

## Research Project

### Immune-mediated response to nutrition in physiology and pathology

#### Third-party funded project

**Project title** Immune-mediated response to nutrition in physiology and pathology

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Departement Biomedizin / Diabetes Research (Donath)

Bereich Medizinische Fächer (Klinik) / Endokrinologie, Diabetologie und Metabolismus (Donath)

**Department**

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**Status** Completed

We and others have demonstrated a pathologic role of chronic inflammation in metabolism. More recently, other studies point to a physiological role of the immune system in the regulation of metabolism. Indeed, we have shown that IL-6 enhances insulin secretion via GLP-1, that macrophage-derived IL-1 $\beta$  potentiates postprandial insulin secretion, and that IL-33-activated islet-resident innate lymphoid cells promote insulin secretion. These processes hint towards a role of the immune system in the endocrine regulation of metabolism. Overall objectives: While the role of the immune system in metabolism has been extensively investigated in pancreatic islets and insulin sensitive tissues, little attention has been given to a potential role of the innate immune system in 3 additional circumstances influencing metabolism, namely (i) the cephalic phase of insulin secretion, which enhances insulin secretion not only while anticipating food, but also during its resorption; (ii) pregnancy, which often leads to gestational diabetes; and (iii) the immune cell infiltration of the exocrine pancreas which occurs in patients with type 2 diabetes. Impact: Understanding the physiology and pathophysiology of the role of the immune system in metabolism is critical for guiding the clinical development of immune treatment of type 2 diabetes and its complications.

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**Add publication**

**Add documents**

**Specify cooperation partners**