



Celebrating our 10<sup>th</sup> Year!

You are invited to join other MLD Families, Researchers, and Experts for a weekend of interaction and education at the world's first international

## MLD Family Conference™

**WHAT:** Learn from and talk with the world's premiere MLD researchers, meet, interact, and share with other MLD affected families

**WHERE:** Munich, Germany

**WHEN:** Friday afternoon March 13th through Saturday night March 14th, 2009

**AGENDA &** Conference details, full agenda, and registration information can be found online at: <http://MLDfoundation.eu/munich> (English)

**REGISTRATION:** or <http://MLDfoundation.de/munich-de> (German)  
or if you do not have computer access call +1 503-656-4808

The MLD Foundation celebrates the start of its 10th year serving families with its first ever international **MLD Family Conference™**. Leading researchers in Enzyme Replacement and Gene Therapies will be presenting along with experts in practical day-to-day care for those affected with MLD. Families will have plenty of time to talk with each other and the doctors.

Register early - space is limited. Registration includes meals and two nights lodging. You are encouraged to bring your MLD affected family member. A staffed respite room will be provided. We are trying to accommodate requests for interpretation.

*The MLD Foundation was formed in May 2001 to serve families affected by metachromatic leukodystrophy (MLD), a terminal neuro-metabolic genetic condition affecting people throughout the world. Our slogan, We C.A.R.E.™ reflects our culture of Compassion, Awareness, Research, and Education. For more information about the MLD Foundation and metachromatic leukodystrophy please visit us at [www.MLDfoundation.eu](http://www.MLDfoundation.eu)*

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WE C.A.R.E.

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