

#### OriGene Technologies, Inc.

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# Product datasheet for TA346907

## Cyclin H (CCNH) Mouse Monoclonal Antibody [Clone ID: 4E11-G2-D7]

### **Product data:**

Product Type:	Primary Antibodies
Clone Name:	4E11-G2-D7
Applications:	IP, WB
<b>Recommend Dilution:</b>	WB: 1:1000
Reactivity:	Human
Host:	Mouse
lsotype:	lgG2b
Clonality:	Monoclonal
Immunogen:	The immunogen for CCNH antibody: purified recombinant human Cyclin H protein fragments expressed in E.coli.
Formulation:	Purified mouse monoclonal in buffer containing 0.1M Tris-Glycine (pH 7.4, 150 mM NaCl) with 0.02% sodium azide, 50% glycerol
Purification:	Affinity purified
Predicted Protein Size:	38 kDa
Gene Name:	cyclin H
Database Link:	<u>NP_001230 Entrez Gene 902 Human</u>
Background:	The protein encoded by this gene belongs to the highly conserved cyclin family, whose members are characterized by a dramatic periodicity in protein abundance through the cell cycle. Cyclins function as regulators of CDK kinases. Different cyclins exhibit distinct expression and degradation patterns which contribute to the temporal coordination of each mitotic event. This cyclin forms a complex with CDK7 kinase and ring finger protein MAT1. The kinase complex is able to phosphorylate CDK2 and CDC2 kinases, thus functions as a CDK-activating kinase (CAK). This cyclin and its kinase partner are components of TFIIH, as well as RNA polymerase II protein complexes. They participate in two different transcriptional regulation processes, suggesting an important link between basal transcription control and the cell cycle machinery. A pseudogene of this gene is found on chromosome 4. Alternate
	splicing results in multiple transcript variants.[



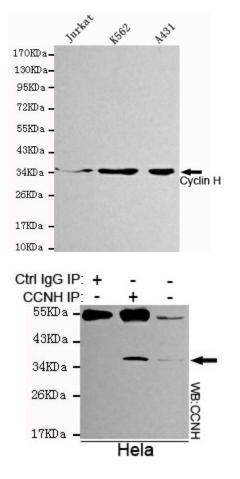
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Protein Families: Druggable Genome, Transcription Factors

Protein Pathways: Cell cycle, Nucleotide excision repair

#### **Product images:**



Western blot detection of Cyclin H in Jurkat, K562 and A431 cell lysates using Cyclin H mouse mAb (1:1000 diluted).Predicted band size: 38KDa.Observed band size: 38KDa.

Immunoprecipitation analysis of Hela cell lysates using Cyclin H mouse mAb.

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