

Annual Thierry Latran Foundation Grantees meeting 2018
SAID BUSINESS SCHOOL - Park End Street
Seminar Room A

Tuesday 19 June

14h30 Welcome coffee

15h	Wim Robberecht	Introduction
15h15 - 15h45	Final presentation N. Charlet-Berguerand	Model-ALS : Novel animal models for Amyotrophic Lateral Sclerosis
15h45 - 16h15	Final presentation G. Schiavo	ALS-GOI: Deficits in axonal transport as a target for pharmacological intervention in ALS
16h15 - 16h45	Final presentation E. Hornstein	miRNAgenetics: Discovery of microRNA-related mutations in human ALS patient genomes
16h45- 17h05	Progress presentation C. Lobsiger	PeriMAC: Peripheral macrophages as a promising target to modulate ALS disease progression.
17h05 - 17h30	Break	
17h30 - 17h50	Progress presentation D. Zytnicki	SPIN-ALS : Can a spinal interneuronopathy contribute to motoneuron degeneration in ALS? An electrophysiological study on patients and mouse models of ALS.
17h50 - 18h10	Progress presentation A. Chio/ P. Van Damme	INSPIRED: Evaluation of disease onset and disease Stage/severity in Patients with amyotrophic lateral sclerosis using FDG-pet imaging
18h10 - 18h30	Progress presentation E. Horstein	Astrocyte secretome: Investigating the role of exosomal RNA in C9orf72 astrocyte toxicity - moving towards gene therapy approaches
18h30 - 18h45	V. de Broglie	The Thierry Latran Foundation
19h	COCKTAIL DINER	The Jam Factory, 4 Hollybush Row - 1 mn

Wednesday 20 June

9h - 9h20	Progress presentation L. Van Den Bosch	ALS-HDAC6: Preclinical testing of histone deacetylase 6 (HDAC6) inhibitors in a FUS mouse model of amyotrophic lateral sclerosis
9h20 - 9h35	Initiation presentation M. Barkats / MG Biferi	GeneTherC9: New gene therapy for C9ORF72-linked ALS-FTD
9h35 - 9h50	Initiation presentation S. Corti / E. Hedlund	IS-ALS: Modulating motor neuron vulnerability using the oculomotor restricted genes IGF-2 and SYT13 for ALS therapy
9h50 - 10h05	Initiation presentation M. Otto	Aggregated-TDP43: Detection of disease specific aggregates of TDP-43 in cerebrospinal fluid of ALS patients for disease specific diagnosis
10h05 - 10h30	Break	
10h30 - 10h45	Initiation presentation K. Talbot	TDPBAC: The contribution of cell autonomous and non-cell autonomous effects of mutant TDP-43 to disease onset and progression in a novel transgenic mouse model of ALS
10h45 - 11h	Initiation presentation C. Verfaillie	ALS_OL : Identification of defects in oligodendrocytes (OL) in familial and sporadic ALS and creation of an all human OL-neuron coculture system for screens of novel drugs that correct OL defects
11h00 - 11h15	Initiation presentation J. Weishaupt	Mutimono: A multicentric approach to monocyte alterations in ALS
11h15 - 11h30	Wim Robberecht	Closing Remarks