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## Summary

|                        |                                |
|------------------------|--------------------------------|
| <b>Production Name</b> | UDG Rabbit Polyclonal Antibody |
| <b>Description</b>     | Rabbit Polyclonal Antibody     |
| <b>Host</b>            | Rabbit                         |
| <b>Application</b>     | WB,                            |
| <b>Reactivity</b>      | Human,Mouse,Rat                |

## Performance

|                     |  |
|---------------------|--|
| <b>Conjugation</b>  | Unconjugated   |
| <b>Modification</b> | Unmodified   |
| <b>Isotype</b>      | IgG  |
| <b>Clonality</b>    | Polyclonal   |
| <b>Form</b>         | Liquid   |
| <b>Storage</b>      | Store at 4°C short term. Aliquot and store at -20°C long term. Avoid freeze/thaw cycles. |
| <b>Buffer</b>       | Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% New type preservative N.       |
| <b>Purification</b> | Affinity purification  |

## Immunogen

|                          |  |
|--------------------------|--|
| <b>Gene Name</b>         | UNG  |
| <b>Alternative Names</b> | UNG; DGU; UNG1; UNG15; Uracil-DNA glycosylase; UDG   |
| <b>Gene ID</b>           | 7374.0   |
| <b>SwissProt ID</b>      | P13051.The antiserum was produced against synthesized peptide derived from human UNG. AA range:191-240 |

## Application

|                         |                                     |
|-------------------------|-------------------------------------|
| <b>Dilution Ratio</b>   | WB 1:500 - 1:2000. ELISA: 1:40000.. |
| <b>Molecular Weight</b> | 35kD                                |

## Background

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**Product Name: UDG Rabbit Polyclonal Antibody**  
**Catalog #: APRab19603**

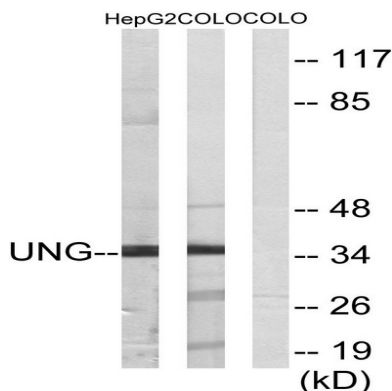


This gene encodes one of several uracil-DNA glycosylases. One important function of uracil-DNA glycosylases is to prevent mutagenesis by eliminating uracil from DNA molecules by cleaving the N-glycosylic bond and initiating the base-excision repair (BER) pathway. Uracil bases occur from cytosine deamination or misincorporation of dUMP residues. Alternative promoter usage and splicing of this gene leads to two different isoforms: the mitochondrial UNG1 and the nuclear UNG2. The UNG2 term was used as a previous symbol for the CCNO gene (GeneID 10309), which has been confused with this gene, in the literature and some databases. [provided by RefSeq, Nov 2010],disease:Defects in UNG are a cause of immunodeficiency with hyper-IgM type 5 syndrome (HIGM5) [MIM:608106]. Hyper-IgM syndrome is a condition characterized by normal or increased serum IgM concentrations associated with low or absent serum IgG, IgA, and IgE concentrations. HIGM5 is associated with profound impairment in immunoglobulin (Ig) class-switch recombination (CSR) at a DNA precleavage step.,function:Excises uracil residues from the DNA which can arise as a result of misincorporation of dUMP residues by DNA polymerase or due to deamination of cytosine.,online information:UNG mutation db,PTM:Isoform 1 is processed by cleavage of a transit peptide.,similarity:Belongs to the uracil-DNA glycosylase family.,subunit:Monomer. Interacts with HIV-1 Vpr.,tissue specificity:Isoform 1 is widely expressed with the highest expression in skeletal muscle, heart and testicles. Isoform 2 has the highest expression levels in tissues containing proliferating cells.,

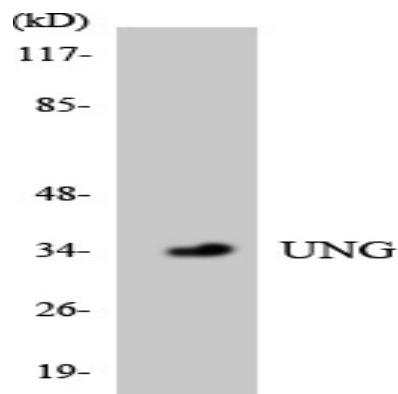
## Research Area

Base excision repair;Primary immunodeficiency;

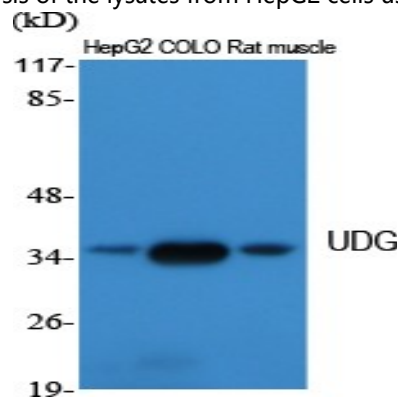
## Image Data



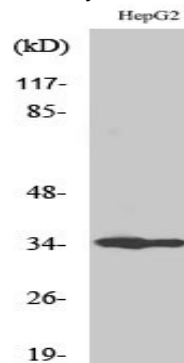
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Western blot analysis of the lysates from HepG2 cells using UNG antibody.



Western Blot analysis of various cells using UDG Polyclonal Antibody. Secondary antibody was diluted at 1:20000



Western Blot analysis of COLO205 cells using UDG Polyclonal Antibody. Secondary antibody was diluted at 1:20000

## Note

For research use only.