Acetylation state of RelA modulated by epigenetic drugs prolongs survival and induces a neuroprotective effect on ALS murine model

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Supplemental Figure 1



Supplemental Figure 1: The evaluation of microglial activation was performed analyzing the area covered by positive Iba1 cells in the lumbar spinal cord of WT (n=3), VEH (n=3) and HD (n=3). (a) Representative images of lumbar spinal cord of WT, VEH and HD, showed a slight decrease of microglial activation. (b) Graph show the percentage of the area covered by positive Iba1 cells. Despite a slight decrease of the positive Iba1 cells were detected not significant statistical differences were found. Magnification 20x, scale bar 100 μ m. Results were analyzed by one-way ANOVA followed by Tukey's multiple comparisons test. Data are expressed as mean±SEM.

Supplemental Figure 2





Blot 1 – RelA







Blot 3 – RelA Ac-K310

Supplemental Figure 2: Immunoprecipitation followed by western blot of RelA subunit of the nuclear fraction isolated from lumbar spinal cord of WT, VEH, LD and HD groups. On the left of *Blot 1 – RelA* and *Blot 3 – RelA Ac-K310* are shown merged blots with the protein molecular weight marker (kDa), on the right are shown the uncropped full length western blots of gels presented in Figure 4a of the main article. The figure *Blot 2 – RelA Ac-K* is displayed with the protein molecular weight marker (kDa). For all blots, the black-lines boxes referred to the cropped parts that are showed in the main article.

Supplemental Figure 3

Figure 6a





Blot 1 – pAMPK











Blot 3 - GAPDH

Supplemental Figure 3: Western blot of cytoplasmic fraction isolated from lumbar spinal cord of WT, VEH, LD and HD groups probed with pAMPK (62kDa), AMPK (62kDa) and GAPDH (37kDa) antibodies. The figure shows the uncropped full length western blots of gels presented in Figure 6a of the main article. The black-lines boxes referred to the cropped parts that are showed in the main article. In the left parts of *Blot 1 – pAMPK*, *Blot 2 – AMPK* and *Blot 3 – GAPDH* are showed merged blots with the protein molecular weight marker (kDa).

Supplemental Figure 4

Figure 7a





Blot 1 – Bcl-xL



Blot 2 – BDNF



Supplemental Figure 4: Western blot of cytoplasmic fraction isolated from lumbar spinal cord of WT, VEH, LD and HD groups probed with Bcl-xL (30kDa), BDNF (14kDa) and β -actin (45kDa) antibodies. The figure shows the uncropped full length western blots of gels presented in Figure 7a of the main article. The black-lines boxes referred to the cropped parts that are showed in the main article. In the left parts of *Blot 1 – Bcl-xL*, *Blot 2 – BDNF* and *Blot 3 – β-actin* are showed merged blots with the protein molecular weight marker (kDa).