MR pathway in retinal health and diseases

Francine Behar-Cohen¹ and Min Zhao²

¹1 Assistance Publique - Hôpitaux de Paris, Hôpital Cochin Ophtalmopole, 75014 Paris, France 2 Centre de Recherche des Cordeliers, Sorbonne Université, Université de Paris, Inserm, From physiopathology of retinal diseases to clinical advances, 75006 Paris, France. ²Centre de Recherche des Cordeliers

September 25, 2021

Abstract

In the retina, mineralocorticoid receptor (MR), expressed in vessels, glial and neuronal cells, is mainly activated by glucocorticoids. Under pathological conditions, ocular MR expression and corticoids change, leading in most cases to MR overactivation. Experimental models using MR agonists or antagonists, administered systemically or intraocularly, acutely or chronically and transgenic models, allowed to identify the deleterious consequences of MR pathway overactivation. Among them, oxidative stress, inflammation, deregulation of hydro-ionic channels, alteration of choroidal vasculature, angiogenesis and cell death, are common to major retinal diseases. Specific MR antagonists showed efficacy in models of diabetic retinopathy, ischaemia, retinal and choroidal angiogenesis and in models of glaucoma. It is highly likely that MR antagonists will find a place in the therapeutic arsenal of age-related macular degeneration, diabetic retinopathy, glaucoma and in pachychoroid associated diseases. Their use in humans is still limited by the need of biomarkers of MR activation and specific ocular formulations.

Hosted file

 $\begin{tabular}{ll} BJP Behar-Cohen.docx & available & at & https://authorea.com/users/436453/articles/538755-mr-pathway-in-retinal-health-and-diseases \\ \end{tabular}$

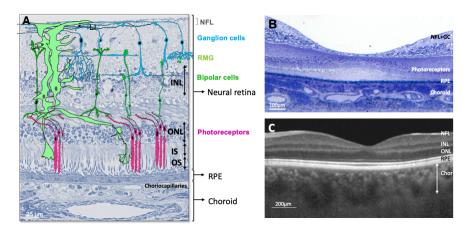
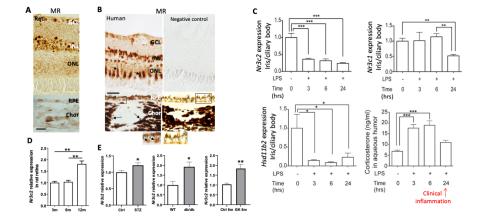


Fig 1



Fiugure 2

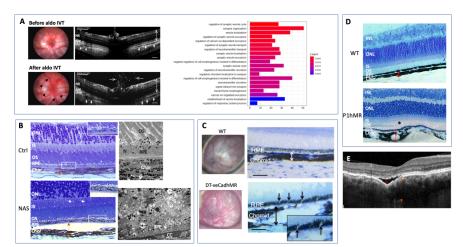


Figure 3

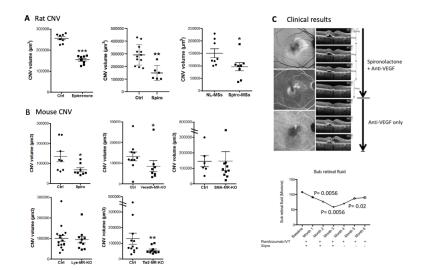


Figure 4

