



# **NEUROMED : Progress Meeting, Toulouse 5 & 6th March 2015**

Institut de Pharmacologie et de Biologie Structurale (IPBS) Centre National de la Recherche Scientifique UMR 5089 DR14 205 route de Narbonne, 31077 Toulouse, France

5 March 2015

13:30 h: Welcome and gathering

## 14:30 h: Announcements and Important Issues (by Javier Sancho)

15:00 h: NEUROMED Science presentations by the six partners: where we are. (20-30 min)
I Javier Sancho. Universidad de Zaragoza. IUI BIFI.
Salvador Ventura. Universidad Autónoma de Barcelona
Maria Joao Saraiva. Instituto de Biología Molecular e Celular. IBMC 16:30 Coffee break
Rui M. M. Brito. Universidad de Coimbra
Pau Bernardó. INSERM. Dèlégation Régionale Languedoc-Roussillon Centre de Biochimie Structurale
Marie L. Maddelein. CNRS UMR5089 IPBS

16:30 Coffee break

18:00 h: Approval of coordinators, steering committees, and other Neuromed management documents and Administrative Tasks

### 20:00 h: NEUROMED dinner

6 March 2015

9h30: Welcome

### 10h : Tasks and delivrables : state of developement of each scientific GT

#### 10h: GT2. Phenylketonuria: organizing the task (by Javier Sancho)

1. Computational design of molecules based on pharmacological chaperone IV using the three-dimensional structure obtained by x ray diffraction

2. Analysis of the folding and assembly of wild PAH and of defective enzymes as well as the effects exerted by

chaperone IV and Kuvan

- 3. Synthesis and in vitro assay of new designed chaperones
- 4. Toxicity and efficacy in animal models

10:45 Coffee break

## 11h: GT3. Parkinson: organizing the task (by Salvador Ventura)

1. Screening of chemical libraries and identification of inhibitors of alpha-synuclein aggregation

2. Analysis of the mechanism of alpha-synuclein inhibitors

3. Design, synthesis and in vitro analysis of fluorescent inhibitors capable of binding to Alpha-synuclein oligomers

4. Effect of alpha-synuclein oligomerization inhibitors in the neurodegenerative process

5. Analysis of fluorescent inhibitors in human specimens from patients with synucleopathies

11:45 Coffee break

## 12h: GT4. Amyloidosis of TTR: organizing the task (by Rui M.M. Brito and Maria Joao Saraiva)

1. Virtual and experimental screening of chemical libraries and identification of new pharmacological chaperones

and inhibitors of TTR aggregation

2. Computational design and production of trans-suppressors of aggregation

3. Experimental analysis of the mode of action of pharmacological chaperones and TTR trans-suppressors found

in actions 1 and 3

4. Design, synthesis and in vitro test of conjugated inhibitors with affinity by TTR

5. Toxicity tests in cell cultures of agents selected in action s1-4

6. Toxicity and efficacy in mice models of the most promising pharmacological chaperones and inhibitors selected in action 5

## 13 h: Lunch

**15h 00 : General Discussion** 

16h End of Progress Meeting