



Today take action!

1. On May 20th let's celebrate the International Blue Cone Monochromacy Day by joining the BCM Registry at www.BCMRegistry.org

The BCM Registry is an online Patient Registry dedicated to a rare genetic retinal disease, [Blue Cone Monochromacy](#). The Registry was created by the [BCM Families Foundation](#), a 501(c)(3) non-profit patient-led organization with the mission to cure BCM.

Anyone diagnosed with Blue Cone Monochromacy is encouraged to join the BCM Patient Registry. Individuals (including minors) with a confirmed genetic diagnosis of Blue Cone Monochromacy are eligible for participation in the BCM Patient Registry. You will be requested to upload your medical records and to indicate the name of your clinician.

Find out more at <https://www.bcmregistry.org/>

Join the Registry

2. On May 20th let's celebrate the International Blue Cone Monochromacy Day by donating to the BCM Families Foundation.

BCM Families Foundation is a non-profit organization entirely dedicated to the cure of this rare eye disease. 100% of donations collected go directly to support the most promising medical research aimed at treating or curing BCM.

Donate



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info@bcmfamilies.org

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