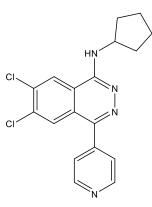


## Catalog # 10-4018

A-196

CAS# 1982372-88-2 Cyclopentyl-(6,7-dichloro-4-pyridin-4-yl-phthalazin-1-yl)-amine Lot # FBS1100



A-196 is a potent and selective (over 29 other methyltransferases) inhibitor of the protein lysine methyltransferases SUV420H1( $IC_{50} = 25 \text{ nM}$ ) and SUV420H2 ( $IC_{50} = 144 \text{ nM}$ ). It substantially inhibited non-homologous end-joining (NHEJ)-directed DNA repair but not homology-directed repair(HDR) in cells treated with ionizing radiation. A-196 is active in cells and displayed no toxicity in several cell lines.

1) Bromberg et al. (2017), The SUV4-20 inhibitor A-196 verifies a role for epigenetics in genomic integrity; Nat.Chem.Biol. 13 317

## PHYSICAL DATA

Molecular Weight:	359.25
Molecular Formula:	C18H16Cl2N4
Purity:	>98%
	NMR: (Conforms)
Solubility:	Soluble in DMSO (5 mg/ml)
Physical Description:	White solid
Storage and Stability:	Store as supplied at -20° for up to 1 year from the date of purchase. Store solutions
	at -20°C for up to 1 month.

Materials provided by Focus Biomolecules are for laboratory research use only and are not intended for human or veterinary applications.

Focus Biomolecules LLC 400 Davis Drive, Suite 600 Plymouth Meeting PA 19462 www.focusbiomolecules.com