

Unveiling dynamics of human macrophage specification during prenatal development

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Human Embryo. Credit: Ed Uthman, MD/Wikipedia

Researchers led by Prof. Li Hanjie from the Shenzhen Institute of Advanced Technology (SIAT) of the Chinese Academy of Sciences have unveiled the dynamics of human macrophage specification across 19 different tissues from early embryonic stages.

The study was published in [Cell](#) on Sept. 12.

Macrophages, i.e., pivotal immune [cells](#), have long been enigmatic in terms of their diversity and roles during [human development](#). Based on findings in rodents, the scientific community has gained some insights into the diversity, developmental origins, and tissue-specific formation of macrophage subtypes. However, it was not clear whether these findings are conserved in humans.

In this study, Prof. LI's team combined single-cell transcriptome sequencing, bioinformatic techniques, immunofluorescence, and in vitro functional assays to construct a high-resolution spatiotemporal dynamic map of human prenatal immune system development across 18 gestational stages (post-conceptual weeks four to 26) and 19 tissues.

Based on this map, the team focused on the most tissue-specific lineage, macrophages. They revealed the origin of differentiation, spatial localization, functional characteristics, and transcriptional regulation mechanisms of multiple macrophage subtypes during development.

"Key finding of our study is the identification of 15 distinct macrophage subtypes, with particular attention to two novel populations: microglia-like cells and proangiogenic macrophages (PraM)," said Prof. Li.

Surprisingly, microglia-like cells, reminiscent of those in the central nervous system, were found in unexpected locations such as the fetal epidermis, testicles, and heart. These cells were found to influence neural crest cell differentiation, contributing to early tissue development.

The research also highlighted the strategic placement of proangiogenic macrophages in perivascular areas across various fetal organs. These [macrophages](#), likely originating from the yolk sac, play a critical role in vascular development during prenatal stages.

"Our study offers a comprehensive map of human macrophage diversity and developmental processes, revealing their multifaceted roles in [development](#)," said Dr. Wang Zeshuai, first author of the study.

"These findings are poised to revolutionize our understanding of these immune cells and offer promising avenues for potential therapeutic interventions."

More information: Hanjie Li et al, An immune cell atlas reveals the dynamics of human macrophage specification during prenatal development, *Cell* (2023). [DOI: 10.1016/j.cell.2023.08.019](https://doi.org/10.1016/j.cell.2023.08.019). [www.cell.com/cell/fulltext/S0092-8674\(23\)00908-X](https://www.cell.com/cell/fulltext/S0092-8674(23)00908-X)

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