

# **Do you or your child suffer from Lymphatic Malformation?**

## **Register for the upcoming online LM Registry**

The LM Registry will be the first online global patient database. It will gather much needed information regarding the parents' medical history, social/environmental/work related exposure, and possible other causes. Information found will also supplement current research and genetic testing.

We are requesting patients (over the age of 18) and/or parents to supply contact information so you may be notified when registry is operational. Our goal is to have the database running by December 2011. The online LM registry will protect patient's medical records and patient's privacy.

**For information:**

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