

Figure S1. Comparisons of expression level of GTPBP10 in different brain regions of HD patients (HD) and normal controls (CTRL). *P*-values were calculated with two tailed *t*-tests. (a) GTPBP10 is significantly down-regulated in the cingulate cortex regions of HD patients compared to normal controls. The data was from AI-Dalahmah et al. 2020[1]. (b) GTPBP10 is down-regulated in motor cortex (BA4) regions of HD patients compared to normal controls (data from GSE79666)[2]. (c) GTPBP10 is down-regulated in HD-PC (BA9) compared to normal controls (data from GSE64810)[3]. (d) GTPBP10 is down-regulated in peripheral venous blood samples of HD patients compared to normal controls (data from GSE61405)[4].



Figure S2. The comparisons for the targets of the original and mutated hsa-mir-499a-3p (hsa-mir-499a_73g) and hsa-mir-3622a-5p (hsa-mir-3622a_21a), respectively. (a) The targets of hsa-mir-499a-3p and hsa-mir-499a_73g. (b). The targets of hsa-mir-3622a-5p and hsa-mir-3622a_21a.

References

- Al-Dalahmah, O., et al., Single-nucleus RNA-seq identifies Huntington disease astrocyte states. Acta Neuropathol Commun, 2020. 8(1): p. 19.
- Lin, L., et al., *Transcriptome sequencing reveals aberrant alternative splicing in Huntington's disease*. Hum Mol Genet, 2016. 25(16): p. 3454-3466.
- Labadorf, A., et al., RNA Sequence Analysis of Human Huntington Disease Brain Reveals an Extensive Increase in Inflammatory and Developmental Gene Expression. PLoS One, 2015. 10(12): p. e0143563.
- 4. Andrade-Navarro, M.A., et al., *RNA Sequencing of Human Peripheral Blood Cells Indicates* Upregulation of Immune-Related Genes in Huntington's Disease. Front Neurol, 2020. **11**: p. 573560.