Retinal microglia in the mouse model of oxygen-induced retinopathy

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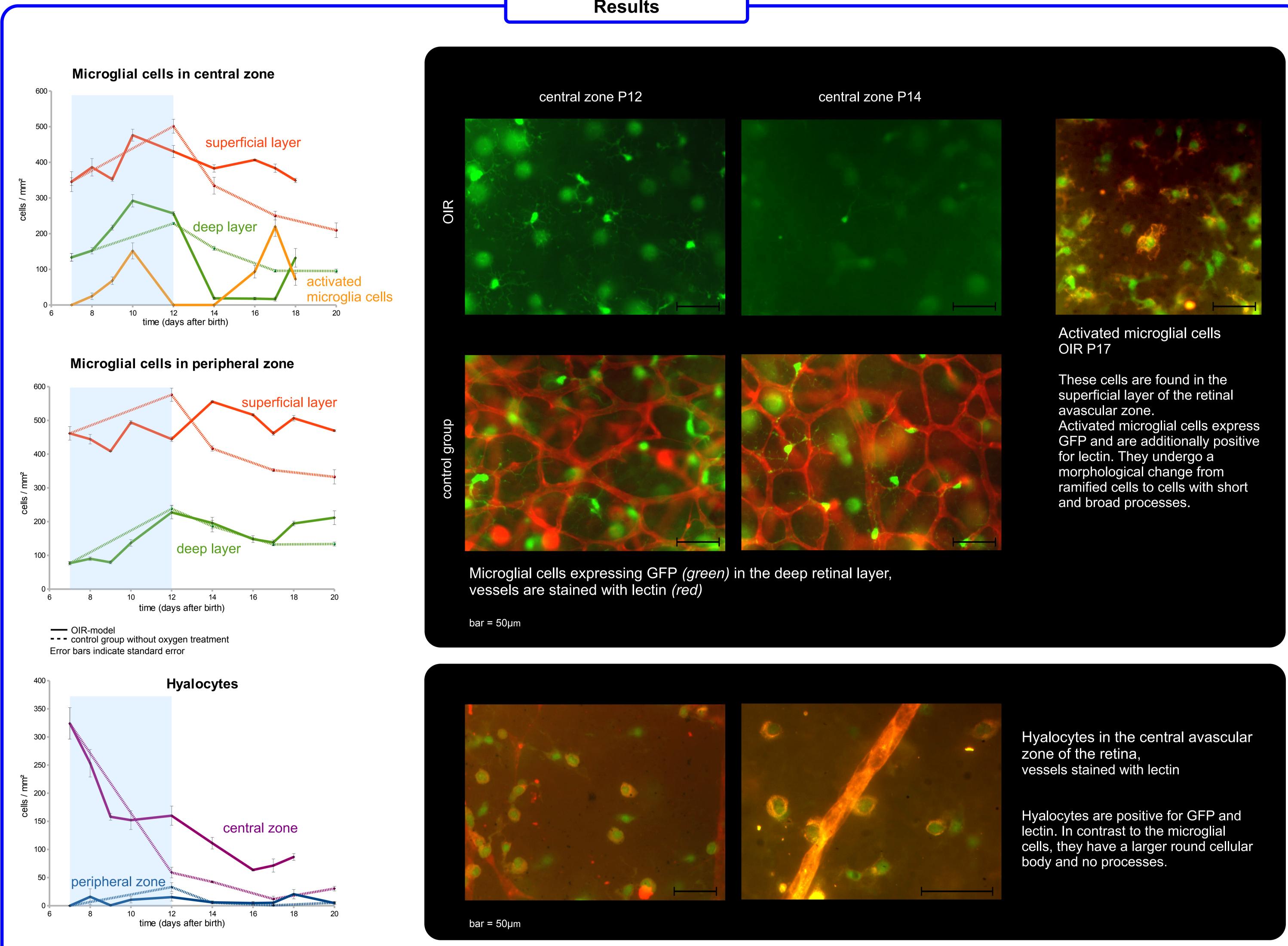


Background

Retinal neovascularization has been intensively investigated in the mouse model of oxygen-induced retinopathy (OIR). Here, we studied the contribution of microglial cells to cell loss and reactive angiogenesis in mice expressing green fluorescent protein (GFP) under the microglial cell-specific promotor Cx3cr1.

Conclusion

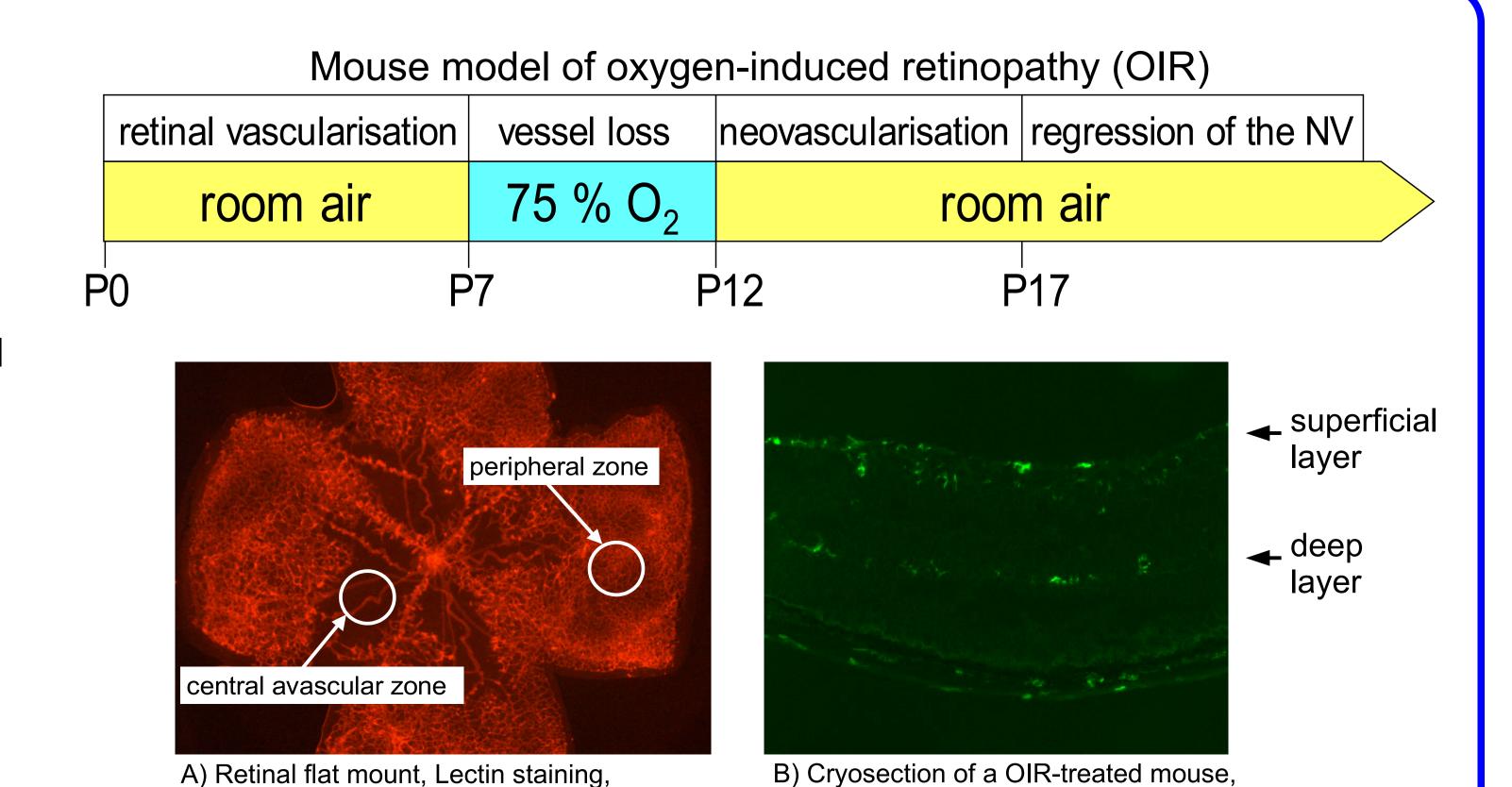
- Microglial cell density is always higher in the superficial than in the deep retinal layer.
- Hyalocytes density decreases with retinal vascularisation.
- The density of microglial cells in the deep layer decreases significantly after return to normal room air.
- It is unclear if microglial cells move from the deep to the superficial layer.
- Activated microglial cells are observed in the superficial retinal layer from P8 to P10 and from P16 to P18.
- Thus, microglial cells may be involved in the formation of the central avascular zone as well as in retinal revascularisation.



OIR P17

Methods

- Animals: heterozygous Cx3cr1-GFP mice (background: C57Bl/6)
- Retinal flat-mounts were stained with lectin and investigated by fluorescence microscopy
- Numbers of microglial cells and hyalocytes in 12 fields were determined from postnatal day 7 to day 20
- Cells were counted:
 - A) in two zones of the retina (central and peripheral zone)
 - B) in two layers of the retina (superficial and deep layer)
 - C) hyalocytes in vitreous



OIR P17, GFP+ microglial cells (green)