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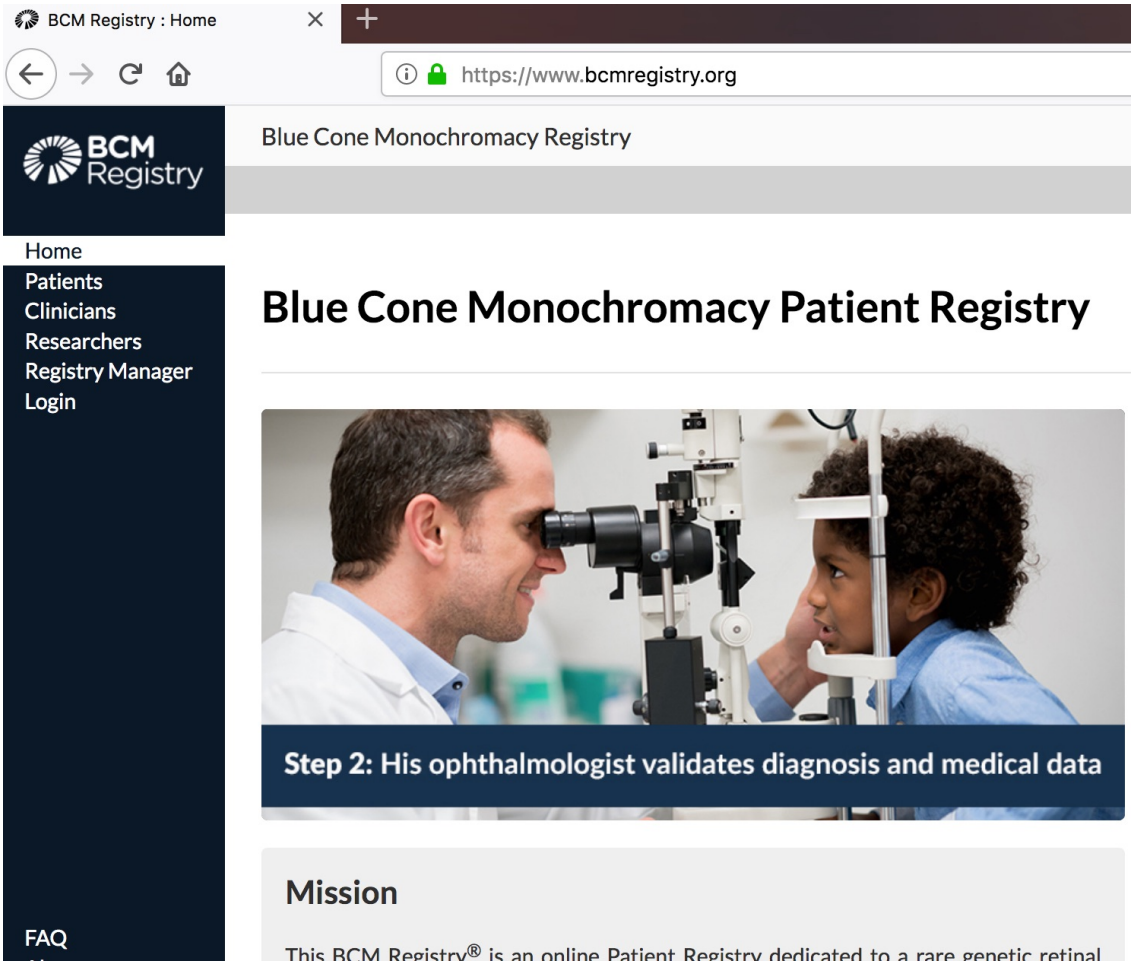
BCM FAMILIES FOUNDATION

Toward the cure of Blue Cone Monochromacy

PO Box 7711 - Jupiter FL 33468-7711 USA

Register now



It is with extreme joy that we inform you that our BCM Patient

Registry is ready and active online.

While working on creating this Registry during the past months, you were on our mind. The Registry is for you and for your children; it has taken tremendous efforts, including your contribution and support. We want to thank you for this.

If you are a person who has a genetic diagnosis of Blue Cone Monochromacy, we invite you to join the BCM Registry. The information collected in the Registry is expected to advance our understanding of the disease and to foster research into the treatment of BCM.

The overall aim of this project is to increase our knowledge about the clinical manifestations of the disease, including the prevailing clinical features and if and how they occur differently in the affected population, its natural history and long-term outcomes with or without treatments.

Individuals (including minors) with a confirmed genetic diagnosis of Blue Cone Monochromacy are eligible for participation in the BCM Registry.
During the registration process, you will be requested to upload your medical records (i.e. genetic test) and to indicate the name of your clinician.

You will find all the details on how to enroll, FAQs and contacts on the Registry webpage:

www.BCMRegistry.org

Some advice for your registration:

- **select a username that doesn't contain your name or your surname;**
- **only patients can register, female carriers cannot;**
- **if you are enrolling a child, make the registration under his name.**

Download the BCM Registry Flyer here:

Register now

Project supported with the Otto per Mille funds of Chiesa Valdese

Our mailing address is:
info@bcmfamilies.org