn researchers discovered that the production of CXCR5 induced by TGF- $\beta$  is independent of the transcription factor Bcl6, but requires the transcription factor c-Maf. Remarkably, although Tfh and Th17 cells partially undergo common developmental stages, c-Maf acts as a switching factor for Tfh versus Th17 cell fates. Th17 cells are another special

## Medical Director and Chairman of the Board

Tel: +49 228 287-10900 Fax: +49 228 287-9010900 wolfgang.holzgreve@ukbonn.de

#### Communication and media

Viola Röser Management

Tel: +49 228 287-10469 viola.roeser@ukbonn.de

Bonn University Hospital Communication and media Venusberg Campus 1 Geb. 02 53127 Bonn





type of CD4<sup>+</sup> T helper cells and play an important role in bacterial infections and autoimmune diseases.

"Overall, our data clarify important aspects of the long-unclear prerequisites and molecular pathways for the development of Tfh cells. They also highlight the diverse functions of the transforming growth factor TGF- $\beta$ . Furthermore, these data indicate that Tfh cell development in mice and humans may not be as different as we previously assumed," says Prof. Baumjohann from the Medical Clinic III for Hematology, Oncology, Immuno-Oncology and Rheumatology at the UKB, who is a member of the Cluster of Excellence ImmunoSensation<sup>2</sup> and the Transdisciplinary Research Area (TRA) "Life & Health" at the University of Bonn. "Importantly, our findings may have implications for the development of new therapeutic strategies that enhance Tfh cells during vaccinations and infections or inhibit them in autoimmune and allergic diseases."

**Funding:** The study was funded in part by the German Research Foundation (Deutsche Forschungsgemeinschaft, DFG), the Cluster of Excellence ImmunoSensation<sup>2</sup>, and the Bonn-Melbourne Research Excellence Fund of the Universities of Bonn and Melbourne.

#### **Publication:**

Yinshui Chang, Luisa Bach et al; TGF-β specifies T<sub>FH</sub> versus T<sub>H</sub>17 cell fates in murine CD4<sup>+</sup> T cells through c-Maf; Science Immunology; DOI: https://doi.org/10.1126/sciimmunol.add4818

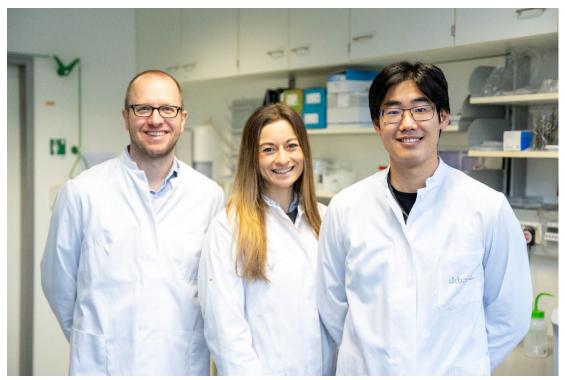
#### Scientific contact:

Prof. Dr. Dirk Baumjohann Medical Clinic III for Hematology, Oncology, Immuno-Oncology and Rheumatology Bonn University Hospital Cluster of Excellence ImmunoSensation<sup>2</sup> & TRA "Life & Health", University of Bonn Phone: 49 (0)228 287 51185 E-mail: dirk.baumjohann@uni-bonn.de

#### Image material:







## Caption: New findings on the immune system:

(from left) Prof. Dirk Baumjohann, Luisa Bach and Dr. Yinshui Chang clarify the mystery surrounding the development of follicular T helper cells.

Picture credits: University Hospital Bonn (UKB) / Alessandro Winkler

#### **Press contact:**

Dr. Inka Väth Deputy Press Officer at the University Hospital Bonn (UKB) Communications and Media Office at Bonn University Hospital Phone: (+49) 228 287-10596 E-mail: inka.vaeth@ukbonn.de

**About Bonn University Hospital:** The UKB treats around 500,000 patients per year, employs around 9,000 staff and has total assets of 1.6 billion euros. In addition to the 3,500 medical and dental students, 550 people are trained in numerous healthcare professions each year. The UKB is ranked first among university hospitals (UK) in NRW in the science ranking and in the Focus clinic list and has the third highest case mix index (case severity) in Germany. In 2022 and 2023, the F.A.Z. Institute recognized the UKB as the most desirable employer and training champion among public hospitals in Germany.