# **PRODUCT** INFORMATION



Cdk2 Inhibitor II

Item No. 15154

CAS Registry No.:	222035-13-4	0
Formal Name:	4-[2-(5-bromo-1,2-dihydro-2-oxo-3H-indol-	°≈ <mark>s −</mark> NH2
	3-ylidene)hydrazinyl]-benzenesulfonamide	
Synonyms:	Cyclin-dependent kinase 2 Inhibitor II,	
	SC-221409	
MF:	C <sub>14</sub> H <sub>11</sub> BrN <sub>4</sub> O <sub>3</sub> S	
FW:	395.2	N —N
Purity:	≥95%	Br H
UV/Vis.:	λ <sub>max</sub> : 213, 257, 395 nm	
Supplied as:	A crystalline solid	
Storage:	-20°C	
Stability:	≥4 years	Н

Information represents the product specifications. Batch specific analytical results are provided on each certificate of analysis.

## Laboratory Procedures

Cdk2 inhibitor II is supplied as a crystalline solid. A stock solution may be made by dissolving the Cdk2 inhibitor II in the solvent of choice, which should be purged with an inert gas. Cdk2 inhibitor II is soluble in the organic solvent DMSO at a concentration of approximately 5 mg/ml.

## Description

The cyclin-dependent kinase 2 (Cdk2) works with cyclins A or E to regulate S phase and G<sub>2</sub>-M transition during the cell cycle.<sup>1</sup> Cdk2 inhibitor II is a 3-(benzylidne)indolin-2-one analog that selectively and potently inhibits Cdk2 (IC<sub>50</sub> = 60 nM).<sup>2</sup> It is cell permeable, reversible, and ATP-competitive.<sup>2</sup>

## References

- 1. Funk, J.O. Cell cycle checkpoint genes and cancer. Encyclopedia of life sciences. John Wiley & Sons (2005).
- 2. Davis, S.T., Benson, B.G., Bramson, H.N., et al. Prevention of chemotherapy-induced alopecia in rats by CDK inhibitors. Science 291(5501), 134-137 (2001).

WARNING THIS PRODUCT IS FOR RESEARCH ONLY - NOT FOR HUMAN OR VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

### SAFETY DATA

This material should be considered hazardous until further information becomes available. Do not ingest, inhale, get in eyes, on skin, or on clothing. Wash thoroughly after handling. Before use, the user must review the complete Safety Data Sheet, which has been sent via email to your institution.

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1180 EAST ELLSWORTH RD ANN ARBOR, MI 48108 · USA PHONE: [800] 364-9897 [734] 971-3335 FAX: [734] 971-3640 CUSTSERV@CAYMANCHEM.COM WWW.CAYMANCHEM.COM