

Biomarkers discovery in the progressive bulbar palsy/amyotrophic lateral sclerosis clinical spectrum: combined tissue and fluid-phase proteomics

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Country

United Kingdom

Title of project or programme

Biomarkers discovery in the progressive bulbar palsy/amyotrophic lateral sclerosis clinical spectrum: combined tissue and fluid-phase proteomics

Source of funding information

The Wellcome Trust

Total sum awarded (Euro)

€ 114,191

Start date of award

01/01/2014

Total duration of award in years

2

Keywords

Research Abstract

The major obstacle in the development of novel therapeutic strategies for PBP and ALS is the poor understanding of their pathophysiology. The diagnostic delay of more than 15 months from disease onset is due to the lack of firm diagnostic biomarkers and reduces significantly the

therapeutic window for any neuron-rescuing strategy. This has led to failures of clinical trials and can possibly explain the negligible effects of Riluzole, the only licensed drug for ALS. The aim of this study is to establish a panel of accessible biomarkers aiding diagnosis and capable of measuring treatment efficacy in PBP. Such a panel of biomarkers will also aid the development of more targeted therapies and better drug selection throughout the treatment phase of disease. Using an unbiased, sensitive proteomic approach based on Tandem Mass Tag@ (TMT@) labeling technology, developed by Proteome Sciences (PS), we will profile affected brain tissue and white cells collected longitudinally from PBP/ALS patients and search for similar fluid-phase biomarkers in matched longitudinal plasma samples. This study will pre-configure biomarkers assays targeting antigens and immunological factors to be tested in routine clinical practice, strengthening the potential for an early diagnosis and accurate monitoring of the rate of disease progression.

Further information available at:

Types:

Investments < €500k

Member States:

United Kingdom

Diseases:

N/A

Years:

2016

Database Categories:

N/A

Database Tags:

N/A