

**Phospho-CCNB1(S35) Blocking Peptide**  
**Synthetic peptide**  
**Catalog # BP3882a****Specification**

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**Phospho-CCNB1(S35) Blocking Peptide - Product Information**

Primary Accession [P14635](#)  
Other Accession [NP\\_114172.1](#)

**Phospho-CCNB1(S35) Blocking Peptide - Additional Information**

**Gene ID** 891

**Other Names**

G2/mitotic-specific cyclin-B1, CCNB1, CCNB

**Target/Specificity**

The synthetic peptide sequence is selected from aa 28-42 of HUMAN CCNB1

**Format**

Peptides are lyophilized in a solid powder format. Peptides can be reconstituted in solution using the appropriate buffer as needed.

**Storage**

Maintain refrigerated at 2-8°C for up to 6 months. For long term storage store at -20°C.

**Precautions**

This product is for research use only. Not for use in diagnostic or therapeutic procedures.

**Phospho-CCNB1(S35) Blocking Peptide - Protein Information**

**Name** CCNB1

**Synonyms** CCNB

**Function**

Essential for the control of the cell cycle at the G2/M (mitosis) transition.

**Cellular Location**

Cytoplasm. Nucleus. Cytoplasm, cytoskeleton, microtubule organizing center, centrosome

**Phospho-CCNB1(S35) Blocking Peptide - Protocols**

Provided below are standard protocols that you may find useful for product applications.

- [Blocking Peptides](#)

**Phospho-CCNB1(S35) Blocking Peptide - Images****Phospho-CCNB1(S35) Blocking Peptide - Background**

The protein encoded by this gene is a regulatory protein involved in mitosis. The gene product complexes with p34(cdc2) to form the maturation-promoting factor (MPF). Two alternative transcripts have been found, a constitutively expressed transcript and a cell cycle-regulated transcript, that is expressed predominantly during G2/M phase. The different transcripts result from the use of alternate transcription initiation sites. [provided by RefSeq].

**Phospho-CCNB1(S35) Blocking Peptide - References**

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van Zon, W., et al. J. Cell Biol. 190(4):587-602(2010)  
Harley, M.E., et al. EMBO J. 29(14):2407-2420(2010)  
Olson, J.E., et al. Breast Cancer Res. Treat. (2010) In press :  
Nantajit, D., et al. PLoS ONE 5 (8), E12341 (2010) :