Datasheet for ABIN7298276
anti-Cyclin L1 antibody (C-Term)

## 2 Images

## Overview

| Quantity: | $100 \mu \mathrm{~L}$ |
| :--- | :--- |
| Target: | Cyclin L1 (CCNL1) |
| Binding Specificity: | C-Term |
| Reactivity: | Human, Mouse, Rat, Dog, Cow, Pig |
| Host: | Robbit |
| Clonality: | This Cyclin L1 antibody is un-conjugated |
| Conjugate: | Western Blotting (WB), Immunohistochemistry (IHC) |
| Application: |  |

Product Details

| Immunogen: | KLH-conjugated synthetic peptide encompassing a sequence within the C-term region of <br> human Cyclin L1. |
| :--- | :--- |
| Specificity: | Recognizes endogenous levels of Cyclin L1 protein. |
| Characteristics: | Rabbit polyclonal antibody to Cyclin L1 |
| Purification: | The antibody was purified by immunogen affinity chromatography. |
| Target Details | Cyclin L1 (CCNL1) |
| Target: | Cyclin L1 (CCNL1 Products) |
| Cyclin-L1, Cyclin-L |  |

Target Details

| Gene ID: | 57018, 56706, 100909712 |
| :---: | :---: |
| UniProt: | Q9UK58, Q52KE7, Q9R1Q2 |
| Application Details |  |
| Application Notes: | WB (1:500-1:1000), IH (1:100-1:200) |
| Restrictions: | For Research Use only |
| Handling |  |
| Format: | Liquid |
| Buffer: | Liquid in 0.42 \% Potassium phosphate, $0.87 \%$ Sodium chloride, $\mathrm{pH} 7.3,30 \%$ glycerol, and 0.01 \% sodium azide. |
| Preservative: | Sodium azide |
| Precaution of Use: | This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only. |
| Storage: | $-20^{\circ} \mathrm{C}$ |
| Storage Comment: | Shipped at $4^{\circ} \mathrm{C}$. Upon delivery aliquot and store at $-20^{\circ} \mathrm{C}$ for one year. Avoid freeze/thaw cycles. |
| Expiry Date: | 12 months |

Images


Western Blotting
Image 1. Western blot analysis of Cyclin L1 expression in HL60 (A), NIH3T3 (B), Jurkat (C), THP1 (D), K562 (E) whole cell lysates.


## Immunohistochemistry <br> Image 2. Immunohistochemical analysis of Cyclin L1 staining in human breast cancer 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.

