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A photograph of a neon sign that reads "DO SOMETHING GREAT" in all caps. The sign is composed of several rectangular panels, each containing a portion of the text, and is set against a dark background.

**What are you waiting for?
Join the BCMFF patients' registry!
www.BCMRegistry.org**

Why should you register?

BCM affects only 1 person out of 100,000 and, at present, has no cure. Because of the low number of patients and the fact that they

the disease is limited and so is the likelihood to develop innovative disease-modifying therapies soon.

The BCM Registry was established to fill these gaps.

Who can register?

Adults and children affected by BCM may participate in the BCM Registry. You must have genetic confirmation of BCM (DNA Test Report).

Please note that BCM Family Foundation funds the DNA test at the Tubinga Laboratory so that patients don't need to pay such an expensive test by their own.

How to register?

The registration procedure is easy and fast.

- You will need to create a username and a password and to provide an email address.
- Once your email address is confirmed, you will be guided through the registration process that requires you to:

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Give your consent for participation in

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the Registry;

- Provide your or your child's contact details;
 - Select your clinician;
 - Upload your or your child's BCM DNA test report.
- Once your account is complete you will enter in a “waiting-list” until your clinician confirms the BCM diagnosis and validates your medical data.

At that point you or your child become a Registry Patient and will contribute to advance the research to identify a cure for the BCM!

For further information please visit <https://www.bcmregistry.org/patients/>

The BCM Patient Registry is possible through the generosity of [Otto per Mille Chiesa Valdese](#)

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