



Cologne, April 2008

## INVITATION

### 6. Meeting of the German Primary Hyperoxaluria Self support group

Dear Patients and Parents!

I would like to invite you to our annual "Patients- and Parents" meeting, which is again held at the Clementine Kinderkrankenhaus in Frankfurt. You know that PH is a very rare disease. This should be reason for all patients and parents in Europe and worldwide to fight together for a timely and certain diagnosis and most importantly effective treatment opportunities for the primary, but also for the secondary hyperoxalurias.

This year physicians from several European countries also take part on the meeting. They work together for the research on primary Hyperoxaluria and recently founded the European Hyperoxaluria Consortium

Our meeting takes place at:

**Clementine-Kinderkrankenhaus, Theobald-Christ-Str. 16, 60316 Frankfurt/Main**

Date: **17. Mai 2008,** Time: **10.00 a.m.**

### Meeting Itinerary

**10.00-10.30 am** Get together

**10.30-11.30 am** News from the bench:

**Dr. B. Beck: Genotype/Phenotype**

**Dr. M. Feldkötter: Non PH I/II patients**

**11.30-11.45 am** PH Patients in Poland / Dr. P. Sikora

**11.45-12.00 am**                      **Secondary hyperoxalurias / B. Hoppe**

**12.00-1.00 pm**                      **Lunch break**

**1.00-5.00 pm**                      **1) Self support group meeting**

**2) Industry presentations**

**a) Altus presentation: Enzyme**

**treatment**

**b) AMT presentation: Gene therapy?**

**~ 5 pm**                                      **Adjourn**

If you require more information, please feel free to contact me any time:

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[info@ph-selbsthilfe.org](mailto:info@ph-selbsthilfe.org)

Im looking forward to see you at the meeting!

With my best regards

Mike Dreibrodt  
President  
German PH-self support group