## Product Information

## Size:

50ul
Reactivity:
Human, Mouse, Rat

## Source:

Rabbit
Isotype:
IgG

## Applications:

ELISA, IHC
Recommended dilutions:
ELISA:1:1000-1:5000, IHC:1:50-1:200

## Protein Background:

Required for $V(D)$, recombination, the process by which exons encoding the antigenbinding domains of immunoglobulins and T-cell receptor proteins are assembled from individual $V$, ( $D$ ), and J gene segments. V(D)J recombination is initiated by the lymphoid specific RAG endonuclease complex, which generates site specific DNA double strand breaks (DSBs). These DSBs present two types of DNA end structures: hairpin sealed coding ends and phosphorylated blunt signal ends. These ends are independently repaired by the non homologous end joining (NHEJ) pathway to form coding and signal joints respectively. This protein exhibits single-strand specific $5^{\prime}-3{ }^{\prime}$ exonuclease activity in isolation and acquires endonucleolytic activity on $5^{\prime}$ and $3^{\prime}$ hairpins and overhangs when in a complex with PRKDC. The latter activity is required specifically for the resolution of closed hairpins prior to the formation of the coding joint.

## Gene ID:

CIB1
Uniprot
Q99828

## Synonyms:

calcium and integrin binding 1 (calmyrin)

## Immunogen:

Synthetic peptide of human CIB1.

## Storage:

-20\° C, pH7.4 PBS, 0.05\% NaN3, 40\% Glycerol


The image on the left is immunohistochemistry of paraffin-embedded Human thyroid cancer tissue using PACO19474(CIB1 Antibody) at dilution $1 / 50$, on the right is treated with synthetic peptide. (Original magnification: $x-200$ ).

The image on the left is immunohistochemistry of paraffin-embedded Human lung cancer tissue using PACO19474(CIB1 Antibody) at dilution $1 / 50$, on the right is treated with synthetic peptide. (Original magnification: $x$-200).

