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## DATASHEET

CMIQ

### Product overview

<b>Name</b>	CMIQ
<b>Cat No</b>	HB0077
<b>Biological action</b>	Inhibitor
<b>Purity</b>	>95%
<b>Description</b>	Potent, selective PKA inhibitor

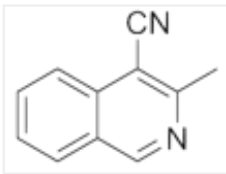
### Biological Data

<b>Biological description</b>	Potent and selective protein kinase A (PKA) inhibitor ( $IC_{50} = 30 \text{ nM}$ ). Membrane permeable. Blocks ATP-binding site of PKA. Restores morphine-related behaviours in morphine tolerance.
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### Solubility & Handling

<b>Storage instructions</b>	+4 °C
<b>Solubility overview</b>	Soluble in DMSO (50mg/ml)
<b>Important</b>	This product is for RESEARCH USE ONLY and is not intended for therapeutic or diagnostic use. Not for human or veterinary use.

### Chemical Data

<b>Chemical name</b>	PKA Inhibitor XI
<b>Molecular Weight</b>	168.2
<b>Chemical structure</b>	
<b>Molecular Formula</b>	$C_{11}H_8N_2$
<b>CAS Number</b>	161468-32-2

### References

#### PKC and PKA inhibitors reinstate morphine-induced behaviors in morphine tolerant mice.

Smith FL *et al* (2006) Pharmacol Res 54(6)

**PubMedID** [17056270](#)

#### Selective inhibition of cyclic AMP-dependent protein kinase by isoquinoline derivatives.

Lu ZX *et al* (1996) Biol Chem Hoppe Seyler 377(6)

**PubMedID** [8839983](#)

Epac mediates a cAMP-to-PKC signaling in inflammatory pain: an isolectin B4(+) neuron-specific mechanism.

