

## Supplemental Material

### Haplotypes of DNA repair and cell cycle control genes, x-ray exposure, and risk of childhood acute lymphoblastic leukemia

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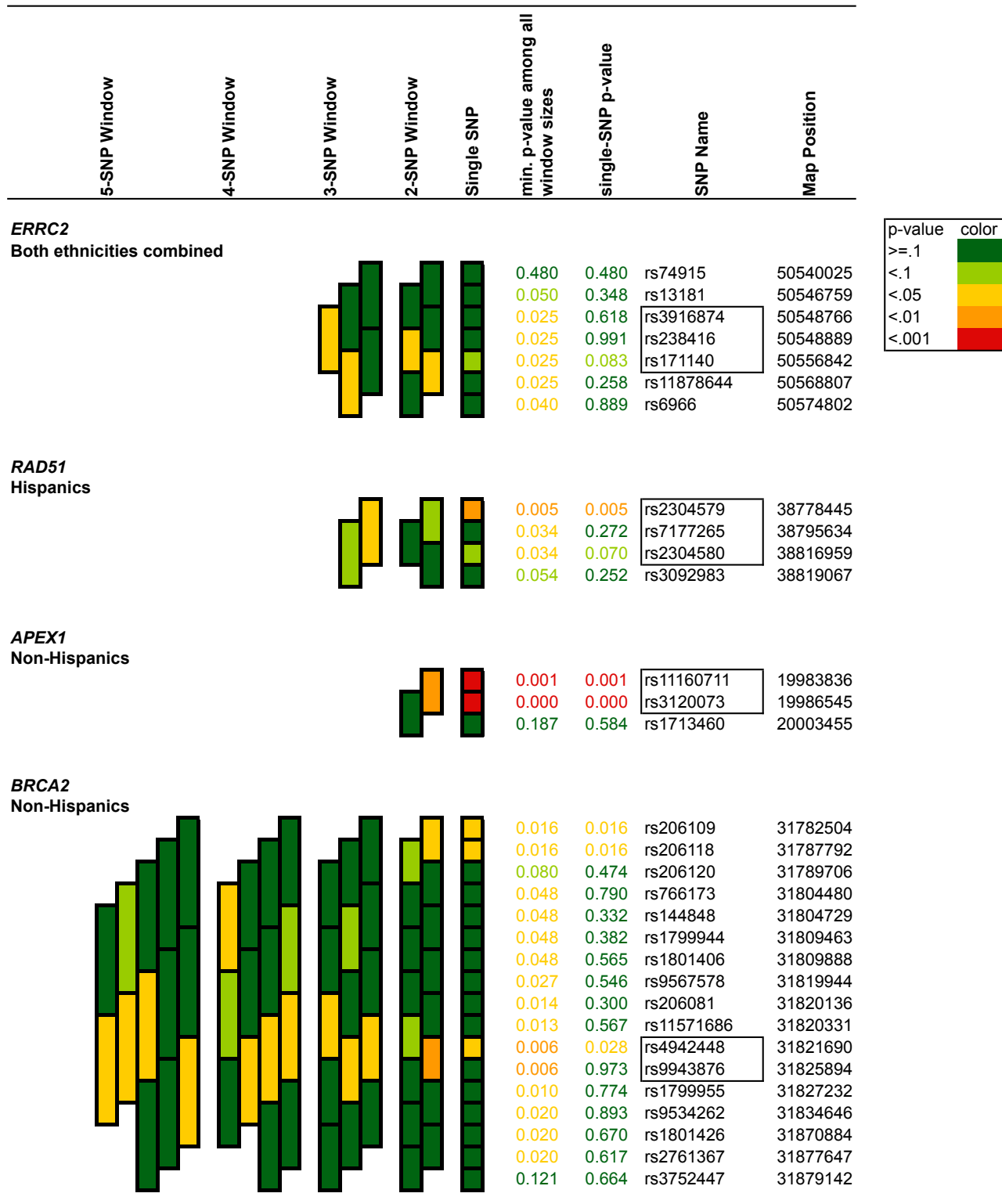
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**Supplementary Figure 1.** Significant ( $p \leq 0.05$ ) haplotype sliding window results for DNA repair and cell cycle control genes and childhood ALL. Outlined blocks show smallest multi-SNP p-values. These include a 3-SNP haplotype association for *ERCC2* ( $p=0.025$ ), a 3-SNP haplotype association for *RAD51* among Hispanics ( $p=0.034$ ), a 2-SNP haplotype association for *APEX1* among non-Hispanics ( $p=0.001$ ), and a 2-SNP haplotype association for *BRCA2* among non-Hispanics ( $p=0.006$ ). The risk estimates from haplotype trend regression of these are in Table 2.



**Supplementary Figure 2.** Significant ( $p \leq 0.05$ ) haplotype sliding window results for DNA repair and cell cycle control genes and childhood ALL, by disease subtype. Outlined blocks show windows with the smallest multi-SNP p-values. These include: for  $t(12;21)$  translocation-positive ALL, a 6-SNP haplotype association for *NBN* ( $p=0.044$ ) and a 6-SNP haplotype association for *XRCC4* ( $p=0.007$ ); for any structural changes (including  $t(12;21)$  translocations), a 3-SNP haplotype association for *XRCC4* ( $p=0.011$ ); for high hyperdiploid ALL and ALL with any numerical ploidy changes, the same 2-SNP haplotype window for *CDKN2A* ( $p=0.003$  and  $0.001$ , respectively). Risk estimates from haplotype trend regression of these are in Table 3.

