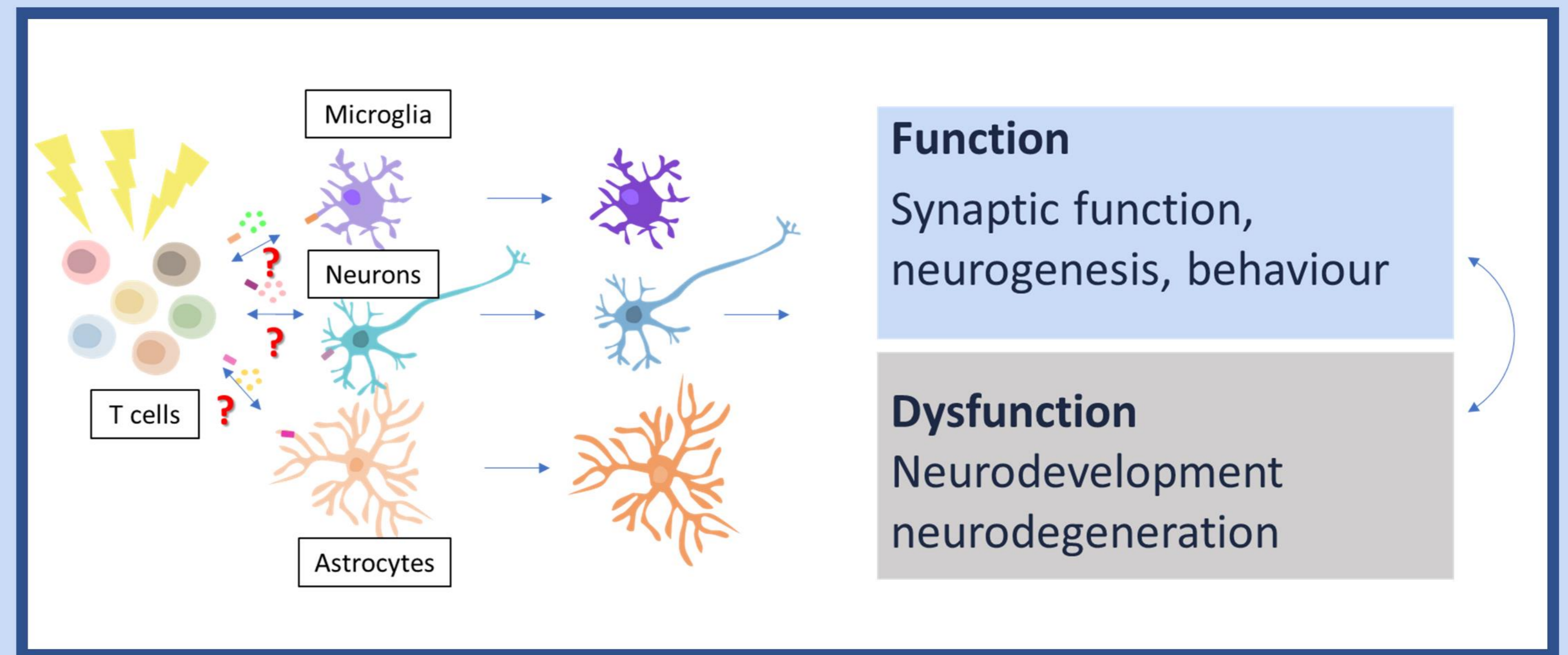


The **NIM lab** studies the complex interactions between the immune and nervous systems. The role of the immune system in brain physiology and pathology is a rapidly evolving topic in science. T cells represent the central component of adaptive immune responses, serving as vital protectors against infections and contributors to immune tolerance. Although the significance of T cells in neuroinflammatory and neurodegenerative processes is gaining recognition, their presence and potential functions in a healthy brain are still not fully elucidated. Our dual objective is to explore the **contribution of T cells to mechanisms of brain plasticity and their impact on neurodegenerative and neurodevelopmental disorders**, with the mission of identifying new biomarkers and therapeutic targets/agents. Capitalizing on my multidisciplinary training we are forging a translational approach that integrates immunology, neuroscience, biochemistry, and behavioural neuroscience to study the involvement of T cells in brain physiology and pathology. Our ultimate aim is to gain mechanistic insights into how T cells influence brain plasticity and diseases and to pinpoint molecular targets for innovative therapeutics. We have devised a comprehensive research pipeline that commences with in-depth immunological profiling for the identification of novel immunological pathways. This is followed by rigorous functional validation to assess the role of the identified circuits. We will employ genetic models and AAV-based gene therapy to target specific pathways, administer biologics, or modulate the presence of beneficial immune infiltrates in the brain.



Project 1: Role of CD4⁺-T cell in mechanisms of brain plasticity and neuroprotection

<p>T cell modulation</p> <p>Enriched environment T cells depletion</p>	<p>Immune phenotyping</p> <p>scRNA seq hpFlow Cite seq</p>
<p>Computational analysis</p> <p>Pathways Cell-Cell interaction</p>	<p>Gene Targeting</p> <p>Synaptic structure Neurogenesis Behaviour</p>

Why T cells?

<p>CD4 T cell KO</p> <p>Microglia Maturation Fetal → Adult Synaptic Pruning Learning and Memory, Anxiety</p>	<p>Enriched Gut microbiome</p> <p>SPF Co-housed</p> <p>cells</p> <p>Treg Tconv</p>	<p>Enriched environment</p> <p>Standard Enriched</p> <p>cells</p> <p>Treg Tconv</p>
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Project 2: T cells dependent enviromimetics to reduce inflammation and Neurodegeneration

<p>T cells modulation</p> <p>AD model</p> <p>Enriched environment T cells depletion_upregulation T cells-microglia interaction</p>	<p>Pathology</p> <p>Inflammation Abeta pathology Neurogenesis, Behaviour</p>
<p>Cell states & Interaction</p> <p>T cells Microglia Astrocytes</p>	<p>Gene therapy</p> <p>Inflammation Abeta pathology</p>

Project 3: Role of brain resident T cells in Autism Spectrum Disorder

<p>Immune phenotyping</p> <p>scRNA seq hpFlow</p>	<p>Functional tests</p> <p>Suppression assay Adoptive transfer Bone marrow chimera</p>	<p>Gene therapy</p> <p>Synaptic structure Synaptic transmission Behaviour</p>
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The NIM team

Identify novel mechanism of brain plasticity

Microglia - T cells interaction
Plasticity and neuroprotection

Role of T cells in neuropathology

Immune phenotyping in Autism model (ADNP)
Microglia-T cells interplay in AD
Gut-brain axis in Colitis

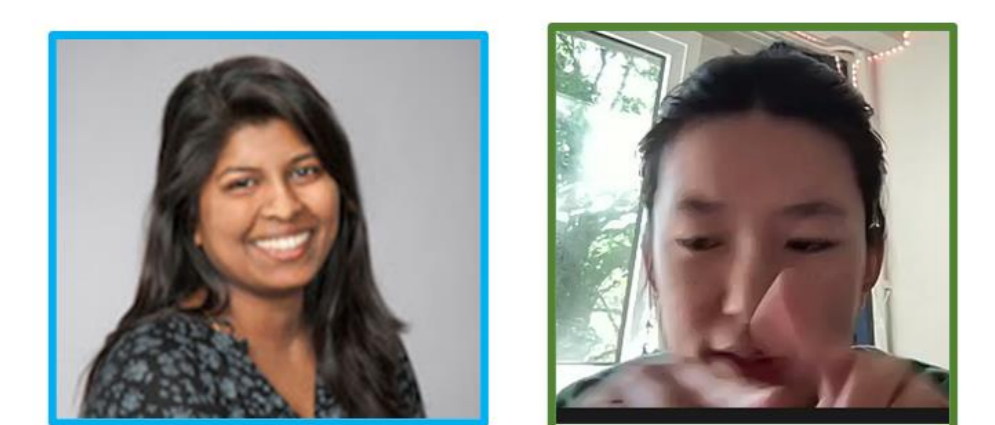
Available Technology:

Immune phenotyping platform: High parametric flow cytometry (Symphony), Spectral flow cytometry (Aurora), scRNA seq, Cite seq, Cytokine profiling

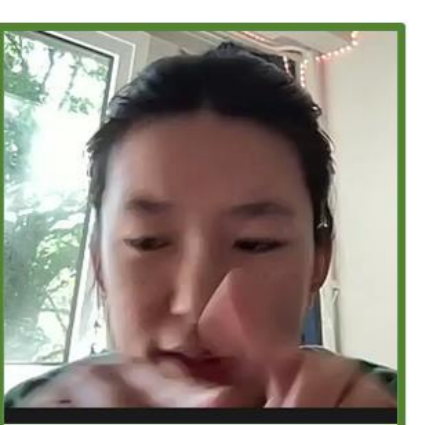
Therapeutic tools development: Gene therapy, enviromimetics, Targeting T cells dependent mechanisms

Inflammation/Disease models: Neurodegeneration (AD), Autism (FXS), Neuroinflammation (MS), Colitis

If you like our research and want to talk more about it and learn about neuroimmunology please contact: Emanuela.Pasciuto@uantwerpen.vib.be
Website: <https://pasciutolab.sites.vib.be/en>



Arthi Shanmugavadivu



Soniya Lima



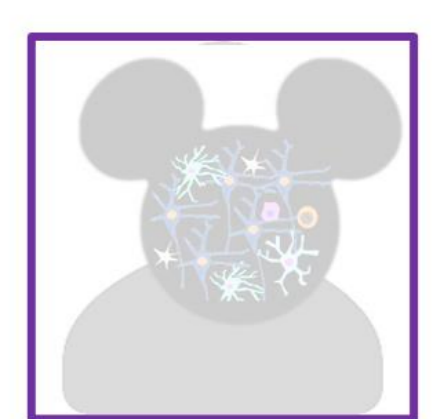
Lisa Zanoletti



Mobina Alemi



Emanuela Pasciuto
Lab Head



TBD

Key Publications

1. Pasciuto, E., et al. Neuron, 2015.
2. Briz, V.*, Restivo, L.*, Pasciuto E.*, et al. Nature Communications, 2017.
3. Pasciuto, E., et al. Cell, 2020.
4. Yshii, L. *, Pasciuto, E. *, et al. Nat Immunol, 2022.