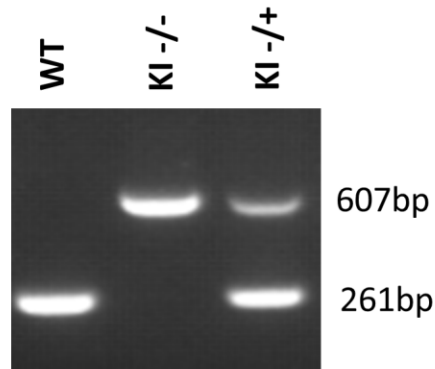


* TATCCA → GCTGCA(YP187/188AA)

U1-F:GTGGCAGTATAGATTCTCACC

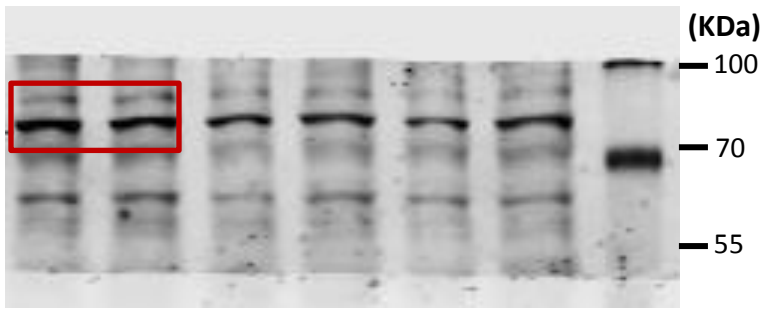
U1-R:CTACCTGGCAGCAAAGC

NEO-R:TATCGCCTTCTTGACGAGTTC

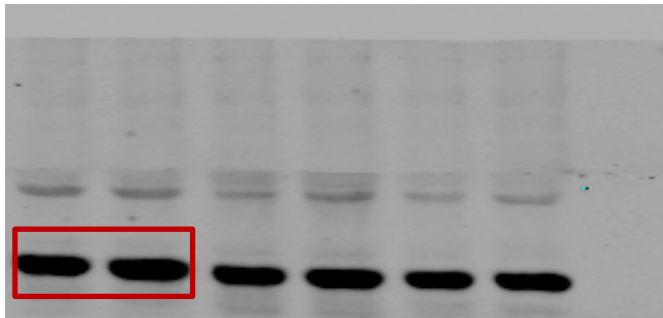


Supplementary Figure 1. Strategy for genotyping of Uhrf1 YP187/188AA KI mice by PCR. The top panel illustrates the genomic difference between WT and KI mice. The middle panel shows the primer sequences for PCR reaction. The lower panel shows a representative genotyping result for WT, homozygous KI $-/-$ and heterozygous KI $-/+$ mouse.

Figure 1C



α Uhrf1



α GAPDH

Figure 1D

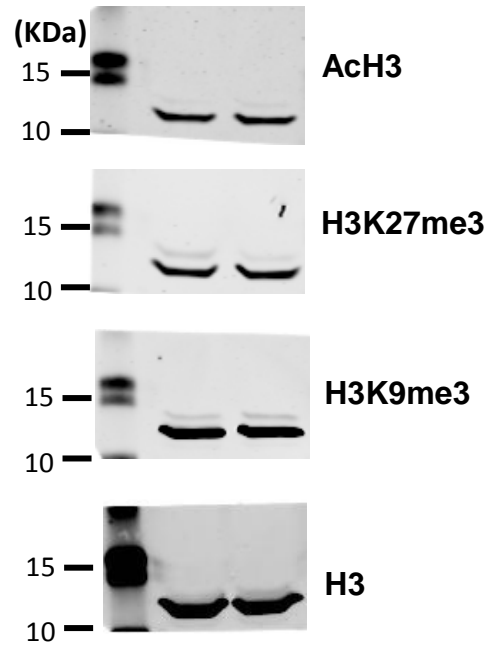
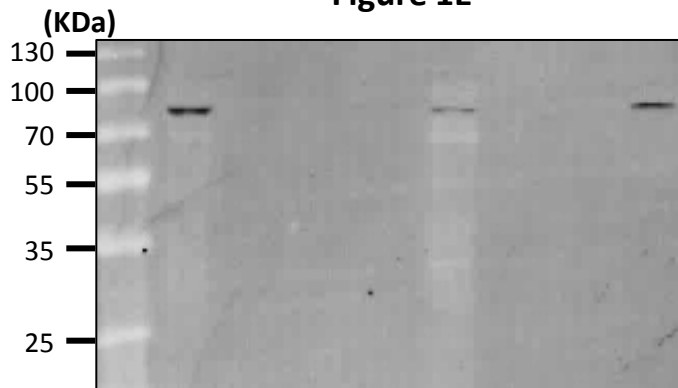
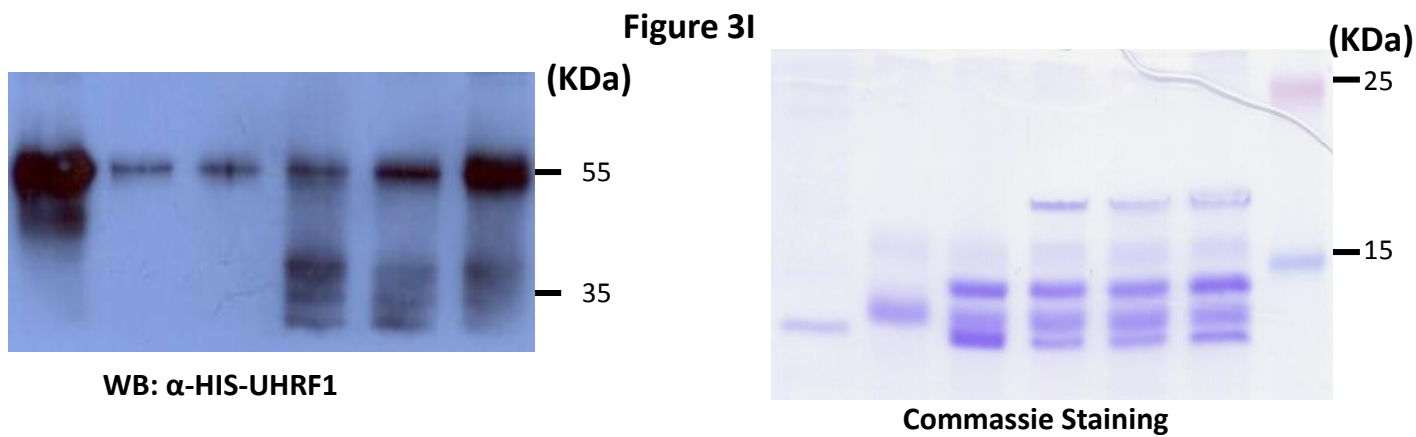
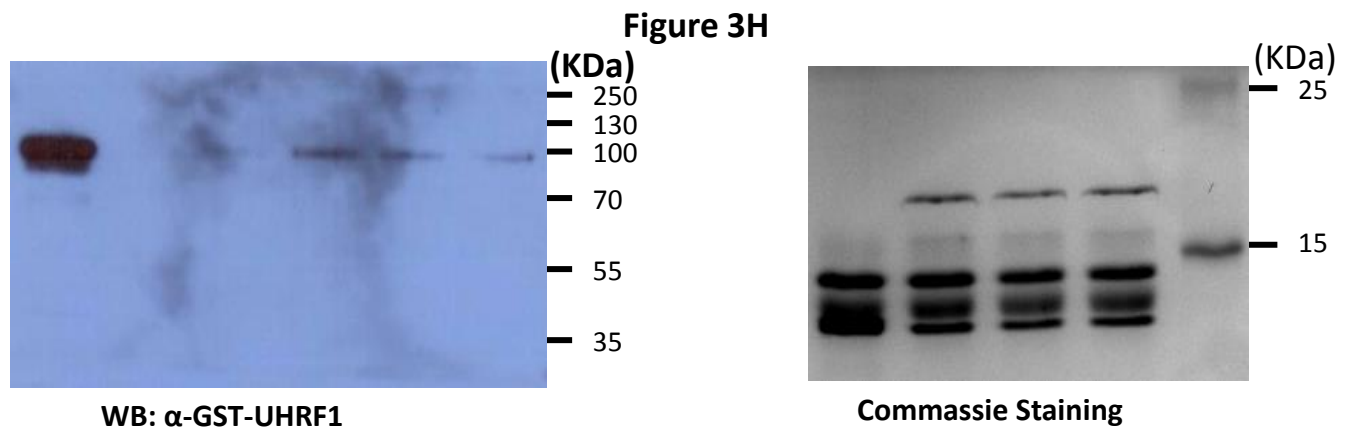
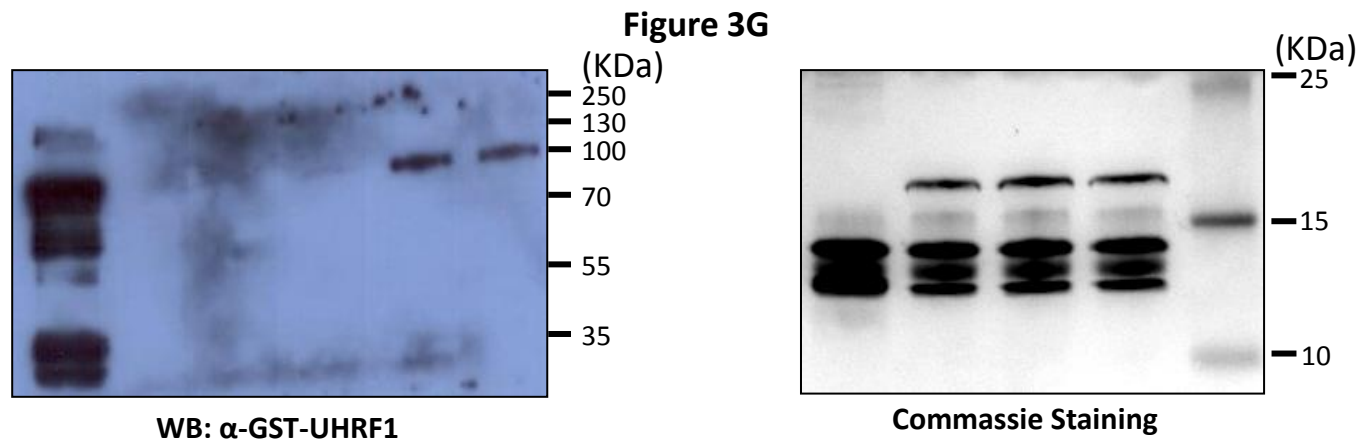
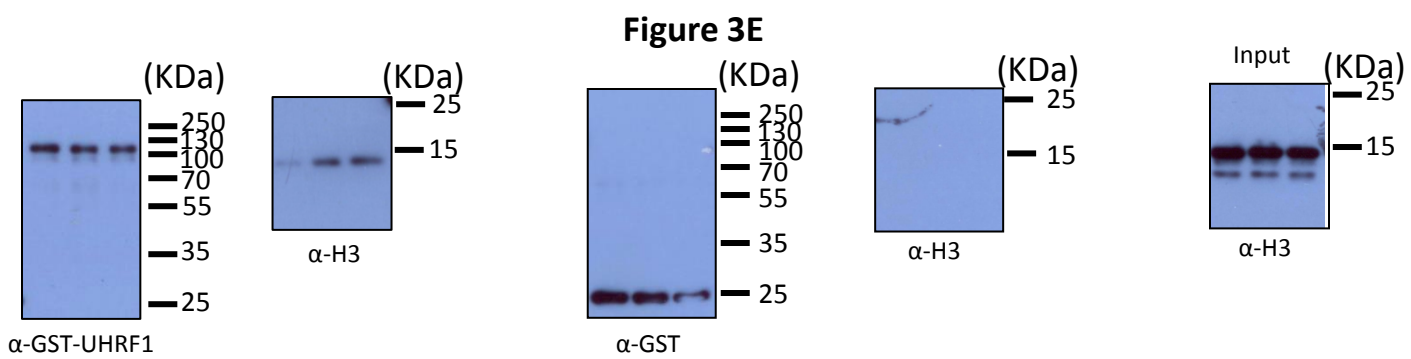


Figure 1E



α Uhrf1

Supplementary Figure 2. Whole uncropped Western blots for Figure 1C, 1D and 1E. Molecular weight markers are as indicated. Labeling and order are consistent with Figure 1.



Supplementary Figure 3. Whole uncropped Western blots for Figure 3. Molecular weight markers are as indicated. Labeling and order are consistent with Figure 3.

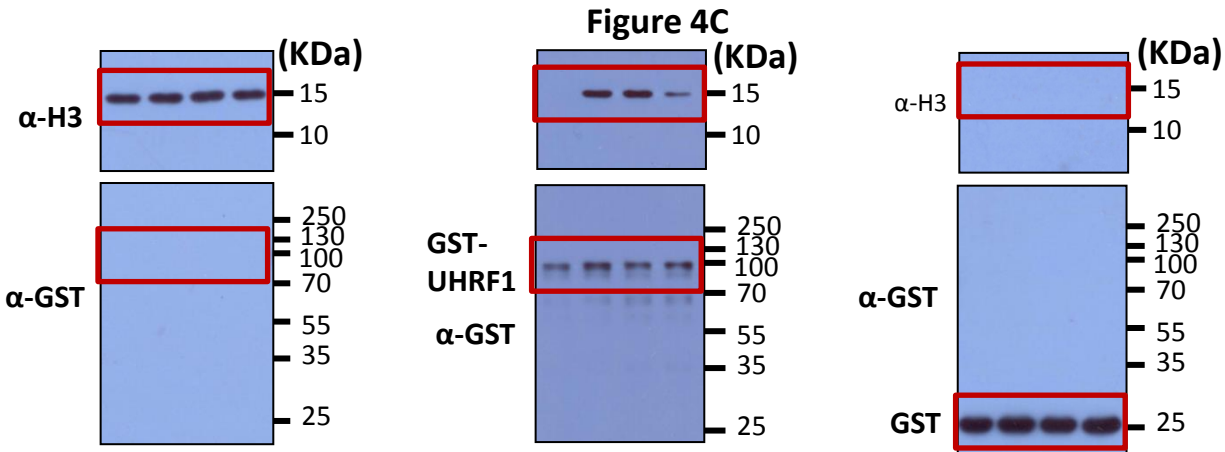


Figure 4D

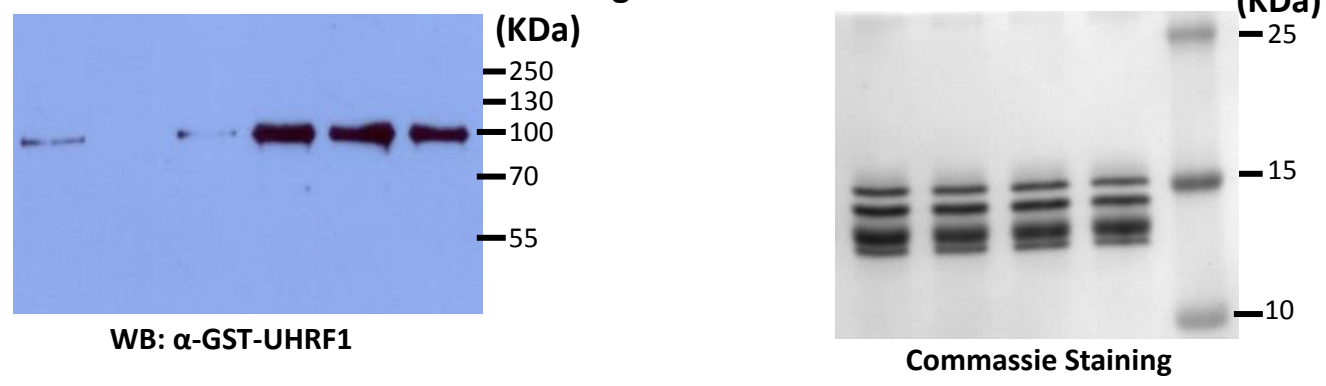
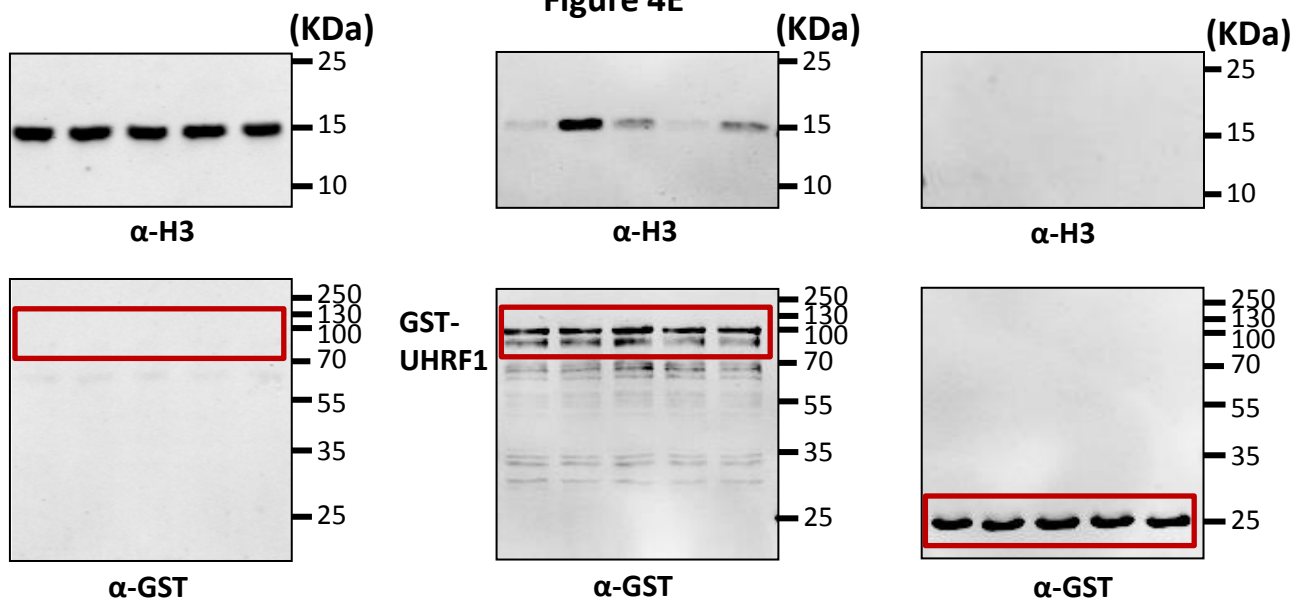
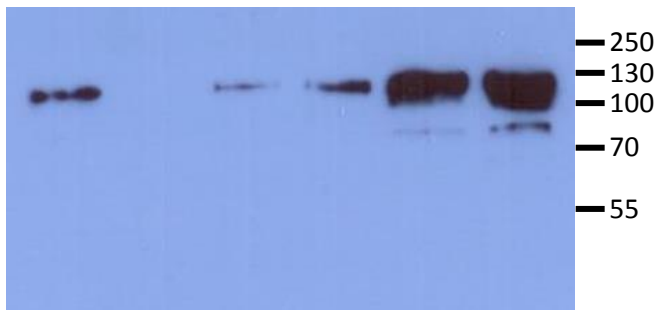


Figure 4E

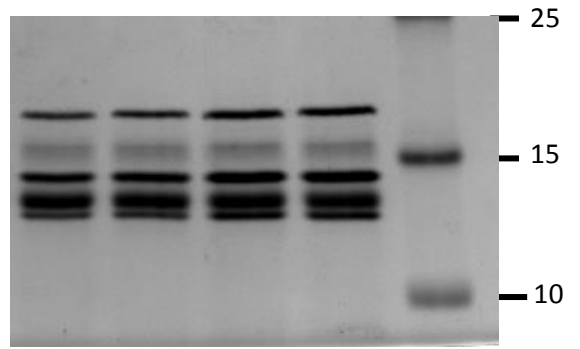


Supplementary Figure 4. Whole uncropped Western blots for Figure 4. Molecular weight markers are as indicated. Labeling and order are consistent with Figure 4.

Figure 5B



WB: α -GST-UHRF1

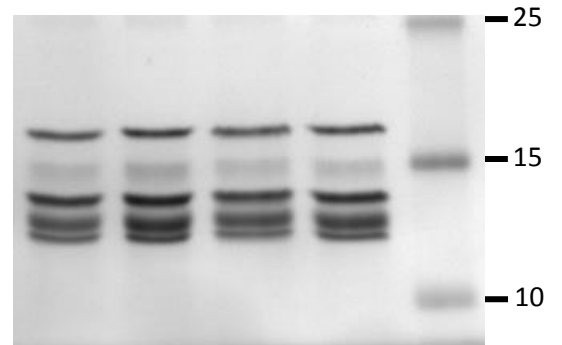


Commassie Staining

Figure 5D

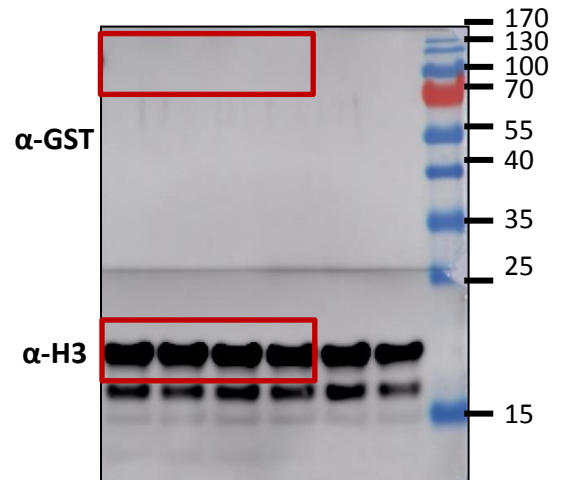
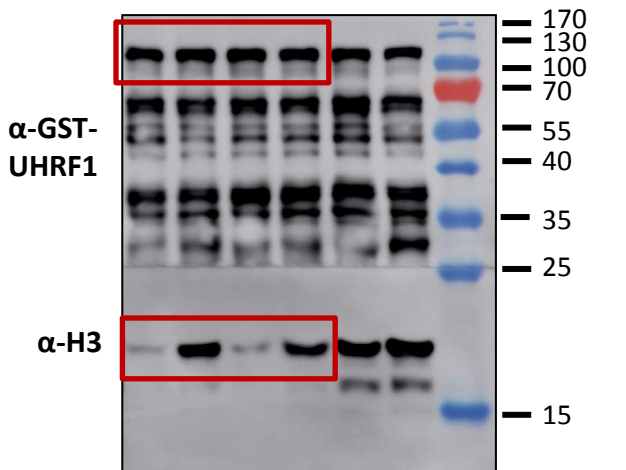


WB: α -GST-UHRF1



Commassie Staining

Figure 5F



Supplementary Figure 5. Whole uncropped Western blots for Figure 5. Molecular weight markers are as indicated. Labeling and order are consistent with Figure 5.