## abcam

## Product datasheet

## Anti－Nucleophosmin（phospho S125）antibody［EPR1856］ ab109546

重组 RabMAb



| 产品名称 | Anti－Nucleophosmin（phospho S125）抗体［EPR1856］ |
| :---: | :---: |
| 描述 | 兔单克隆抗体［EPR1856］to Nucleophosmin（phospho S125） |
| 宿主 | Rabbit |
| 经测试应用 | 适用于：Flow Cyt（Intra），WB，IHC－P |
| 种属反应性 | 与反应：Human |
|  | 预测可用于：Mouse，Rat－ |
| 免疫原 | Synthetic peptide．This information is proprietary to Abcam and／or its suppliers． |
| 阳性对照 | WB：HeLa cell lysate IHC－P：Human breast or kidney tissue |
| 常规说明 | This product is a recombinant monoclonal antibody，which offers several advantages including： |
|  | －High batch－to－batch consistency and reproducibility |
|  | －Improved sensitivity and specificity |
|  | －Long－term security of supply |
|  | －Animal－free production |
|  | For more information see here． |
|  | Our RabMAb ${ }^{\circledR}$ technology is a patented hybridoma－based technology for making rabbit monoclonal antibodies．For details on our patents，please refer to $\underline{\text { RabMAb }}{ }^{\circledR}$ patents． |


| 性能 |  |
| :--- | :--- |
| 形式 | Liquid |
| 存放说明 | Shipped at $4^{\circ} \mathrm{C}$ ．Store at $-20^{\circ} \mathrm{C}$ ．Stable for 12 months at $-20^{\circ} \mathrm{C}$. |
| 存储溶液 | $\mathrm{pH}: 7.20$ |
|  | Preservative： $0.05 \%$ Sodium azide |
|  | Constituents： $0.1 \%$ BSA， $40 \%$ Glycerol（glycerin，glycerine）， $9.85 \%$ Tris glycine， $50 \%$ Tissue |
|  | culture supernatant |
| 纯度 | Protein A purified |
| 克隆 | 单克隆 |
| 克隆编号 | EPR1856 |

## 应用

## The Abpromise guarantee Abpromise ${ }^{\text {TM 承诺保证使用ab109546于以下的经测试应用 }}$

＂应用说明＂部分下显示的仅为推荐的起始稀释度；实际最佳的稀释度 $/$ 浓度应由使用者检定。

| 应用 | Ab评论 | 说明 |
| :--- | :--- | :--- |
| Flow Cyt（Intra） |  | $1 / 100$. <br> ab172730－Rabbit monoclonal lgG，is suitable for use as an <br> isotype control with this antibody． |
| WB |  | （1） |
| IHC－P |  | $1 / 10000-1 / 50000$. Predicted molecular weight： 33 kDa. |

## 靶标

功能

## 疾病相关

## 序列相似性

翻译后修饰
Involved in diverse cellular processes such as ribosome biogenesis，centrosome duplication， protein chaperoning，histone assembly，cell proliferation，and regulation of tumor suppressors p53／TP53 and ARF．Binds ribosome presumably to drive ribosome nuclear export．Associated with nucleolar ribonucleoprotein structures and bind single－stranded nucleic acids．Acts as a chaperonin for the core histones $\mathrm{H} 3, \mathrm{H} 2 \mathrm{~B}$ and H 4 ．Stimulates APEX1 endonuclease activity on apurinic／apyrimidinic（AP）double－stranded DNA but inhibits APEX1 endonuclease activity on AP single－stranded RNA．May exert a control of APEX1 endonuclease activity within nucleoli devoted to repair AP on rDNA and the removal of oxidized rRNA molecules．In concert with BRCA2， regulates centrosome duplication．Regulates centriole duplication：phosphorylation by PLK2 is able to trigger centriole replication．Negatively regulates the activation of EIF2AK2／PKR and suppresses apoptosis through inhibition of EIF2AK2／PKR autophosphorylation．Antagonizes the inhibitory effect of ATF5 on cell proliferation and relieves ATF5－induced G2／M blockade （PubMed：22528486）．

| 疾病相关 | A chromosomal aberration involving NPM1 is found in a form of non－Hodgkin lymphoma． |
| :--- | :--- |
|  | Translocation $t(2 ; 5)(p 23 ; q 35)$ with ALK．The resulting chimeric NPM1－ALK protein homodimerize |
| and the kinase becomes constitutively activated． |  |
|  | A chromosomal aberration involving NPM1 is found in a form of acute promyelocytic leukemia． |
|  | Translocation $t(5 ; 17)(q 32 ; q 11)$ with RARA． |
|  | A chromosomal aberration involving NPM1 is a cause of myelodysplastic syndrome（MDS）． |
|  | Translocation $t(3 ; 5)(q 25.1 ; q 34)$ with MLF1． |
|  | Defects in NPM1 are associated with acute myelogenous leukemia（AML）．Mutations in exon 12 |
| affecting the C－terminus of the protein are associated with an aberrant cytoplasmic location． |  |
| 序列相似性 | Belongs to the nucleoplasmin family． |
| 翻译后修饰 | Acetylated at C－terminal lysine residues，thereby increasing affinity to histones． |
|  | ADP－ribosylated． |

required for centriole duplication and is sufficient to trigger centriole replication．Phosphorylation at Ser－4 by PLK1 takes place during mitosis．Phosphorylated by CDK2 at Ser－125 and Thr－199． Phosphorylation at Thr－199 may trigger initiation of centrosome duplication．Phosphorylated by CDK1 at Thr－199，Thr－219，Thr－234 and Thr－237 during cell mitosis．When these four sites are phosphorated，RNA－binding activity seem to be abolished．May be phosphorylated at Ser－70 by NEK2．The Thr－199 phosphorylated form has higher affinity for ROCK2．CDK6 triggers Thr－199 phosphorylation when complexed to Kaposi＇s sarcoma herpesvirus（KSHV）V－cyclin，leading to viral reactivation by reducing viral LANA levels．

Sumoylated by ARF．
细胞定位
Nucleus，nucleolus．Nucleus，nucleoplasm．Cytoplasm，cytoskeleton，microtubule organizing center，centrosome．Generally nucleolar，but is translocated to the nucleoplasm in case of serum starvation or treatment with anticancer drugs．Has been found in the cytoplasm in patients with primary acute myelogenous leukemia（AML），but not with secondary AML．Can shuttle between cytoplasm and nucleus．Co－localizes with the methylated form of RPS10 in the granular component（GC）region of the nucleolus．Colocalized with nucleolin and APEX1 in nucleoli． Isoform 1 of NEK2 is required for its localization to the centrosome during mitosis．

图片


Westem blot－Anti－Nucleophosmin（phospho S125） antibody［EPR1856］（ab109546）


Flow Cytometry（Intracellular）－Anti－Nucleophosmin （phospho S125）antibody［EPR1856］（ab109546）

All lanes：Anti－Nucleophosmin（phospho S125）antibody ［EPR1856］（ab109546）at 1／10000 dilution

Lane 1 ：HeLa cell lysate
Lane 2 ：HeLa cell lysate treated with Alkaline Phosphatase

Lysates／proteins at $10 \mu \mathrm{~g}$ per lane．

Predicted band size： 33 kDa

Overlay histogram showing HeLa cells stained with ab109546（red line）．The cells were fixed with $80 \%$ methanol（ 5 min ）and then permeabilized with $0.1 \%$ PBS－Tween for 20 min ．The cells were then incubated in 1x PBS／10\％normal goat serum／0．3M glycine to block non－specific protein－protein interactions followed by the antibody（ab109546，1／100 dilution）for 30 min at $22^{\circ} \mathrm{C}$ ．The secondary antibody used was Alexa Fluor ${ }^{\circledR} 488$ goat anti－rabbit lgG $(\mathrm{H}+\mathrm{L})(\mathbf{a b 1 5 0 0 7 7})$ at $1 / 2000$ dilution for 30 min at $22^{\circ} \mathrm{C}$ ．Isotype control antibody（black line）was rabbit lgG（monoclonal）$\left(1 \mu \mathrm{~g} / 1 \times 10^{6}\right.$ cells）used under the same conditions．Unlabelled sample（blue line）was also used as a control．Acquisition of $>5,000$ events were collected using a 20 mW Argon ion laser（488nm）and 525／30


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Nucleophosmin (phospho S125) antibody [EPR1856] (ab109546)


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Nucleophosmin (phospho S125) antibody [EPR1856] (ab109546)
ab109546, at a $1 / 100$ dilution, staining Nucleophosmin in formailnfixed, paraffin-embedded Human breast tissue.

Perform heat mediated antigen retrieval via the pressure cooker method before commencing with IHC staining protocol.
ab109546, at a 1/100 dilution, staining Nucleophosmin in formalinfixed, paraffin-embedded Human kidney tissue.

Perform heat mediated antigen retrieval via the pressure cooker method before commencing with IHC staining protocol.


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