Supporting Information

Small molecule compounds that inhibit antioxidant response gene expression in an inducer-dependent manner

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The supporting information contains five pages comprised of four figures.

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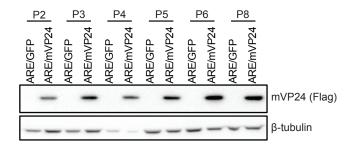


Figure S1 (related to Figure 1). Analysis of cell lysates from ARE/GFP and ARE/mVP24 HEK-293T cells at indicated passages (P) for Flag-mVP24 expression. Western blots were performed with anti-Flag antibody.

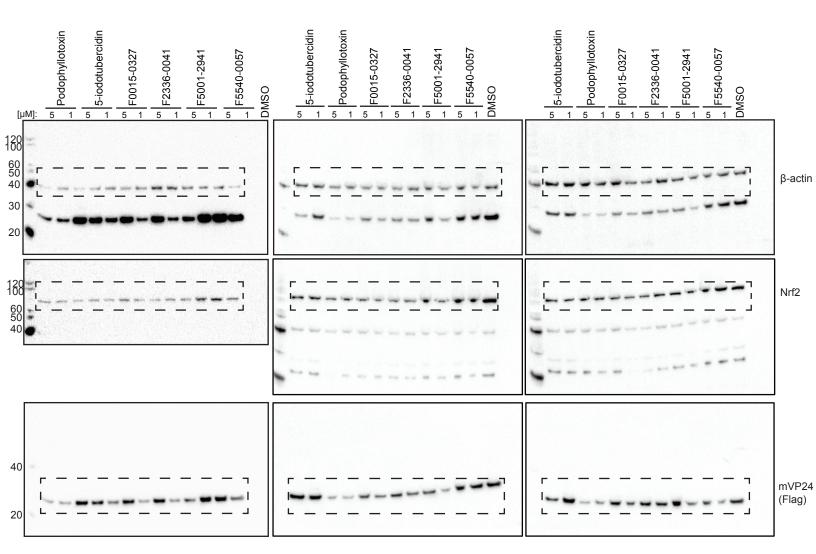


Figure S2 (related to Figure 4). Western blots used for the analysis of relative expression of Flag-tagged mVP24, Nrf2 and β -actin in ARE/mVP24 HEK293T cells treated with the indicated compounds at 5 and 1 μM for 24 h. Western blots were probed with individual antibodies sequentially; anti-Flag first, followed by anti- β -actin, and anti-Nrf2 last. Dashed boxes indicate the bands used in the analysis. The order of the podophyllotoxin and 5-iodotubercidin samples are different in the left-most image, compared with the center and right images.

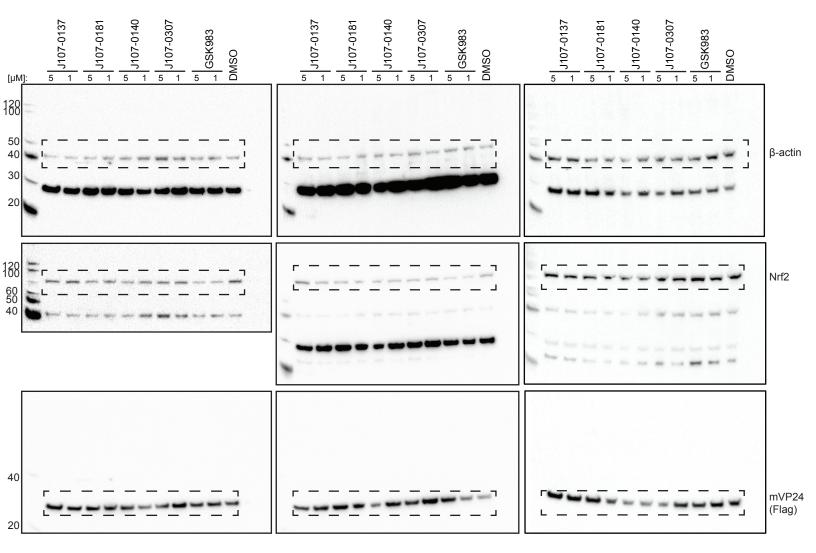


Figure S3 (related to Figure 5). Western blots used for the analysis of relative expression of Flag tagged mVP24, Nrf2 and β -actin in ARE/mVP24 HEK293T cells treated with the indicated compounds at 5 and 1 μM for 24 h. Western blots were probed with individual antibodies sequentially; anti-Flag first, followed by anti- β -actin, and anti-Nrf2 last. Dashed boxes indicate the bands used in the analysis.

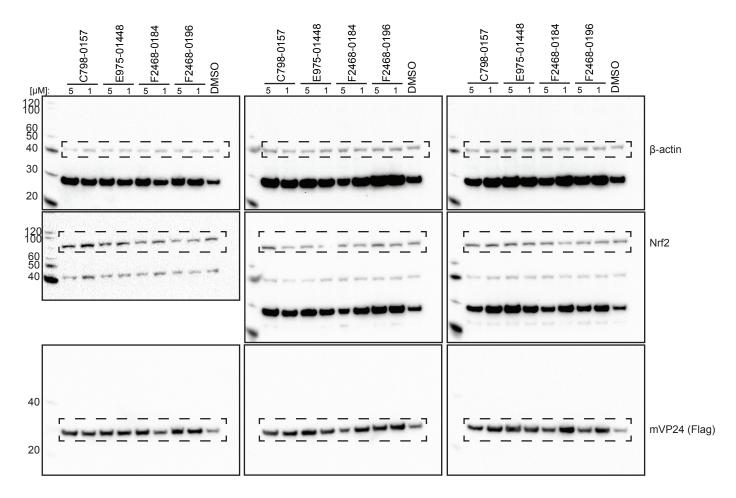


Figure S4 (related to Figure 6). Western blots used for the analysis of relative expression of Flag tagged mVP24, Nrf2 and β -actin in ARE/mVP24 HEK293T cells treated with the indicated compounds at 5 and 1 μM for 24 h. Western blots were probed with individual antibodies sequentially; anti-Flag first, followed by anti- β -actin, and anti-Nrf2 last. Dashed boxes indicate the bands used in the analysis.