

LUNCH SYMPOSIUM

“Emerging Therapies for Pompe Disease”

Pompe Disease is also called Acid Maltase Deficiency or Glycogen Storage Disease Type II

Date: 11 May 2005 (during Myology 2005)

Location: Nantes, France

Lunch will be served before the session from 12h30 to 13h20

Chairman: Dr. Pascal Laforêt (Paris, France)

- 13h20 – 13h30: Introduction (Dr. P. Laforêt, Paris - France)
- 13h30 – 13h45 : Infantile-Onset Population: Natural History and Results with Enzyme Replacement Therapy
(Prof. M. Nicolino, Lyon - France)
- 13h45 – 14h00: Natural Course in Late-Onset Pompe Disease - Data from an International Survey and First Results on Enzyme Therapy
(Dr. A. van der Ploeg, Rotterdam – The Netherlands)
- 14h00 – 14h15: Management of Late-Onset Patients in France and Outcomes of an International Late-Onset Prospective Observational Study
(Dr. P. Laforêt, Paris - France)
- 14h15 – 14h30: Outcome of the 1st ENMC Workshop: Defining the Standards of Medical Management in Pompe Disease
(Prof. V. Straub, Newcastle - UK)
- 14h30 – 14h45: Gene Therapy for Pompe Disease: Pathway to Clinical Studies
(Dr. B. Byrne, Florida - USA)
- 14h45 – 15h00: Panel discussion