

Product Data Sheet

Anti-Nucleophosmin Antibody

Catalog # Source Reactivity Applications

CPA9781 Mouse H WB, IH

Description Mouse monoclonal antibody to Nucleophosmin

Immunogen KLH-conjugated synthetic peptide encompassing a sequence within human

Nucleophosmin. The exact sequence is proprietary.

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of Nucleophosmin protein.

Clonality Monoclonal

Conjugation

Form Mouse IgG1. Liquid in PBS containing 50% glycerol, 0.2% BSA and 0.01% sodium

azide.

Dilution WB (1/500 - 1/1000), IH (1/100 - 1/300)

Gene Symbol NPM1

Alternative Names NPM; Nucleophosmin; NPM; Nucleolar phosphoprotein B23; Nucleolar protein

NO38; Numatrin

Entrez Gene 4869 (Human)

SwissProt P06748 (Human)

Storage/Stability Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid

freeze/thaw cycles.

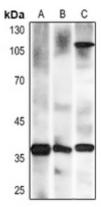
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, RNAi- RNA interference Species reactivity key: H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb-Rabbit, S- Sheep, Z- Zebrafish

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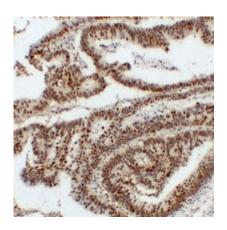
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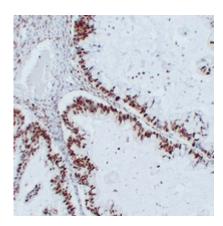
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Western blot analysis of Nucleophosmin expression in HepG2 (A), Hela (B), A431 (C) whole cell lysates. (Predicted band size: 32 kD; Observed band size: 37 kD)



Immunohistochemical analysis of Nucleophosmin staining in human colon carcinoma formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.



Immunohistochemical analysis of Nucleophosmin staining in human ovarian mucinous formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

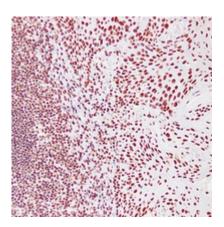
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Immunohistochemical analysis of Nucleophosmin staining in human tonsil formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

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