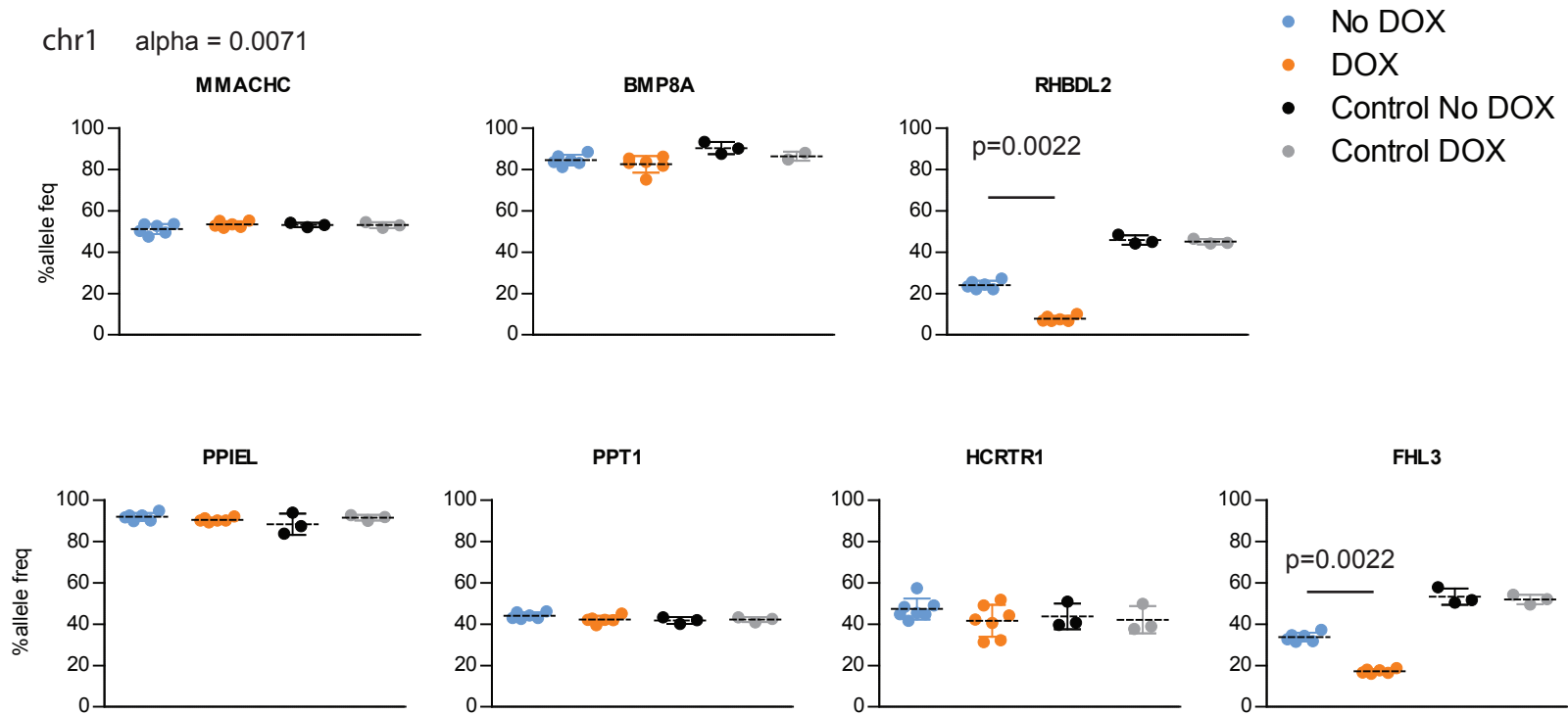


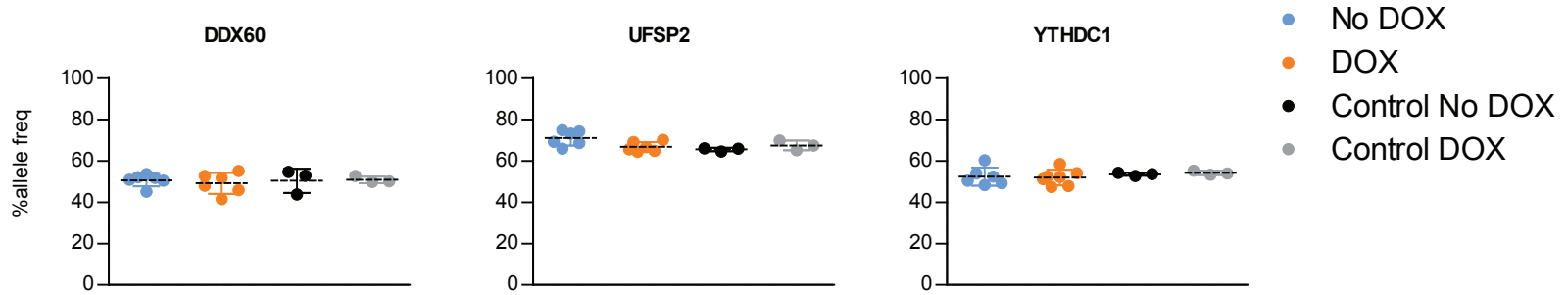
Additional Data File 3. Candidate gene silencing assays for each integration site.

a

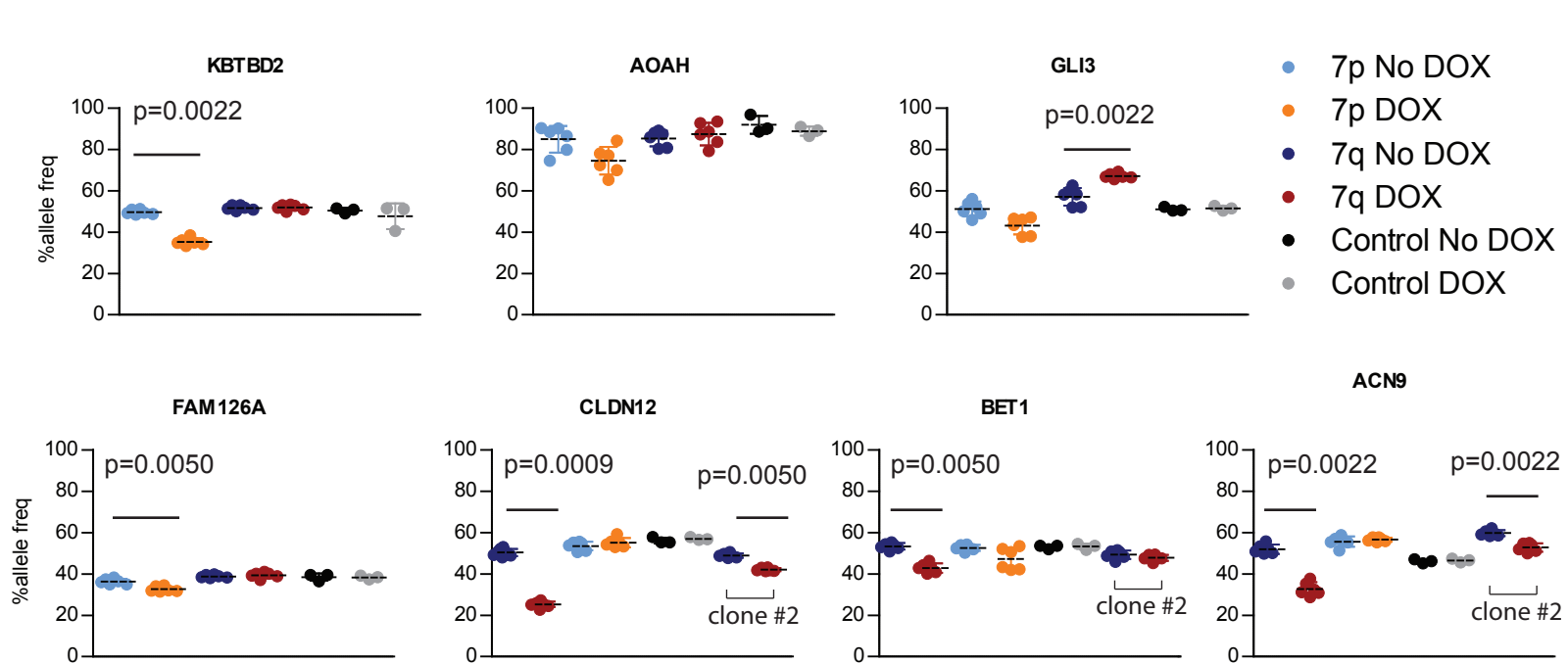
chr1 alpha = 0.0071



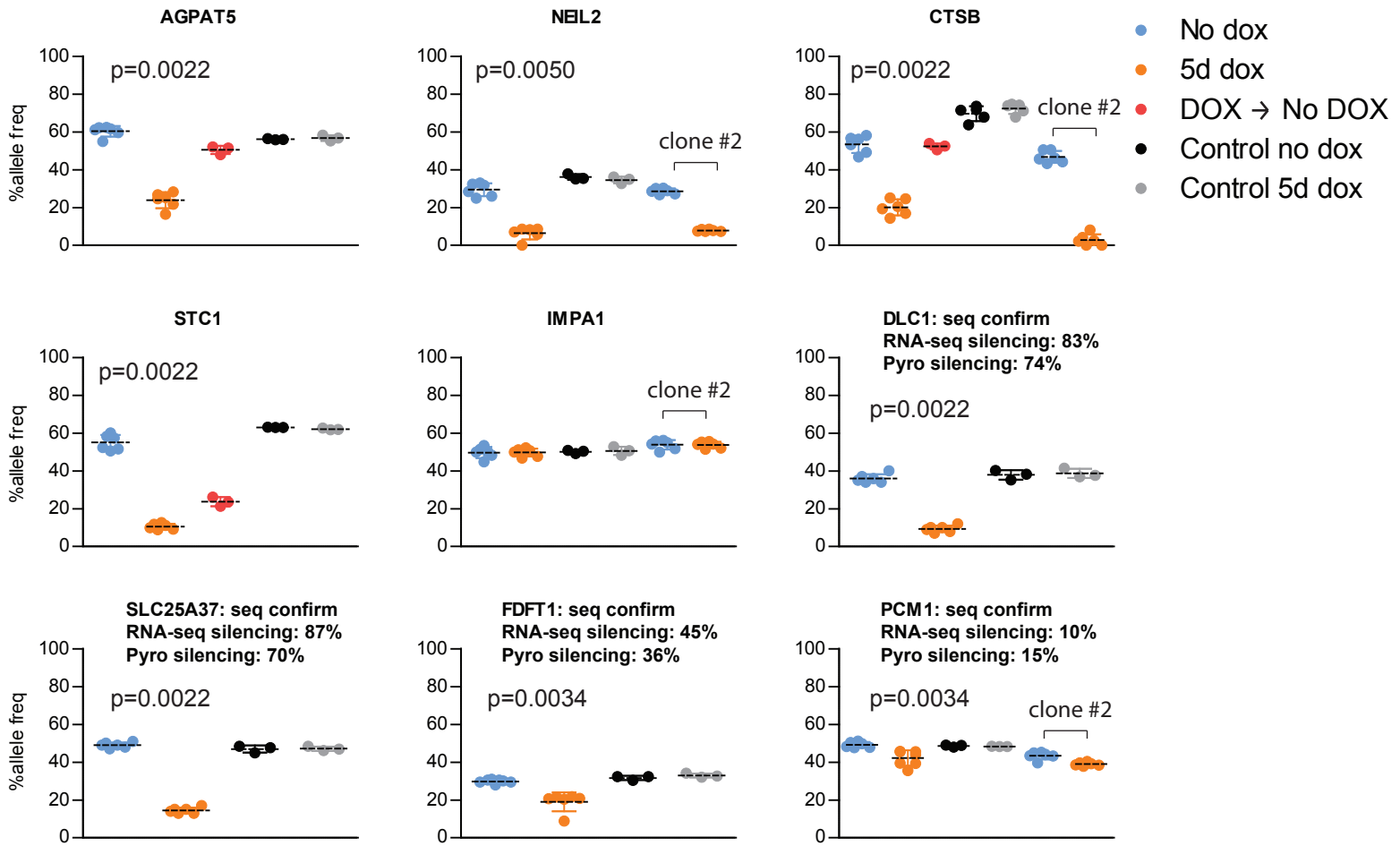
chr4



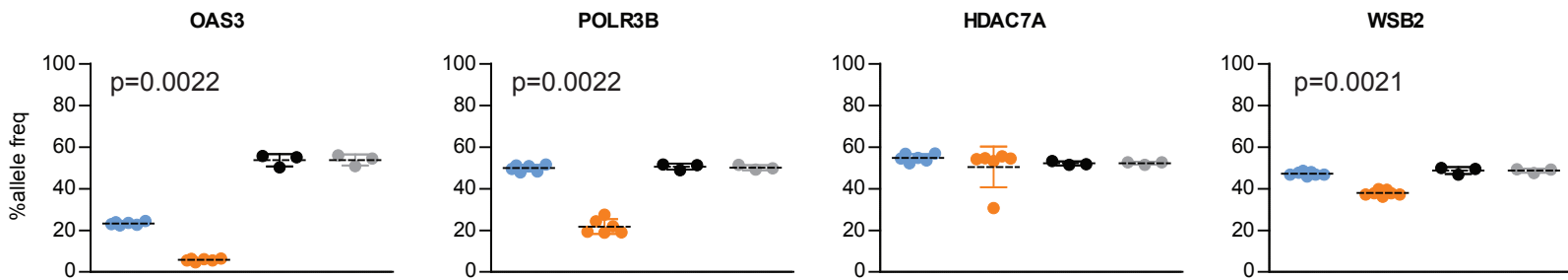
chr7 alpha = 0.0071



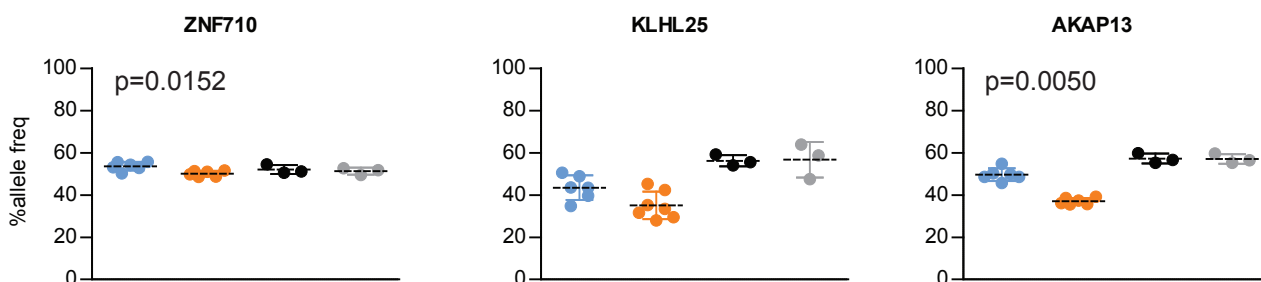
chr8 alpha = 0.0056



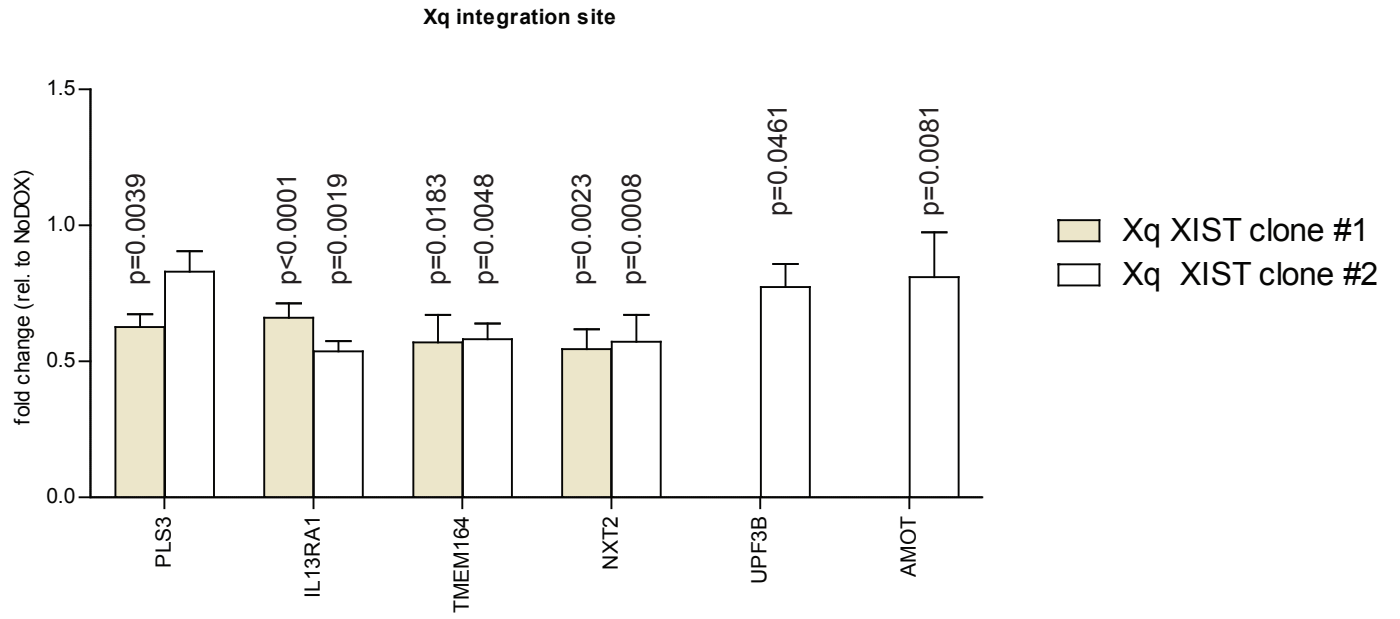
chr12 alpha = 0.0125



chr15 alpha = 0.0167

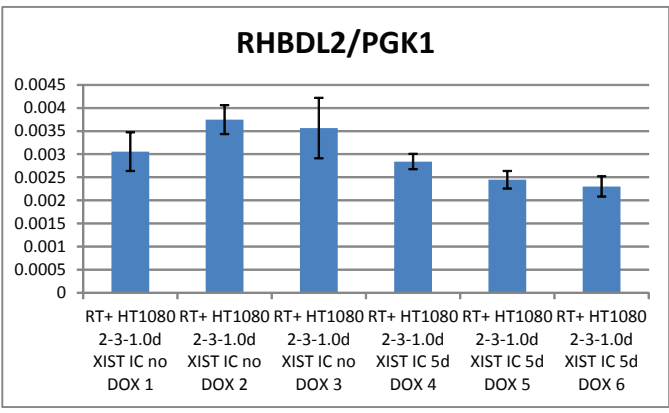
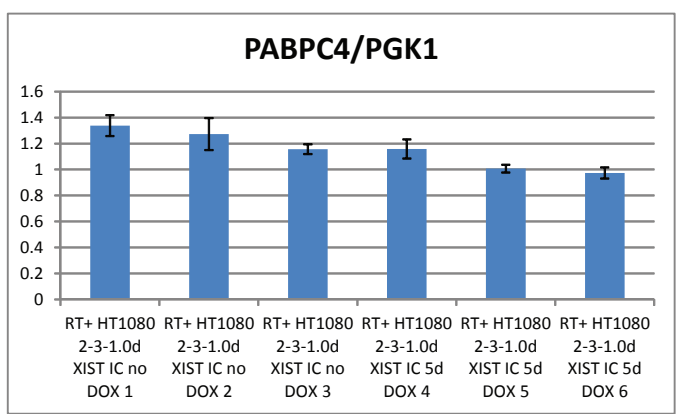
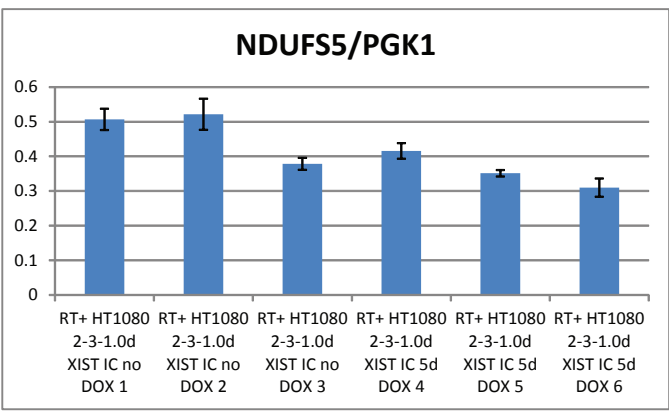


b

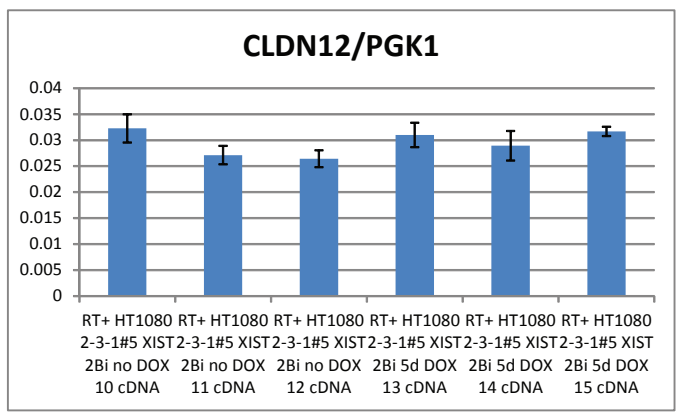
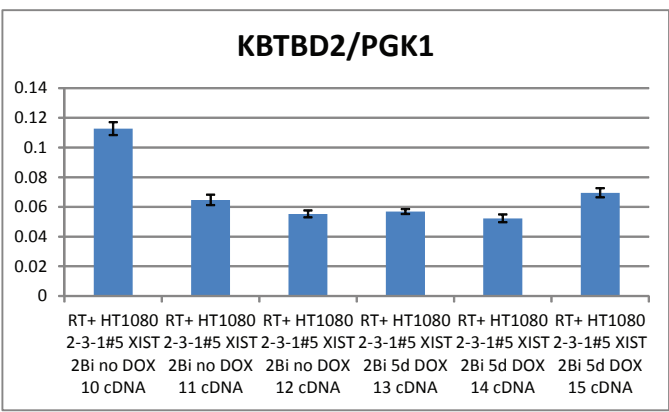


C

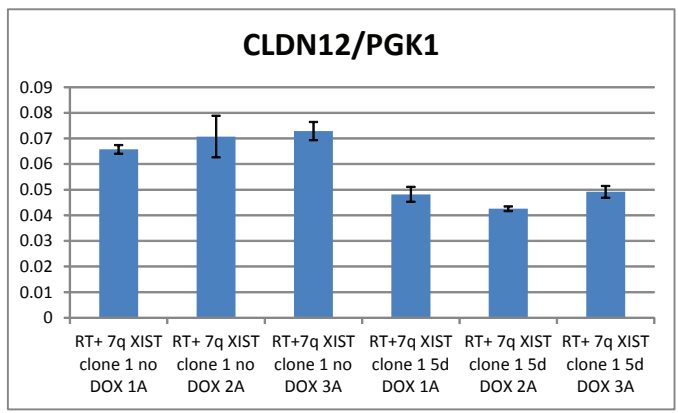
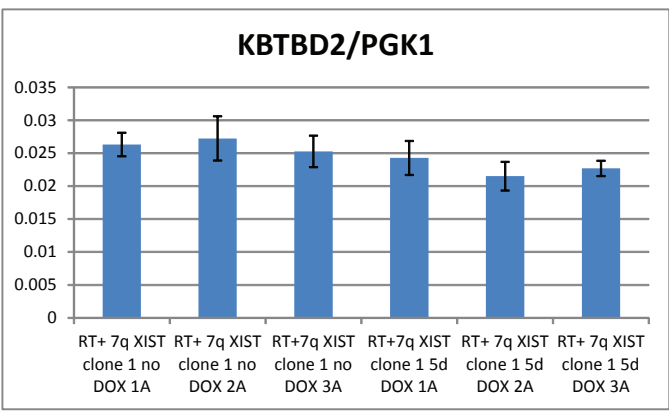
1p

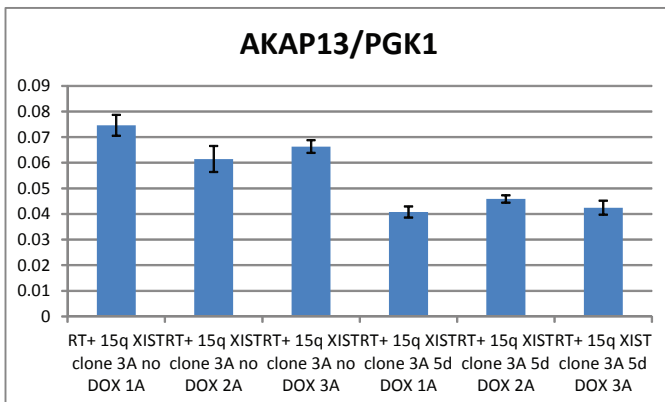
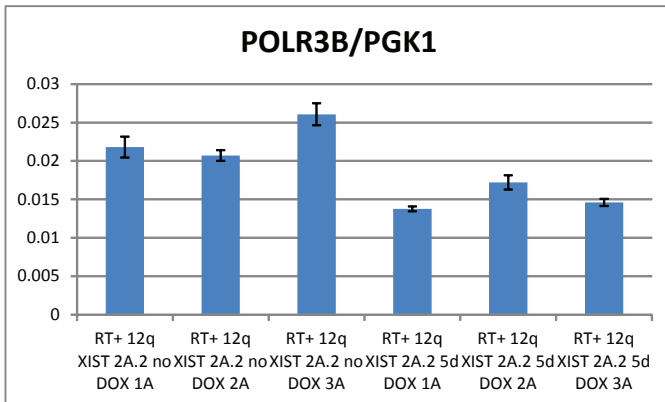
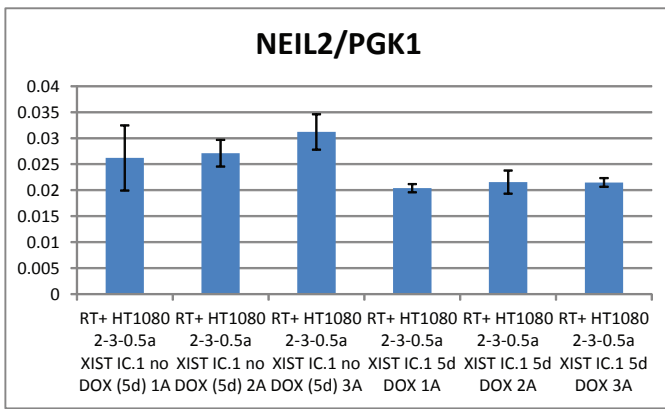


7p



7q





Additional Data File 3. Candidate gene silencing assays for each integration site.

a. Allelic pyrosequencing of three cDNAs from No DOX and DOX were compared in duplicate pyrosequencing reactions. As a control cDNA from a different integration was also assessed. One assay for each integration site was included in Figure 2.

b. For the X-chromosome integration site q-RT-PCR was used to determine silencing as the cells are hemizygous.

c. Q-RT-PCR assays were also performed to validate some autosomal silencing and compare with pyrosequencing and RNA-seq.