

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™

HOTEL ROOM
AVAILABILITY AND
PRICING SUBJECT
TO CHANGE AFTER
FEBRUARY 12TH

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

The MLD Foundation is headquartered in West Linn, Oregon, USA and is a registered (501)(3) non-profit organization. MLD Family Conference and We C.A.R.E. are trademarks of the MLD Foundation.

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WE C.A.R.E.

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