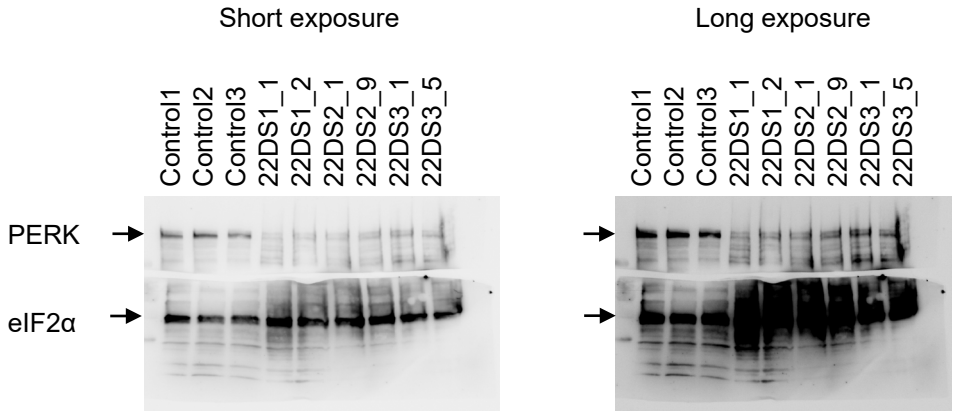
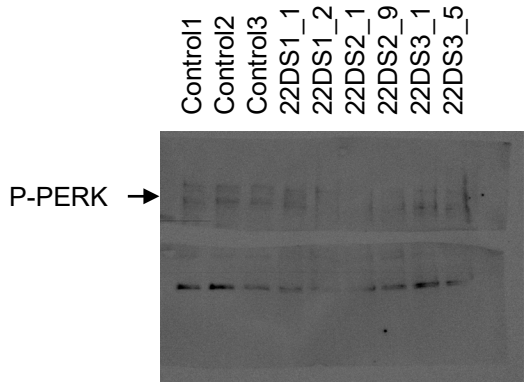
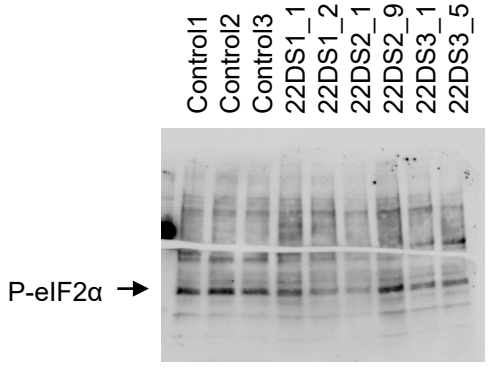


After blotting, the membrane was cut in two. The cut membrane was used for detecting PERK and eIF2α expression. The left was used for eIF2α and the right was used for PERK.

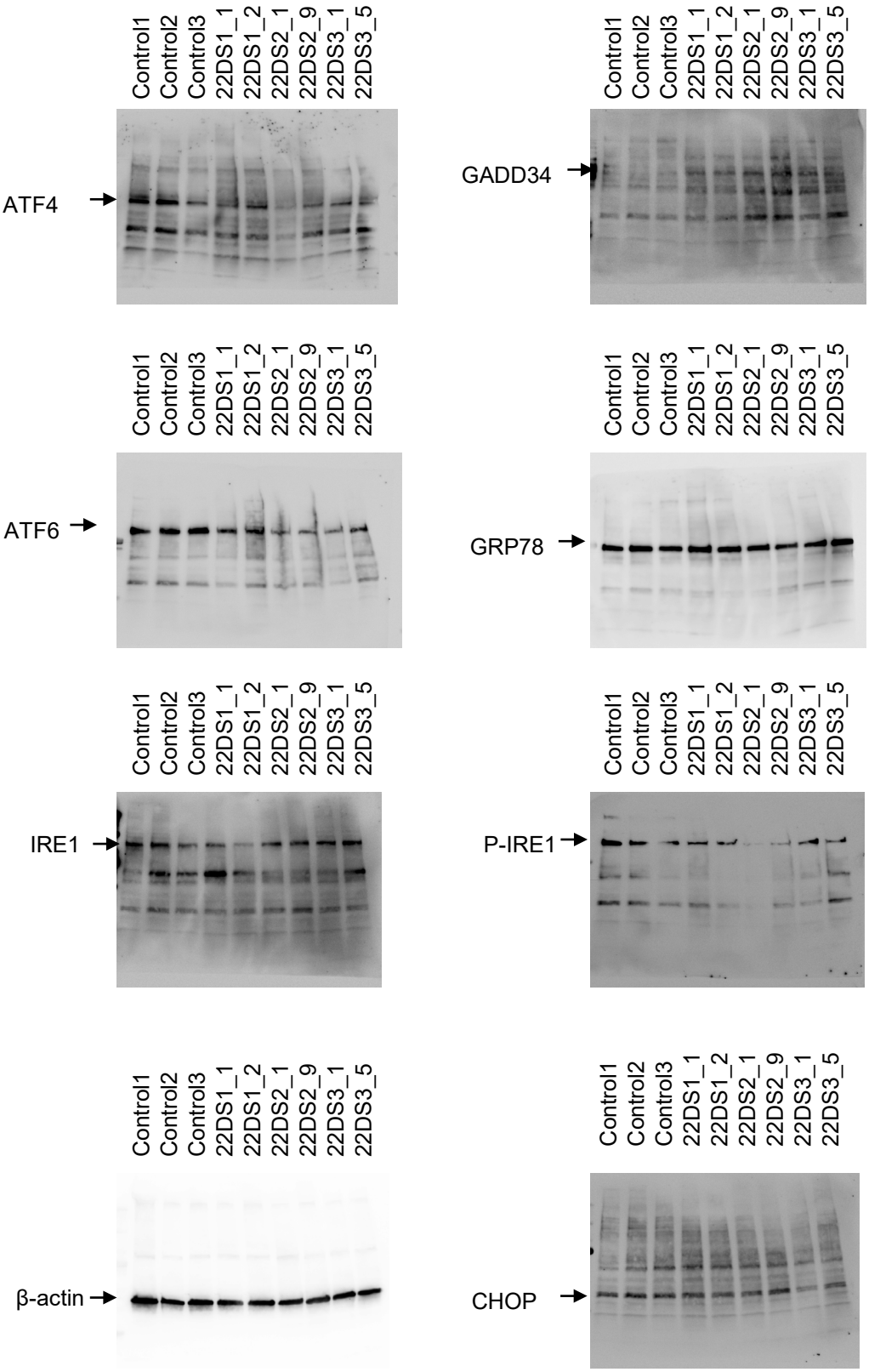


After blotting, the membrane was cut in two. The upper cut membrane was used for detecting P-PERK.

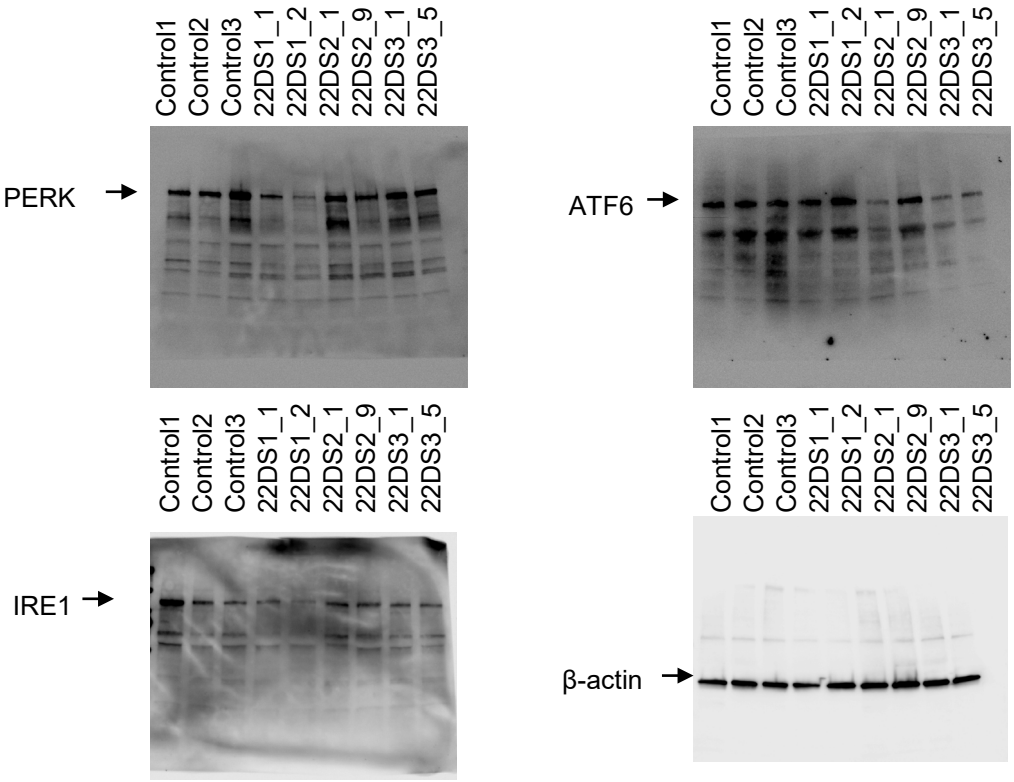


After blotting, the membrane was cut in two. The lower cut membrane was used for detecting P-eIF2α.

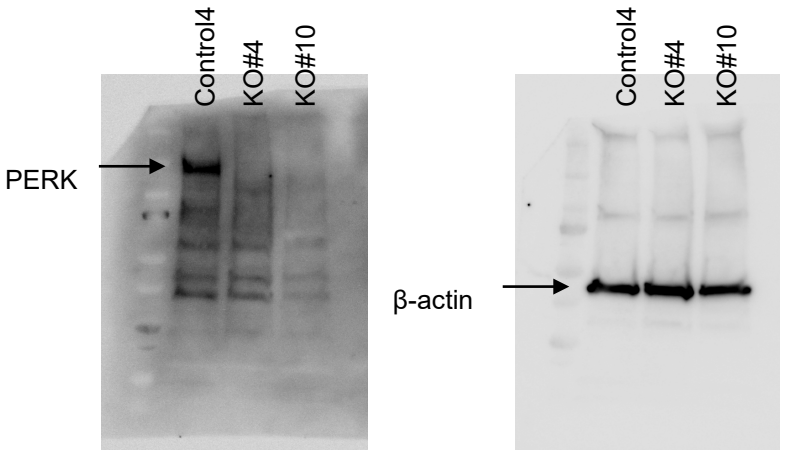
Full blots for Figure 2a



(continued) Full blots for Figure 2a

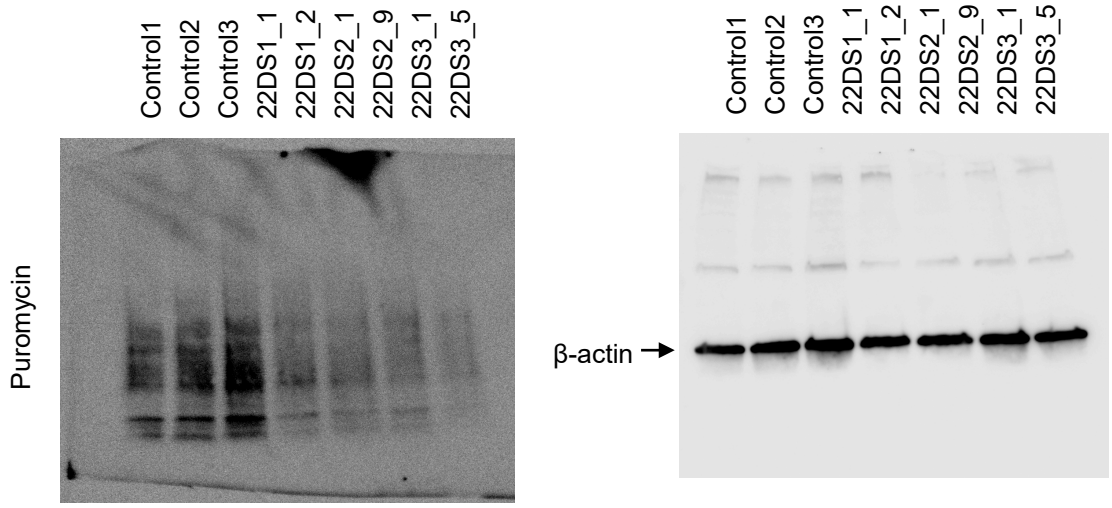


Full blots for Figure 3i



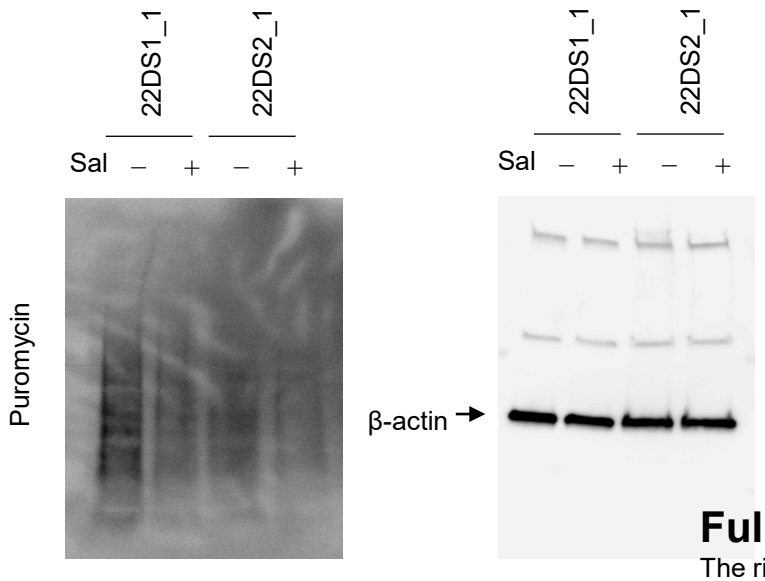
Full blots for Figure 4g

The right membrane was the same as the left.



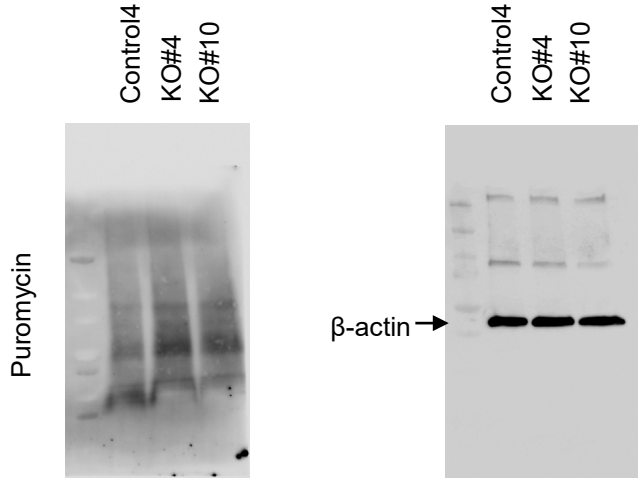
Full blots for Figure 6a

The right membrane was the same as the left.



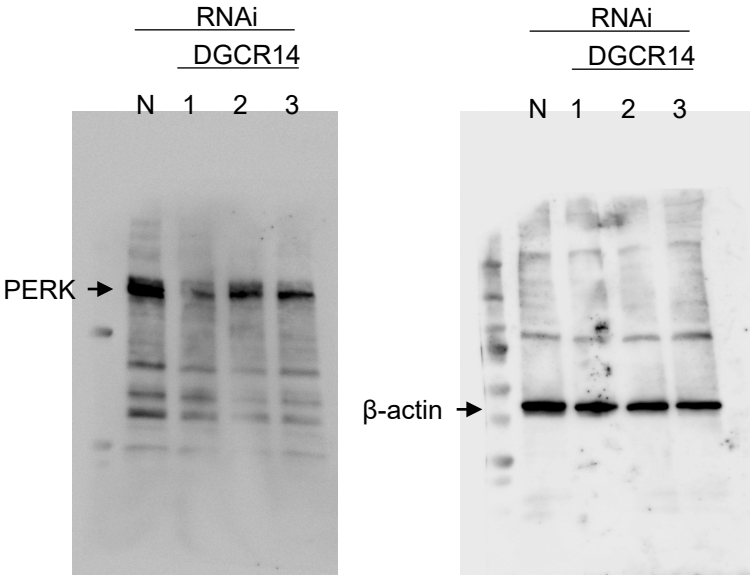
Full blots for Figure 6c

The right membrane was the same as the left.



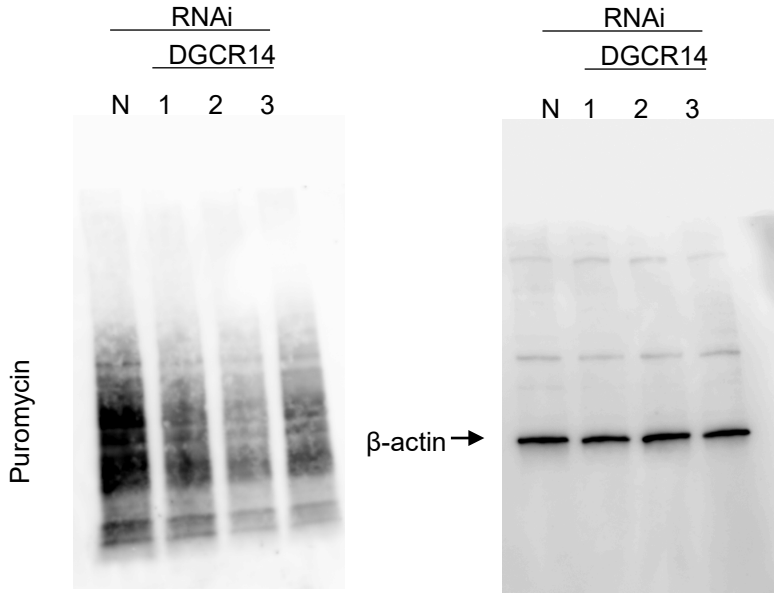
Full blots for Figure 6d

The right membrane was the same as the left.



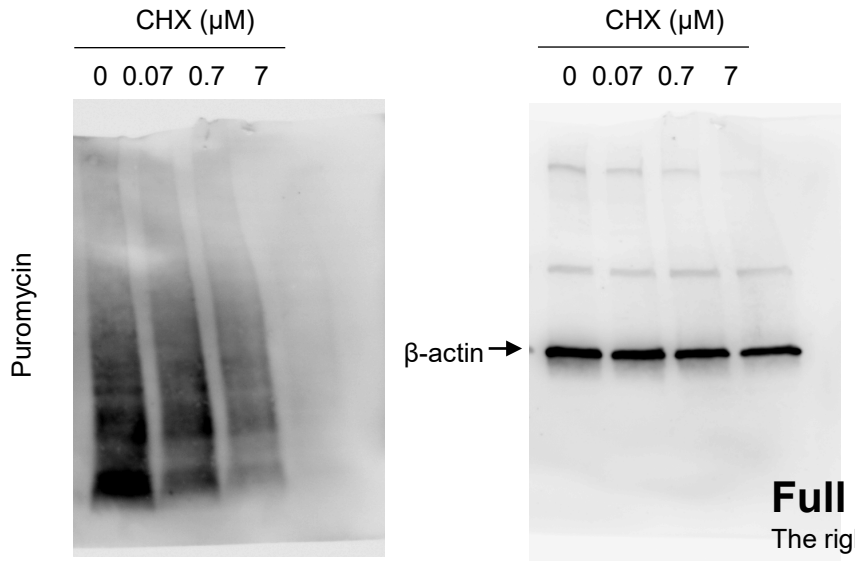
Full blots for Figure 8b

The right membrane was the same as the left.



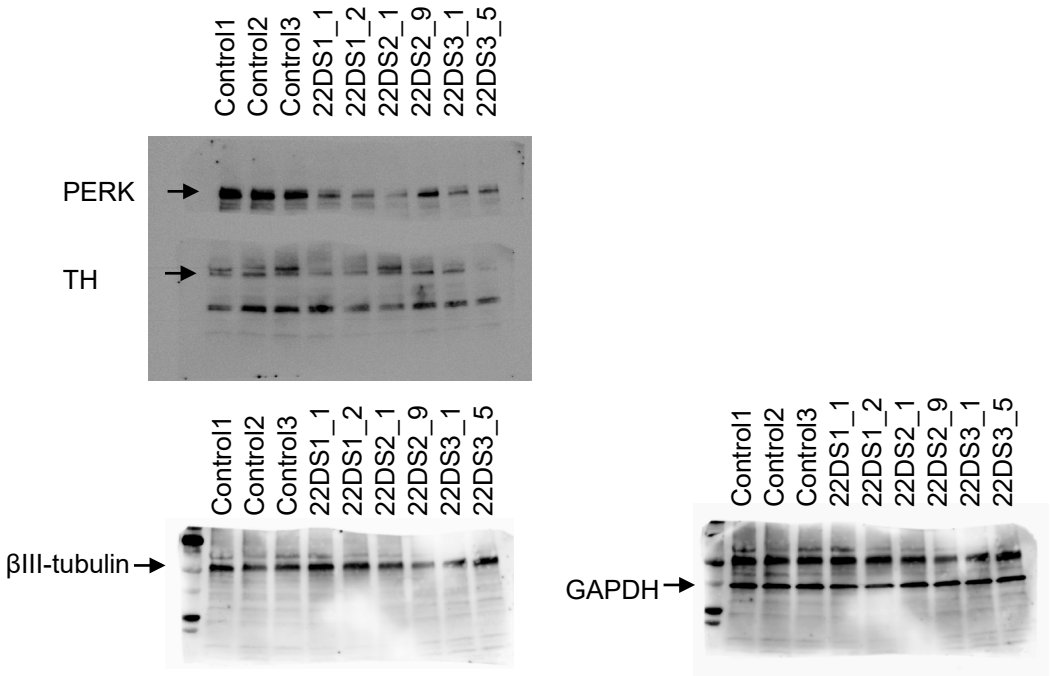
Full blots for Figure 8e

The right membrane was the same as the left.



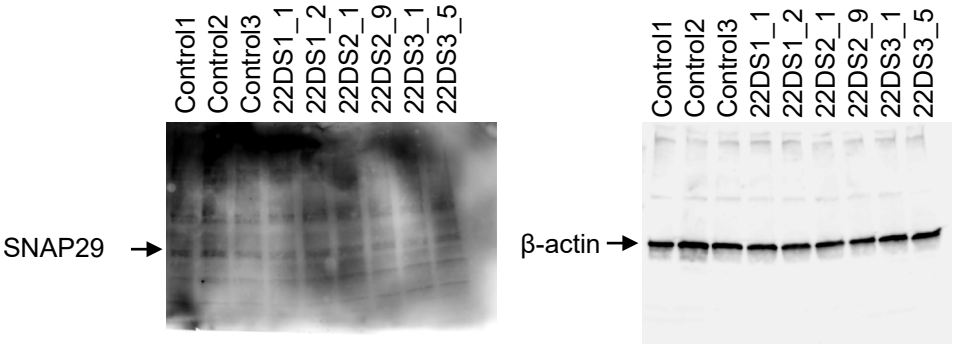
Full blots for Figure 8g

The right membrane was the same as the left.



Full blots for Supplementary Figure 6a

After blotting, the membrane was cut in two. The cut membrane was used for detecting PERK and TH expression. The upper was used for PERK and the lower was used for TH, βIII-tubulin, and GAPDH.



Full blots for Supplementary Figure 6c

The right membrane was the same as the left one.