

## CENPC | Validation File

**TARGET** Homo sapiens centromeric protein C1 (CENPC)

**CLONE NAME** AL61A

**DESCRIPTION** mouse monoclonal

**ANTIGEN USED** hCENPC Nterm-GST

**ISOTYPE** IgG1

**SPECIES REACTIVITY** human

**LOCALIZATION** nuclear

**POSITIVE CONTROL** HeLa cell line

**STORAGE BUFFER** Tissue culture supernatant: 0.02% sodium azide

**STORAGE** Aliquot and store at 4C. Do not freeze

 Recommended

 Inconclusive

 Not Recommended

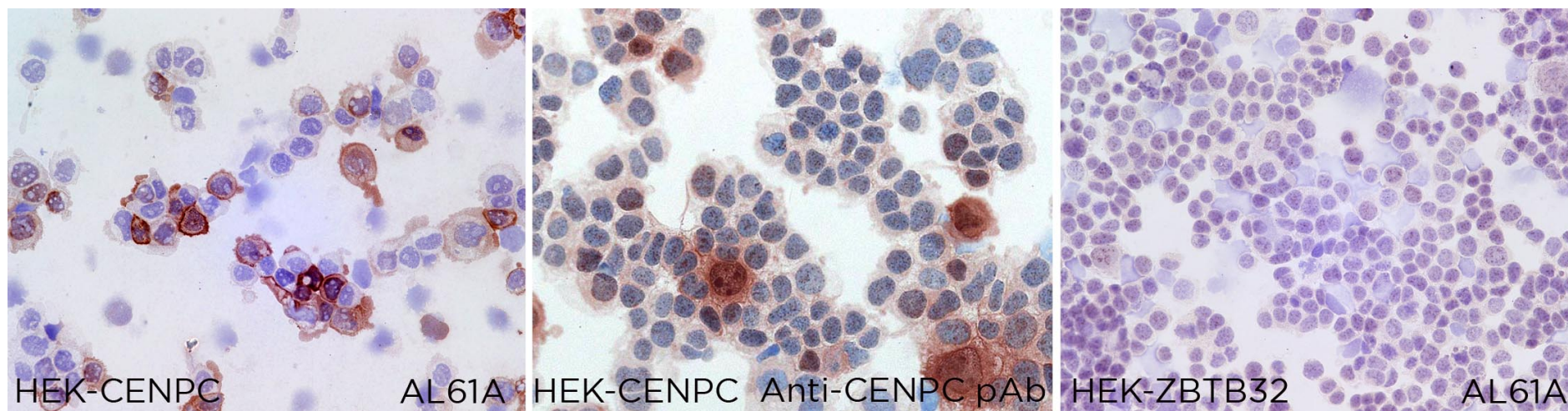
 Not Tested

# APPLICATIONS

## ● | ICC | Immunocytochemistry

AL61A mAb is able to detect human CENPC protein in immunocytochemistry.

To confirm that AL61A mAb recognizes human CENPC protein, immunocytochemistry on cytopins preparations of human CENPC expressed in HEK293T was performed. Labeling with the anti-CENPC pAb confirmed the efficiency of transfection. Cytopsin preparation of human ZBTB32 protein was used as a negative control.



● | IP | **Immunoprecipitation**

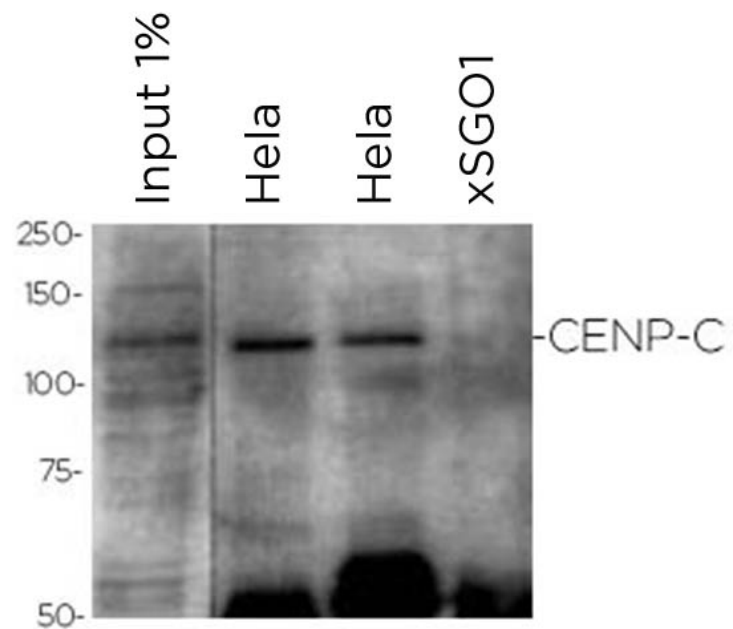
AL61A mAb is able to detect human CENP-C1 protein by WB.

**DILUTION** 1:10 (supernatant)

Predicted molecular weight: **106kDa**  
Observed molecular weight: **106kDa**

**LANES**

Lane 1 input 1% (+)  
Lane 2 HeLa nuclear extract (+)  
Lane 3 HeLa nuclear extract (+)  
Lane 4 xSGO1 protein (-)

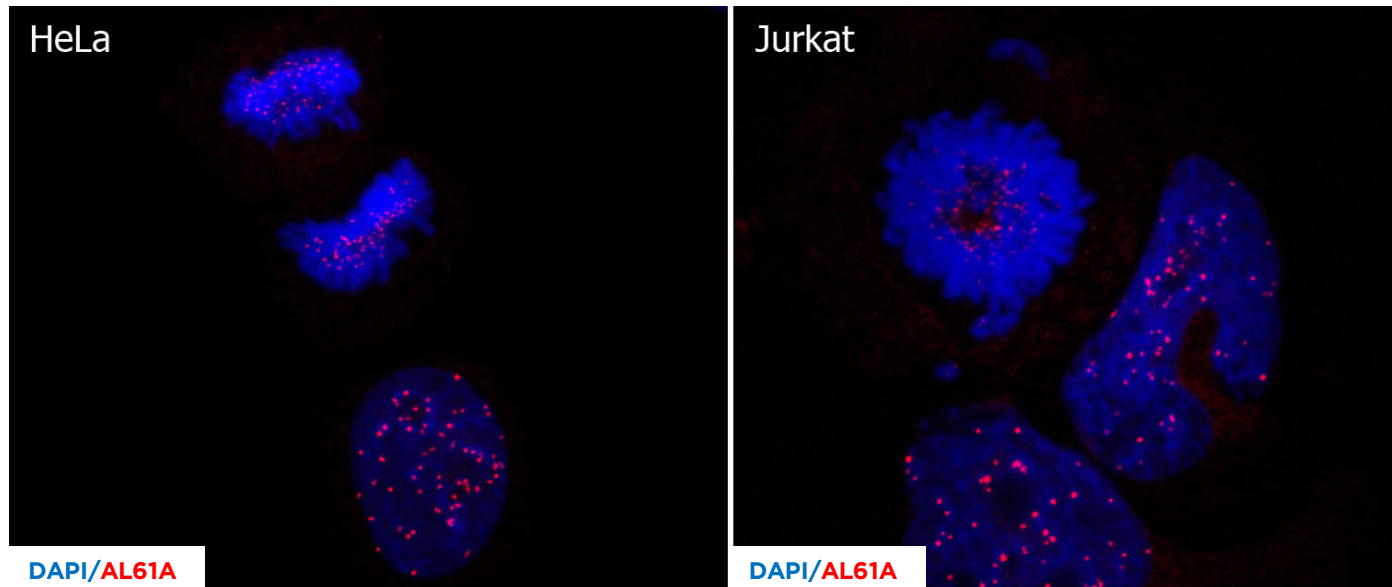


● | IF | **Immunofluorescence (paraffin)**

Antibody AL61A can be used to detect CENP-C1 in immunofluorescence.

**TISSUE SAMPLE** HeLa and Jurkat cell lines

**DILUTION** No Dilution (Neat supernatant)



● | WB | **Western Blotting** Not done

● | IHC-F | **Immunohistochemistry (frozen)** Not done

● | IHC-P | **Immunohistochemistry (paraffin)** Not done

● | FC | **Flow Cytometry** Not done

**SOLD BY:** Millipore