

Therapy for retinal degeneration in ciliopathies

Pharmacological therapy to slow significantly retinal degeneration linked to protein trafficking defect inducing a Unfolded Protein Response

KEYWORDS

*Photoreceptor cell,
Retinitis pigmentosa,
Cellular protein overload,
Endoplasmic reticulum
stress,
Apoptosis,
rare diseases*

PATENTS

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INVENTORS

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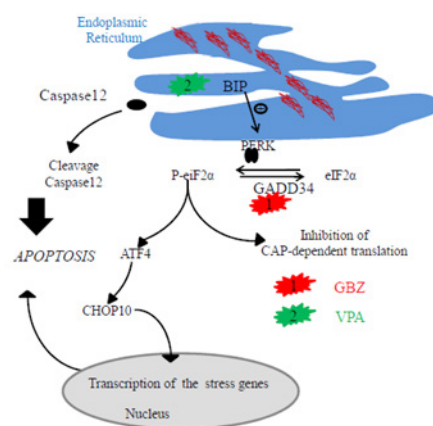
Laboratory
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TECHNOLOGY

Impaired intraciliary transport observed in Bbs12^{-/-} murine model

- results in protein retention in the endoplasmic reticulum
- this leads to a proapoptotic unfolded protein response (UPR) triggering death of the photoreceptors
- BiP, and PERK-mediated phosphorylation of eIF2α have been identified as 2 key UPR actors (Mockel et al., 2012)
- GV-ReT is a combination of clinically validated molecules (Valproic Acid and Guanabenz) for the treatment of cilia-related retinal degeneration



APPLICATION

- Pharmacological therapy for ciliopathy related retinal degeneration

INNOVATION ADVANTAGES

- A combination of two approved drugs, with a well defined MoA
- Effective in protecting retinal cells from dying and significantly increasing light detection capacity
- Aims at preserving the vision loss and slowing the retinal defects to cure secondarily with a more targeted therapy
- Positions on early onset of the retinal degeneration
- Applicable to different paradigms of retinal degeneration : UPR-inducing ciliopathies, retinal degeneration linked to protein trafficking defect inducing a UPR

DEVELOPMENT STATUS

- Compelling ex vivo and in vivo data establishing the efficacy of this drug combination
- Efficacy assessed on different cilia-related retinal degeneration (Bardet Biedl syndrome, Leber Congenital Amaurosis)

Partnership : program available for out-licensing

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