Cure ADOA Foundation

The Cure ADOA Foundation is committed to help people with a severe hereditary disease of the optic nerve, named autosomal dominant optic atrophy (ADOA). In ADOA, the optic never gets thinner and this causes low vision or even blindness. In addition, some people also develop other symptoms like hearing loss or muscular spasms, this is called the ADOA-plus syndrome.

Treatment and cure

Since November 2018, the Cure ADOA Foundation has been committed to support fellow sufferers and their families. Our goals: enable scientific research, awareness of the condition, interaction between everyone involved and create a trusted place for patients. Our ultimate goal is to prevent further disease development and in the ideal situation: find a cure.

Do you know someone with ADOA?

We want everyone who is suffering of ADOA, or their loved ones, to be able to reach out to us. We have a website (Dutch, English, German) www.adoa.eu and a closed Facebook Group (mainly Dutch, but foreign people are also welcome). You can register with us as a patient by emailing us at info@adoa.eu. If you are an ophthalmologist, clinical geneticist, family member or loved one, you can also sign up for our newsletter. Extensive information about ADOA and the foundation can be found on our website www.adoa.eu .

This way, we hope to support everyone who is involved with ADOA!

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Algemeen Nut Beogende Instelling ANBI