## **Supplementary Material**

## Impaired levels of gangliosides in the Corpus Callosum of

## **Huntington Disease animal models**

Alba Di Pardo<sup>1</sup>, Enrico Amico<sup>1</sup>, and Vittorio Maglione<sup>\*</sup>

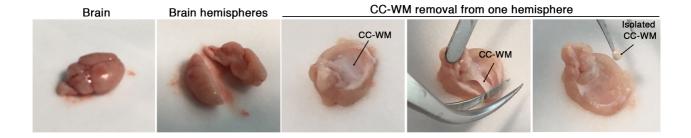
IRCCS Neuromed, Localita' Camerelle 86077 Pozzilli (IS), Italy

<sup>1</sup> These authors contributed equally to this work

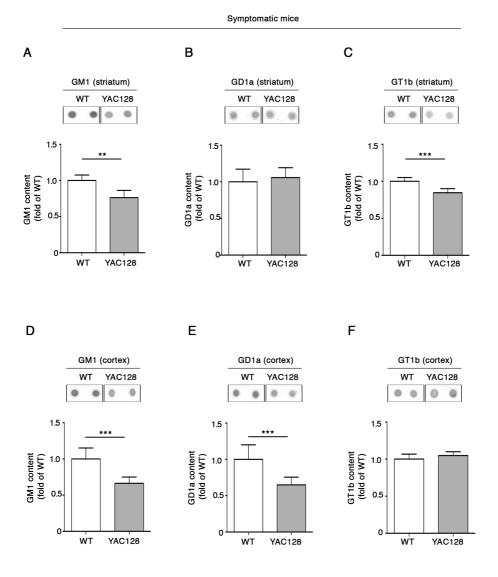
## \*Correspondence

Vittorio Maglione

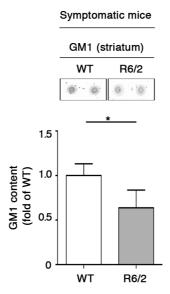
E-mail: vittorio.maglione@neuromed.it



Supplementary Figure 1. Corpus Callosum (CC-WM) isolation from mouse brain.



**Supplementary Figure 2. Brain ganglioside content is aberrant in striatal and cortical tissues of symptomatic YAC128 HD mice.** Representative dot blottings and densitometric analysis of GM1, GD1a and GT1b gangliosides in striatum (A-C) and cortex (D-E) from symptomatic (9 month old) YAC128 mice and age-matched WT littermates. Ganglioside spots were visualized by ECL. Data are represented as the mean±SD, n= 5 for each group of mice. \*\*P<0.001; \*\*\*P<0.0001 (non-parametric Mann–Whitney U-test).



Supplementary Figure 3. Levels of ganglioside GM1 are reduced in the striatum of symptomatic R6/2 HD mice.

Representative dot blottings and densitometric analysis of GM1 in striatal tissues isolated from symptomatic R6/2 mice and WT controls. Ganglioside spots were visualized by ECL. Data are represented as the mean±SD, n= 7 for each group of mice. \*P<0.05 (non-parametric Mann–Whitney U-test).