



# SZABO SCANDIC

Part of Europa Biosite

## Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

Weitere Information auf den folgenden Seiten!  
See the following pages for more information!



### Lieferung & Zahlungsart

siehe unsere [Liefer- und Versandbedingungen](#)

### Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

### SZABO-SCANDIC Handels GmbH

Quellenstraße 110, A-1100 Wien

T. +43(0)1 489 3961-0

F. +43(0)1 489 3961-7

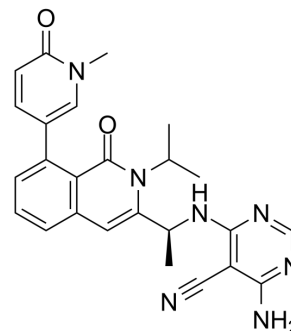
[mail@szabo-scandic.com](mailto:mail@szabo-scandic.com)

[www.szabo-scandic.com](http://www.szabo-scandic.com)

[linkedin.com/company/szaboscandic](https://www.linkedin.com/company/szaboscandic) 

## IPI-3063

Cat. No.:	HY-111510		
CAS No.:	1425043-73-7		
Molecular Formula:	C <sub>25</sub> H <sub>25</sub> N <sub>7</sub> O <sub>2</sub>		
Molecular Weight:	455.51		
Target:	PI3K		
Pathway:	PI3K/Akt/mTOR		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	6 months
		-20°C	1 month



## SOLVENT & SOLUBILITY

### In Vitro

DMSO : ≥ 62.5 mg/mL (137.21 mM)

\* "≥" means soluble, but saturation unknown.

	Solvent Concentration	Mass	1 mg	5 mg	10 mg
Preparing Stock Solutions	1 mM		2.1953 mL	10.9767 mL	21.9534 mL
	5 mM		0.4391 mL	2.1953 mL	4.3907 mL
	10 mM		0.2195 mL	1.0977 mL	2.1953 mL

Please refer to the solubility information to select the appropriate solvent.

## BIOLOGICAL ACTIVITY

Description	IPI-3063 is a potent and selective PI3K p110δ inhibitor with an IC <sub>50</sub> of 2.5 ± 1.2 nM.			
IC <sub>50</sub> & Target	p110δ 2.5 nM (IC <sub>50</sub> )	p110α 1170 nM (IC <sub>50</sub> )	p110β 1508 nM (IC <sub>50</sub> )	p110γ 2187 nM (IC <sub>50</sub> )
In Vitro	<p>IPI-3063 inhibits p110α, p110β, and p110γ with IC<sub>50</sub>s of 1171±533 nM, 1508±624 nM, and 2187±1529 nM, respectively. IPI-3063 potently reduces mouse B cell proliferation, survival, and plasmablast differentiation while increasing antibody class switching to IgG1. IPI-3063 is a p110δ selective compound with an IC<sub>50</sub>=0.1 nM in p110δ-specific cell-based assays and cellular IC<sub>50</sub> values for the other class I PI3K isoforms are at least 1,000-fold higher (IC<sub>50</sub>=1901±1318 nM for p110α, IC<sub>50</sub>=102.8±35.7 nM for p110β, IC<sub>50</sub>=418.8±117.2 nM for p110γ). IPI-3063 is very potent in reducing p-AKT (significant effect at 1 nM). IPI-3063 also reduces p-ERK1/2 with a significant effect at 10 nM. IPI-3063 is very potent, achieving a significant decrease in B cell survival when present at 10 nM<sup>[1]</sup>.</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p>			

## PROTOCOL

### Kinase Assay <sup>[1]</sup>

Human recombinant PI3K- $\alpha$ , PI3K- $\beta$ , PI3K- $\delta$ , and PI3K- $\gamma$  are used. Phosphatidylinositol 4,5 bis phosphate (diC8-PtdIns(4,5)P<sub>2</sub>) is used. PI3K- $\alpha$ ,  $\beta$ , and  $\delta$  are heterodimers consisting of full length p110 $\alpha$ , p110 $\beta$ , or p110 $\delta$  catalytic subunit and the p85 $\alpha$  regulatory subunit. PI3K- $\gamma$  is a monomer of the p110 $\gamma$  catalytic subunit. Samples of kinase (10 nM- $\alpha$ ,  $\beta$ , and  $\delta$ ; 20 nM- $\gamma$ ) are incubated with IPI-3063 for 30 min at room temperature in reaction buffer (15 mM HEPES pH 7.4, 20 mM NaCl, 1 mM EGTA, 0.02% Tween 20, 10 mM MgCl<sub>2</sub>, 0.2 mg/mL bovine- $\gamma$ -globulins) followed by addition of ATP/diC8-PtdIns(4,5)P<sub>2</sub> mixture to give final concentrations of 3 mM ATP and 500  $\mu$ M diC8-PtdIns(4,5)P<sub>2</sub>. Reactions are incubated at room temperature for 2 h, with PI3K activity is assessed. Plates are read on plate reader in luminescence mode<sup>[1]</sup>. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

### Cell Assay <sup>[1]</sup>

Peripheral blood mononuclear cells (PBMCs) are first purified from blood by density gradient centrifugation. Human B cells are then purified from PBMCs by negative selection. B-cell purity is increased from 4% to >70% as measured by FACS analysis using anti-CD19 PE conjugated antibody. Purified B cells are seeded at a final concentration of 0.1 $\times$ 10<sup>6</sup> cells/mL and cultured with 2  $\mu$ g/mL human CD40L+5  $\mu$ g/mL anti-human IgM/IgG+100  $\mu$ g/mL hIL-2+100  $\mu$ g/mL hIL-21. All B cells are cultured in RPMI 1640 supplemented with 10% (vol/vol) heat-inactivated FCS, 5 mM Hepes, 2 mM L-glutamine, 100 U/mL Penicillin, 100  $\mu$ g/mL Streptomycin, 50  $\mu$ M 2-mercaptoethanol. Purified human B cells are pretreated with IPI-3063 (0.1, 1, 10, and 100 nM) for 30 min, then stimulated with human CD40L+anti-human IgM/IgG+human IL-2+human IL-21 for 120 h<sup>[1]</sup>. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

## REFERENCES

[1]. Chiu H, et al. The Selective Phosphoinoside-3-Kinase p110 $\delta$  Inhibitor IPI-3063 Potently Suppresses B Cell Survival, Proliferation, and Differentiation. Front Immunol. 2017 Jun 30;8:747.

**Caution: Product has not been fully validated for medical applications. For research use only.**

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: tech@MedChemExpress.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA