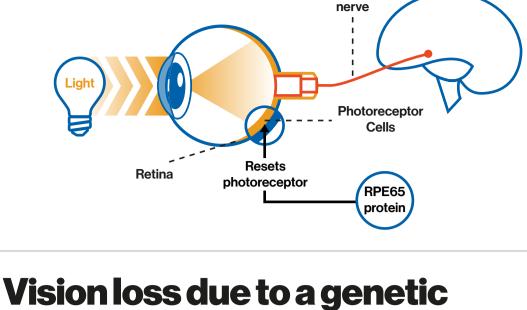
How does Luxturna®* (voretigene neparvovec) work?

What is the visual cycle?

The visual cycle is the process that converts light entering the eye into electrical signals that are transmitted to the brain¹.

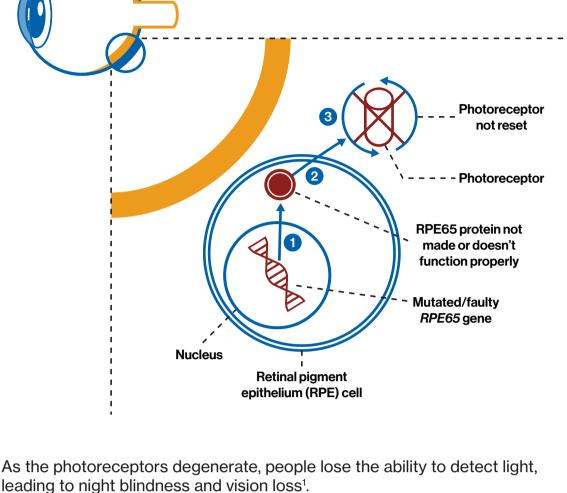
A particle of light (photon) hits a photoreceptor in the retina - the

light-sensitive tissue that lines the back of the eye - triggering electrical signals along the optic nerve to the brain¹. However the photoreceptor needs to be reset to be ready for the next photon¹. A protein called **RPE65** is involved in this task1. **Optic**



mutation in both copies of the RPE65 gene If the RPE65 gene carrying the instructions to make the protein is mutated/faulty, the protein won't be made or will not function properly¹. Therefore, the photoreceptors are unable to reset and will start to

degenerate over time^{1,2}.



What is Luxturna?

and who have enough viable retinal cells. This mutation is ultra-rare, affecting

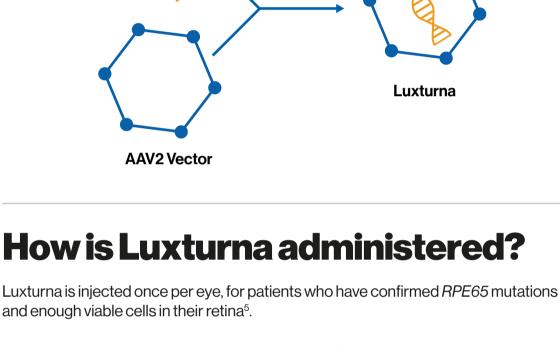
Each person has two copies of the RPE65 gene3. Luxturna is a one-time gene therapy for patients with vision loss due to a genetic mutation in both copies of the RPE65 gene,

approximately 1 in 200,000 people worldwide⁴. Luxturna provides a working copy of the RPE65 gene to act in place of the mutated

RPE65 gene

Luxturna consists of a piece of DNA - containing a working copy of the RPE65 gene that is packaged inside a transporter known as a vector, which is made from a modified, inactivated virus². Working copy of the

RPE65 gene². This working gene has the potential to restore vision and improve sight².



Retina

Luxturna

A specialised retinal surgeon injects the gene therapy behind the retina,

The Luxturna vector enters the RPE cells and delivers a working copy of

enters the **RPE cells**

Delivers working copy of RPE65 gene to the nucleus

What does Luxturna do in the eye?

the RPE65 gene to the cell nucleus⁵. The cells then begin producing the RPE65 protein⁵.

into the subretinal space⁵.

RPE cell

RPE cells

Luxturna vector

RPE65 protein expressed How does Luxturna restore vision and improve sight? Luxturna provides a working copy of the RPE65 gene to act in place of the mutated RPE65 gene². The working RPE65 gene creates the RPE65 protein⁵, thus restoring the visual cycle, with the potential to restore vision and improve sight².

Photoreceptor reset **RPE65** protein produced **Functioning** Mutated/faulty RPE65 gene RPE65 gene Retinal pigment **Nucleus** epithelium (RPE) cell

Important Safety Information

Some patients who received Luxturna experienced red or painful eyes sensitivity to light, an eye infection, cataracts, increased pressure in the eye, or temporary visual disturbances, like flashes or floaters, worsening of or blurred vision. Some of these may be related to the procedure used to inject Luxturna. This information is not comprehensive. For full information please see the EU Summary of

1. NIH U.S. National Library of Medicine (2018), Genetics Home Reference. RPE65 gene. Available at: https://ghr.nlm.nih.gov/gene/RPE65. Last accessed November 2018

 Russell S et al. Efficacy and safety of voretigene neparvovec
(AAV2-hRPE65v2) in patients with RPE65- mediated inherited retinal dystrophy: a randomised, controlled, open-label, phase 3 trial. The Lancet 2017; 390:849-860. Available at:

https://www.ncbi.nlm.nih.gov/pubmed/28712537 NIH U.S. National library of medicine. What is a gene? Available at: https://ghr.nlm.nih.gov/primer/basics/gene 4. Novartis. Data on File. 2018 S. Luxturna™ (voretigene neparvovec) Novartis Pharmaceuticals. Approved EU SmPC. Available imminently at: https://www.ema.europa.eu/en/medicines

^{*} Luxturna is a trademark of Spark Therapeutics, Inc. in the United States and is registered in the EU Luxturna is indicated for the treatment of adult and paediatric patients with vision loss due to inherited retinal dystrophy caused by confirmed biallelic RPE65 mutations and who have sufficient viable retinal cells.