

## International Awareness Day

BCM Families Foundation

Thu 5/18/2023 3:47 PM

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**Today take action!**

**1. On May 20<sup>th</sup> let's celebrate the International Blue Cone Monochromacy Day by joining the BCM Registry at [www.BCMRegistry.org](http://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/>

And watch the Registry Manager Tutorial here: <https://youtu.be/3unnr3O5euM>

## Join the Registry

### **2. On May 20<sup>th</sup> let's celebrate the International Blue Cone Monochromacy Day by donating to the BCM Families Foundation.**

BCM Families Foundation is the only non-profit organization totally dedicated to the cure of this rare eye disease. 100% of donations that we collect go exclusively to fund the most promising medical research which have as their purpose the treatment or the cure of BCM.

## Donate



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