



2ND EUROPEAN SYMPOSIUM ON
RARE ANAEMIAS
13-14 MARCH 2008

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Program

Venue:

The Cyprus Institute of Neurology and Genetics
Nicosia, Cyprus



TUESDAY, MARCH 13 2008

09:00 Welcome:
Introduction to the Second European Symposium on Rare Anaemias
Marina Kleanthous

09:15 ENERCA presentation
Joan Lluís Vives Corrons

Session 1:
Current Progress in Prenatal and Neonatal Diagnosis of Haemoglobinopathies

09:30 Prenatal diagnosis for thalassaemia and other haemoglobinopathies. New approaches
Marina Kleanthous (Cyprus)

10:00 Preimplantation genetic diagnosis therapy in beta-thalassaemia and other haemoglobinopathies
Mireille Claustres (France)

10:30 Neonatal screening for sickle cell anaemia and other haemoglobinopathies in Europe
Béatrice Gulbis (Belgium)

11:00 Genetic Counselling on Haemoglobinopathies. New Challenges.
Patricia Aguilar Martinez (France)

11:30 *Coffee break*

Session 2:
Iron overload and chelation therapy

12:00 Advances in iron metabolism
Clara Camaschella (Italy)

12:30 Updating iron chelation therapy in thalassaemia
Maria Domenica Cappellini (Italy)

13:00 Chelation Therapy in Thalassaemia. The Cyprus experience
Soteroula Christou (Cyprus)

13:30 *Lunch*



TUESDAY, MARCH 13 2008

Session 3:

Treatment of haemoglobinopathies. Current situation

- 15:00 Transfusion therapy
Antonio Piga (Italy)
- 15:30 Stem cell transplantation
Franco Locatelli (Italy)
- 16:00 2 selected oral presentations
- 16:30 *Coffee break*
- 17:00 Patients Associations in the prevention of Thalassaemias and other haemoglobinopathies
Androulla Eleftheriou - TIF Representative



FRIDAY, MARCH 14 2008

Session 4:
Red blood cell hereditary disorders

- 09:00 General approach for the identification and diagnosis of hereditary RBC membrane defects
Leticia Ribeiro (Portugal)
- 09:30 Stomatocytosis and allied disorders. An update
Jean Delaunay (France)
- 10:00 RBC enzyme defects. New challenges
Joan LLuis Vives Corrons (Spain)
- 10:30 Standards for laboratory diagnosis of rare anaemias
Andrea Mosca (Italy)
- 11:00 *Coffee break*

Session 5:
Very rare anaemias

- 11:30 Anaemia and porphyria
Carole Beaumont (France)
- 12:00 New inherited microcytic hypochromic anaemias
Achille Iolascon (Italy)
- 12:30 Congenital Dyserythropoietic Anaemias. Diagnosis and therapy.
Hermann Heimpel (Germany)
- 13:00 *Lunch*

Session 6 :
Free Communications Session

- 14:30 - 15:30 Oral presentation of the six top quality selected posters
- 15:30 Sideroblastic anaemias
Mario Cazzola (Italy)