

International mini-symposium Models and Therapies for Rare Diseases

November 23, 2018

Aula de Graus, Facultat de Biologia

17h: Alba Tristan (Institut de Recerca Sant Joan de Déu, Barcelona)

Tyrosine hydroxylase deficiency: studies in patients and in an iPSC cell model

18h: Dr. Isaac Canals (Fac. of Medicine, Lund University, Lund, Sweden)

Stem cell and CRISPR technologies to model neurodegenerative disorders

November 26, 2018

Aula de Graus, Facultat de Biologia

17h: Dr. Cecilia Jimenez (I. de Recerca S. Joan de Déu, CIBERER, Barcelona)

Genetics of neuromuscular diseases and therapeutic advances

18h: Dr. Victor Hernandez (Inst. of Child Health, Univ. College, London, UK)

Gene therapy for multisyndromic ciliopathies. Bardet-Biedl syndrome