**BACKGROUND**

With 181 million base pairs encoding around 1,000 genes, chromosome 5 is about 6% of human genomic DNA. It is associated with Cockayne syndrome through the ERCC8 gene and familial adenomatous polyposis through the adenomatous polyposis coli (APC) tumor suppressor gene. Treacher Collins syndrome is also chromosome 5 associated and is caused by insertions or deletions within the TCOF1 gene. Deletion of the p arm of chromosome 5 leads to Cri du chat syndrome. Deletion of 5q or chromosome 5 altogether is common in therapy-related acute myelogenous leukemias and myelodysplastic syndrome.

**REFERENCES**


**CHROMOSOMAL LOCATION**

Genetic locus: C5orf38 (human) mapping to 5p15.33.

**SOURCE**

CEI (W-17) is an affinity purified goat polyclonal antibody raised against a peptide mapping within an internal region of CEI of human origin.