

FGF Signaling Pathway Inhibitor Kit

A unique collection of 13 inhibitors for the study of FGF signaling pathway

Catalog No. K1040

URL for this product: www.selleckchem.com/kits/fgfr-inhibitor-kit.html

Catalog No.	Size	Price
K1040	1 ea	USD 930

Product Overview

A unique collection of 13 potent and selective inhibitors for the study of FGF signaling pathway. This kit contains the following inhibitors:

- 5 mg of MK-2206 dihydrochloride [S1078](#);
- 5 mg of BIBF1120 (Vargatef) [S1010](#);
- 5 mg of BGJ398 (NVP-BGJ398) [S2183](#);
- 10 mg of SP600125 [S1460](#);
- 50 mg of BEZ235 (NVP-BEZ235) [S1009](#);
- 25 mg of U0126-EtOH (UO126 EtOH) [S1102](#);
- 25 mg of SB 203580 [S1076](#);
- 5 mg of Ruxolitinib (INCB018424) [S1378](#);
- 5 mg of NSC 74859 (S3I-201) [S1155](#);
- 5 mg of Sorafenib (Nexavar) [S1040](#);
- 10 mg of PLX-4720 [S1152](#);
- 5 mg of KX2-391 [S2700](#);
- 5 mg of Sotrastaurin (AEB071) [S2791](#).

Product Details

Formulation:	A unique collection of 13 inhibitors for the study of FGF signaling pathway supplied as lyophilized powder		
Container:	Screw Cap Micro Tubes of WATSON		
Stability:	Powder	-20°C	2 years
	In DMSO	-80°C	6 months
Shipping:	Blue ice		
Packaging:	Inert gas		

Data

Cat. No.	Product Name	Targets and IC ₅₀ s	Suggested Solvent	Amount Supplied
S1078	MK-2206 dihydrochloride	A highly selective inhibitor of Akt . Akt1 , IC ₅₀ =8 nM; Akt2 , IC ₅₀ =12 nM; Akt3 , IC ₅₀ =65 nM.	DMSO 96 mg/mL Water 96 mg/mL	5 mg
S1010	BIBF1120 (Vargatef)	A potent VEGFR , PDGFR and FGFR kinase inhibitor for VEGFR1 , VEGFR2 , VEGFR3 with IC ₅₀ s of 34 nM, 5 nM and 5 nM, respectively.	DMSO 6 mg/mL	5 mg
S2183	BGJ398 (NVP-BGJ398)	A potent and selective FGFR family inhibitor for FGFR1 , FGFR2 , FGFR3 and FGFR4 with IC ₅₀ s of 0.9 nM, 1.4 nM, 1 nM and 60 nM, respectively.	DMSO 2 mg/mL	5 mg

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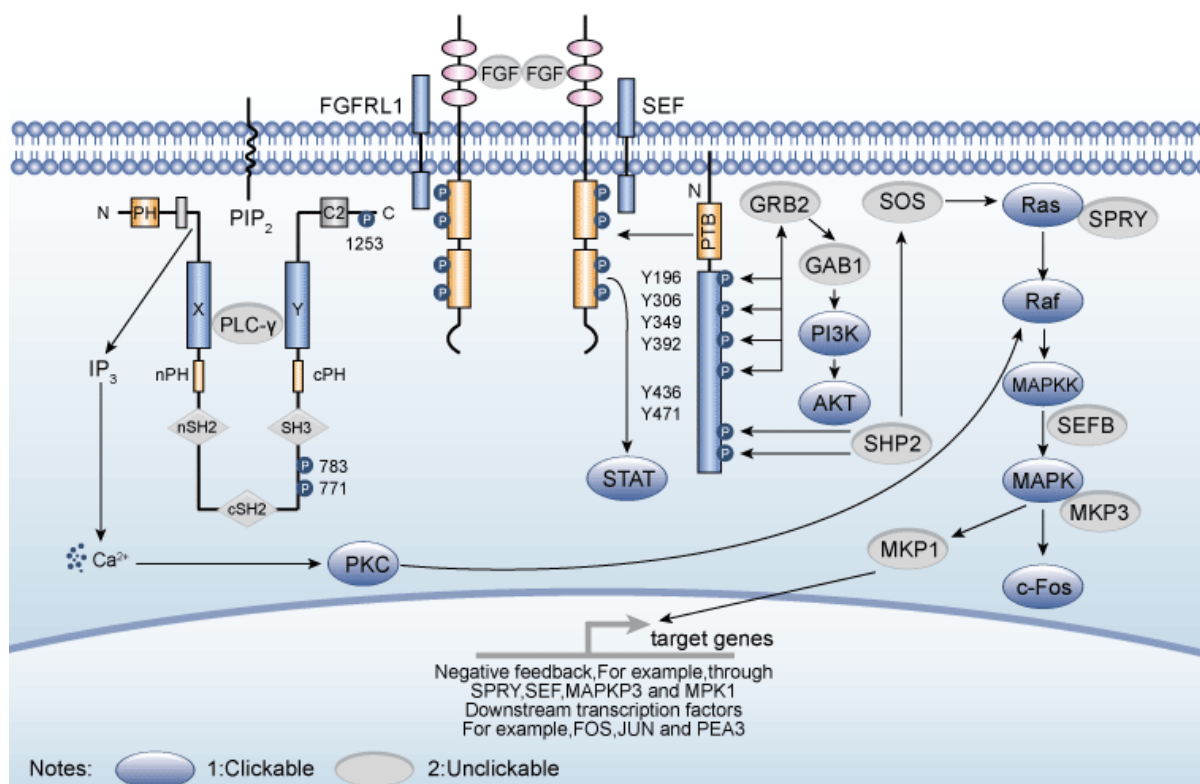
Tel: +1-832-582-8158

Fax: +1-832-582-8590

Email: sales@selleckchem.com

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S1460	SP600125	A broad-spectrum serine/threonine kinase inhibitor of JNK with an IC ₅₀ range from 40 to 90 nM.	DMSO	44 mg/mL	10 mg
S1009	BEZ235 (NVP-BEZ235)	A dual ATP-competitive inhibitor of PI3K and mTOR . p110α, IC ₅₀ =4 nM; p110β, IC ₅₀ =75 nM; p110δ, IC ₅₀ =7 nM; p110γ, IC ₅₀ =5 nM.	DMSO	7 mg/mL	50 mg
S1102	U0126-EtOH (UO126 EtOH)	A chemically synthesized and highly selective inhibitor of both MEK1 and MEK2 with IC ₅₀ s of 70 nM and 60 nM, respectively.	DMSO	85 mg/mL	25 mg
S1076	SB 203580	A p38 MAPK inhibitor with an IC ₅₀ range of 0.3-0.5 μM and blocks PKB phosphorylation with an IC ₅₀ range of 3-5 μM.	DMSO	76 mg/mL	25 mg
S1378	Ruxolitinib (INCB018424)	A JAK family kinase inhibitor of JAK1 , JAK2 and JAK3 with IC ₅₀ s of 2.7 nM, 4.5 nM and 322 nM, respectively	DMSO	61 mg/mL	5 mg
S1155	NSC 74859 (S3I-201)	A chemical probe inhibitor of STAT3 with an IC ₅₀ of 86 μM.	DMSO	73 mg/mL	5 mg
S1040	Sorafenib (Nexavar)	A small molecular inhibitor of VEGFR , PDGFR , c-Raf and B-Raf with IC ₅₀ s of 18 nM, 10 nM, 3 nM and 15 nM, respectively.	DMSO	127 mg/mL	5 mg
S1152	PLX-4720	A potent and selective inhibitor of B-Raf ^{V600E} and c-Raf-1 ^{Y340D/Y341D} with IC ₅₀ s of 13 nM and 6.7 nM, respectively.	DMSO	83 mg/mL	10 mg
S2700	KX2-391	A synthetic, orally bioavailable small molecule and non-ATP competitive Src tyrosine kinase inhibitor with an IC ₅₀ of average 72 nM.	DMSO	86 mg/mL	5 mg
S2791	Sotrastaurin (AEB071)	A potent selective pan-PKC inhibitor and highly inhibits PKCθ with a Ki of 0.22 nM.	DMSO	87 mg/mL	5 mg



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