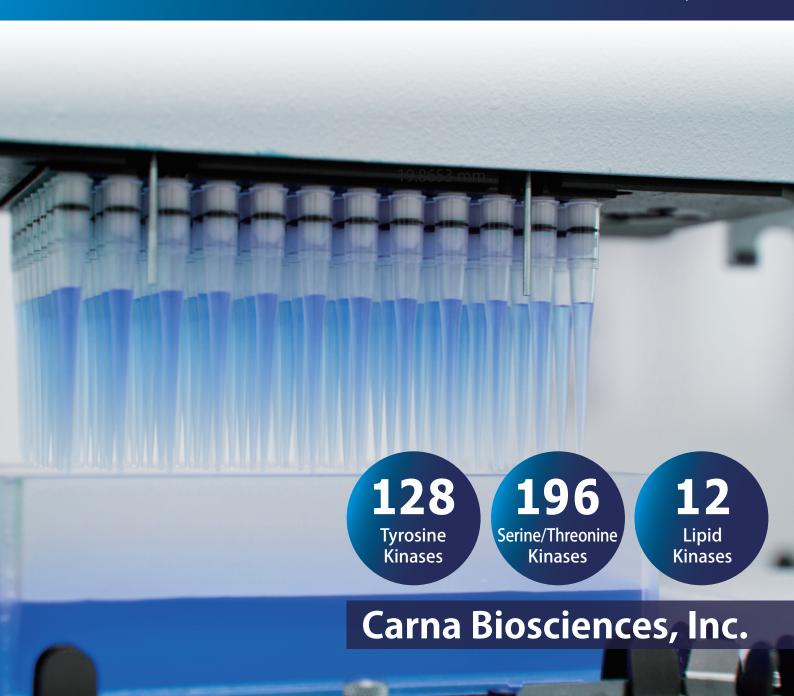
Custom and Pre-Selected Kinase Profiling to fit your Budget and Needs!

Kinase Profiling Book

As of July 1, 2021

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Profiling Assays available from Carna Biosciences, Inc.

As of July 1, 2021

Page	Kinase Name	Assay Platform
4	ABL(ABL1)	MSA
4	ABL(ABL1)[E255K]	MSA
4	ABL(ABL1)[T315I]	MSA
4	ACK(TNK2)	MSA
4 5 5 5 5 5 6	AKT1 AKT2	MSA MSA
5	AKT3	MSA
5	ALK	MSA
5	ALK[C1156Y]	MSA
5	ALK[F1174L]	MSA
6	ALK[G1202R] ALK[G1269A]	MSA MSA
6	ALK[C1203A] ALK[L1196M]	MSA
6	ALK[R1275Q]	MSA
6	ALK[T1151_L1152insT]	MSA
7 7 7 7	EML4-ALK NPM1-ALK	MSA MSA
7	AMPKa1/β1/γ1(PRKAA1/B1/G1)	MSA
7	AMPKα2/β1/γ1(PRKAA2/B1/G1)	MSA
	ARG(ABL2)	MSA
8	AurA(AURKA)	MSA
8 8	AurA(AURKA)/TPX2 AurB(AURKB)/INCENP	MSA MSA
8	AurC(AURKC)	MSA
8	ÄXL	MSA
9	BLK	MSA
9	BMX	MSA MSA
9	BRK(PTK6) BRSK1	MSA MSA
9	BRSK2	MSA
10	BTK	MSA
10	BTK[C481S]	MSA
10	BUB1/BUB3	MSA
10 10	CaMK1α(CAMK1) CaMK1δ(CAMK1D)	MSA MSA
11	CaMK2a(CAMK2A)	MSA
11	CaMK2β(CAMK2B)	MSA
11	CaMK2y(CAMK2G)	MSA
11 11	CaMK2δ(CAMK2D) CaMK4	MSA MSA
12	CDC2/CycB1	MSA
12	CDC7/ASK	MSA
12	CDK2/CycA2	MSA
12	CDK2/CycE1	MSA
12 13	CDK3/CycE1 CDK4/CycD3	MSA MSA
13	CDK 4 /CyCD3	MSA
13	CDK6/CycD3	MSA
13	CDK7/CycH/MAT1	MSA
13 14	CDK9/CycT1 CGK2(PRKG2)	MSA MSA
14	CGK2(PKKG2) CHK1(CHEK1)	MSA MSA
14	CHK2(CHEK2)	MSA
14	CK1a(CSNK1A1)	MSA
14	CK1y1(CSNK1G1)	MSA
15 15	CK1y2(CSNK1G2) CK1y3(CSNK1G3)	MSA MSA
15	CK1γ3(CSNK1G3) CK1δ(CSNK1D)	MSA MSA
15	CK1ɛ(CSNK1E)	MSA
15	CK2a1/β(CSNK2A1/B)	MSA
16 16	CK2α2/β(CSNK2A2/B) CLK1	MSA MSA
16 16	CLKI CLK2	MSA MSA
16	CLK3	MSA
16	CRIK(CIT)	MSA
17	CSK DARK1	MSA
17 17	DAPK1 DCAMKL2	MSA MSA
17	DDR1	MSA
17	DDR2	MSA
18	DYRK1A	MSA
18	DYRK1B DYRK2	MSA MSA
18 18	DYRKZ DYRK3	MSA MSA
18	EEF2K	MSA
19	EGFR	MSA
19	EGFR[C797S/L858R]	MSA MSA
19 19	EGFR[d746-750] EGFR[d746-750/C797S]	MSA MSA
19	EGFR[d746-750/C7975] EGFR[d746-750/T790M]	MSA MSA
20	EGFR[d746-750/T790M/C797S]	MSA
20	EGFR[D770_N771insNPG]	MSA
20	EGFR[L858R]	MSA MSA
20 20	EGFR[L861Q] EGFR[T790M]	MSA MSA
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21			=
Pith			=
EPHA4			=
EPHA5			
PPHA7			=
222 EPHAB MSA 233 EPHB1 MSA 234 EPHB2 MSA 235 EPHB3 MSA 236 ErK1(MAPK3) MSA 241 ErK2(MAPK1) MSA 242 ErK2(MAPK1) MSA 244 ErK2(MAPK7) MSA 244 FER MSA 244 FER MSA 244 FER MSA 245 FES MSA 246 FES MSA 247 FES MSA 255 FGFR1 MSA 255 FGFR2 MSA 266 FGFR3/K650E1 MSA 266 FGFR3/K650E1 MSA 266 FGFR3/K650E1 MSA 266 FGFR3/K555M1 MSA 276 FGFR4/W550E1 MSA 277 FGFR4/W550E1 MSA 278 FGFR4/W550E1 MSA 279 FGR			
EPHB1			
EPHB2			
EPHB4	23		
23 Erk1(MAPK1) MSA 244 Erk2(MAPK1) MSA 244 Erk5(MAPK7) MSA 244 FER MSA 245 FER MSA 246 FES MSA 255 FGFR1 MSA 255 FGFR2 MSA MSA 255 FGFR2 MSA MSA 255 FGFR3 MSA MSA 266 FGFR3 MSA MSA 266 FGFR3 MSASOMI MSA 267 FGFR4 MSA MSA 277 FGFR4 MSA MSA 277 FGFR4 MSA MSA 277 FGFR4 MSA MSA 288 FLT3 MSA			
24 Erk2(MAPKT) MSA 24 Erk5(MAPKT) MSA 24 FER MSA 24 FER MSA 24 FES MSA 25 FGFR1 MSA MSA 25 FGFR2 MSA MSA 25 FGFR2 MSA MSA 26 FGFR3 MSA MSA 26 FGFR3 MSOSDI MSA 27 FGFR4 MSA MSA 27 FGFR4 MSA MSA 27 FGFR4 MSSA MSA 27 FGFR4 MSSA MSA 28 </td <td></td> <td></td> <td></td>			
24 FAK(PTK2) MSA 24 FER MSA 24 FER MSA 25 FGFR1 MSA 25 FGFR1IV561M] MSA 25 FGFR2[V5641] MSA 25 FGFR3[MS650] MSA 26 FGFR3[K550] MSA 26 FGFR3[K555M] MSA 26 FGFR3[K555M] MSA 26 FGFR3[K555M] MSA 26 FGFR3[V555M] MSA 26 FGFR3[V555M] MSA 26 FGFR3[V555M] MSA 26 FGFR4[V550L] MSA 27 FGFR4[V550L] MSA 27 FGFR4[V550L] MSA 27 FGRR4[V550L] MSA 28 FLT1 MSA 28 FLT4 MSA 28 FN[CSF1R] MSA 28 FN[CSF1R] MSA 29 FSK38(GSK3B) MSA 29 </td <td>24</td> <td>Erk2(MAPK1)</td> <td>MSA</td>	24	Erk2(MAPK1)	MSA
24 FER MSA 25 FGFR1 MSA 25 FGFR1 V561M MSA 25 FGFR2 MSA MSA 25 FGFR2 V5641 MSA 25 FGFR3 V5650 MSA 26 FGFR3 K650H MSA 26 FGFR3 K650H MSA 26 FGFR3 V555L MSA 26 FGFR3 V555L MSA 26 FGFR3 V555L MSA 26 FGFR3 V555L MSA 26 FGFR4 V550E MSA 27 FGFR4 V550L MSA 27 FGFR4 V550L MSA 27 FGFR4 V550L MSA 28 FLT3 MSA 28 FLT3 MSA 28 FLT3 MSA 28 FK MSA 29 FYN[isoform a] MSA 29 FYN[isoform b] MSA 29 GSK3a(GSK3B) MSA 30			
25 FGFR1 V561M MSA 25 FGFR2 V564M MSA 25 FGFR2 V564M MSA 25 FGFR3 V565M MSA 26 FGFR3 K650M MSA 26 FGFR3 V555L MSA 26 FGFR3 V555M MSA 26 FGFR3 V555M MSA 26 FGFR4 V55DL MSA 26 FGFR4 V55DL MSA 26 FGFR4 V55DL MSA 26 FGFR4 V55DL MSA 27 FGFR4 V55DL MSA 27 FGR MSA 27 FGR MSA 27 FGR MSA 28 FLT3 MSA 28 FMS(CSF1R) MSA 28 FRK MSA 29 FYN[isoform a] MSA 29 FYN[isoform b] MSA 29 GSK3B(GSK3A) MSA 29 HASpin(GSG2) MSA			
25 FGFR1/V561M MSA 25 FGFR2 MSA 25 FGFR2 MSA 26 FGFR3/K650E MSA 26 FGFR3/K650M MSA 26 FGFR3/K55SL MSA 26 FGFR3/V55SM MSA 26 FGFR4/V55SM MSA 27 FGFR4/V55DE MSA 27 FGFR4/V55DL MSA 27 FGFR4/V55DL MSA 27 FGFR4/V55DL MSA 28 FLT1 MSA 29 FFRK MSA 28 FLT3 MSA 28 FMSCSF1R) MSA 28 FMSCSF1R) MSA 28 FYN[isoform a] MSA 29 FYN[isoform b] MSA 29 GSK38(GSK3A) MSA 29 GSK38(GSK3B) MSA 29 GSK38(GSK3B) MSA 30 HERZ(ERBE2) MSA			
25 FGFR2 MSA 25 FGFR2 MSA 26 FGFR3 MSA 27 FGFR4 MSA 27 FGR MSA 27 FGR MSA 4 FLT3 MSA 28 FLT3 MSA 28 FLT4 MSA 28 FYN[isoform a] MSA 29 FYN[isoform b] MSA 29 GSK3G(SSX3B) MSA 29 GSK3G(SSX3B) MSA 30			
25 FGFR3 MSA 26 FGFR3 K650E MSA 26 FGFR3 K650M MSA 26 FGFR3 V555L MSA 26 FGFR3 V55SM MSA 26 FGFR4 V550E MSA 27 FGFR4 V550E MSA 27 FGFR4 V550L MSA 27 FGR MSA 27 FGR MSA 27 FGR MSA 28 FLT3 MSA 28 FLT4 MSA 28 FKK MSA 28 FKK MSA 28 FYN[isoform a] MSA 29 GSK3a(GSK3A) MSA 29 FYN[isoform b] MSA 29 GSK3a(GSK3B) MSA 29 GSK3a(GSK3B) MSA 29 GSK3a(GSK3B) MSA 29 HCK MSA 30 HERV(ERBB4) MSA 30 HERV(ER	25		
26 FGFR3[K650E] MSA 26 FGFR3[K650M] MSA 26 FGFR3[V555L] MSA 26 FGFR3[V555M] MSA 27 FGR4[N535K] MSA 27 FGFR4[V550E] MSA 27 FGR MSA 27 FGR MSA 28 FLT1 MSA 28 FLT3 MSA 28 FLT4 MSA 28 FKK MSA 28 FYN[isoform a] MSA 29 GSK33G(GSK3A) MSA 29 GSK33G(GSK3B) MSA 29 GSK33G(GSK3B) MSA 29 HACK MSA 30 HER2(ERBB4) MSA 30 HER2(ERBB4) MSA 30 HIPK1 MSA 30 HIPK2 MSA 31 HIPK3 MSA 31 HIPK3 MSA 31 HIPK4	25	FGFR2[V564I]	MSA
26 FGFR3[K650M] MSA 26 FGFR3[V555L] MSA 26 FGFR3[V555M] MSA 26 FGFR4[N535K] MSA 27 FGFR4[N535K] MSA 27 FGRA[V550L] MSA 27 FGR MSA 28 FLT1 MSA 28 FLT3 MSA 28 FTK MSA 28 FTK MSA 28 FTYN[isoform a] MSA 29 GSK3G(GSK3A) MSA 29 GSK3G(GSK3B) MSA 29 GSK3G(GSK3B) MSA 29 HASpin(GSG2) MSA 30 HER2(ERBB2) MSA 30 HER4(ERBB4) MSA 30 HIPK1 MSA 31 HIPK2			
26 FGFR3 V555M MSA 26 FGFR4 27 FGFR4[N535K] MSA 27 FGFR4[V550E] MSA 27 FGFR4[V550E] MSA 27 FGR MSA 27 FGR MSA 27 FLT1 28 FLT3 28 FLT3 28 FK 28 FKK 28 FKK 28 FYN[isoform a] MSA 29 FYN[isoform b] MSA 29 GSK3G(GSK3A) MSA 29 GSK3G(GSK3B) MSA 29 HASPINIGSG2) MSA 30 HER2(ERBB2) MSA 30 HER2(ERBB2) MSA 30 HER4(ERBB4) MSA 30 HER4(ERBB4) MSA 30 HIPK1 MSA 31 HIPK2 MSA 31 HIPK3 MSA 31 HIPK3 MSA 31 HIPK3 MSA 32 IKK6(KBKB) MSA 33 IKK6(KBKE) MSA 34 IKK6(KBKE) MSA 35 </td <td></td> <td></td> <td>MSA</td>			MSA
26 FGFR4 MSA 27 FGFR4[N535K] MSA 27 FGRR4[V550E] MSA 27 FGRR MSA 27 FGR MSA 27 FLT1 MSA 28 FLT3 MSA 28 FLT4 MSA 28 FRK MSA 29 FYN[isoform a] MSA 29 GSK33(GSK33A) MSA 29 GSK33(GSK33A) MSA 29 HCK MSA 30 HER2(ERBB4) MSA 30 HER2(ERBB4) MSA 30 HIPK1 MSA 30 HIPK2 MSA 31 HIPK2 MSA 31 HPK1(MAP4K1) MSA 31 IKK6(CHUK) IMAP <t< td=""><td></td><td></td><td></td></t<>			
27 FGFR4[N550E] MSA 27 FGFR4[V550E] MSA 27 FGRR MSA 27 FGR MSA 28 FLT1 MSA 28 FLT3 MSA 28 FLT4 MSA 28 FSK MSA 28 FYN[isoform a] MSA 29 FYN[isoform b] MSA 29 GSK33(GSK3A) MSA 29 GSK33(GSK3B) MSA 29 HASPIN(GSG2) MSA 30 HER4(ERBB4) MSA 30 HER4(ERBB4) MSA 30 HIPK1 MSA 30 HIPK1 MSA 30 HIPK2 MSA 31 HIPK3 MSA 31 HIPK3 MSA 31 HIPK4 MSA 31 HIPK4 MSA 31 IKK6(IKBKB) MSA 32 INSR MSA			
27 FGRR # MSA 27 FGR MSA 27 FLT1 MSA 28 FLT4 MSA 28 FLT4 MSA 28 FSMS(CSF1R) MSA 28 FSMS(SSF1R) MSA 28 FYN[isoform a] MSA 29 FYN[isoform b] MSA 29 GSK3a(GSK3A) MSA 29 GSK3g(GSK3B) MSA 29 GSK3g(GSK3B) MSA 29 Haspin(GSG2) MSA 29 HASPIN(GSG2) MSA 30 HERZ(ERBB4) MSA 30 HERZ(ERBB4) MSA 30 HERZ(ERBB4) MSA 30 HIPK1 MSA 30 HIPK2 MSA 31 HIPK1 MSA 32 MSA HIPK2 MSA 31 HIPK2 MSA MSA 31 HIPK1 MSA MSA	27	FGFR4[N535K]	MSA
27 FGR MSA 27 FLT1 MSA 28 FLT3 MSA 28 FLT4 MSA 28 FSKS MSA 28 FFK MSA 28 FYN[isoform a] MSA 29 GSK3G(GSK3A) MSA 29 GSK3B(GSK3B) MSA 29 GSK3B(GSK3B) MSA 29 HSA MSA 30 HER2(ERBB2) MSA 30 HER4(ERBB4) MSA 30 HIPK1 MSA 31 HIPK1 MSA 31 HIPK2 MSA 31 HIPK3 MSA 31 HIPK4 MSA 31 HIPK4 MSA 31 HKK6(CHUK) IMA 32			
27 FLT1 MSA 28 FLT3 MSA 28 FMSCCSF1R) MSA 28 FRK MSA 28 FYN[isoform a] MSA 29 GSK3G(GSK3A) MSA 29 GSK3B(GSK3B) MSA 29 GSK3B(GSK3B) MSA 29 HCK MSA 30 HER2(ERBB2) MSA 30 HER2(ERBB2) MSA 30 HER4(ERBB4) MSA 30 HIPK1 MSA 30 HIPK1 MSA 30 HIPK1 MSA 31 HIPK2 MSA 31 HIPK3 MSA 31 HIPK4 MSA 31 HIPK4 MSA 31 IKK6(CHUK) IMAP 32 IKK6(IKBKB) MSA 33 IKK6(IKBKE) MSA 34 IKK6(IKBKE) MSA 33 JAK1 M			
28 FLT4 MSA 28 FMS(CSF1R) MSA 28 FRK MSA 29 FYN[isoform a] MSA 29 GSK3a(GSK3A) MSA 29 GSK3B(GSK3B) MSA 29 HCK MSA 30 HER2(ERBB2) MSA 30 HER4(ERBB4) MSA 30 HIPK1 MSA 30 HIPK1 MSA 30 HIPK1 MSA 30 HIPK2 MSA 31 HIPK2 MSA 31 HIPK3 MSA 31 HIPK3 MSA 31 HIPK4 MSA 31 HIPK1 (MAPK8I) MSA 32 INK8 (MSKB) MSA 33 IRK4 MSA	27	FLT1	MSA
28 FMS(CSF1R) MSA 28 FRK MSA 29 FYN[isoform a] MSA 29 GSK3α(GSK3A) MSA 29 GSK3β(GSK3B) MSA 29 HASpin(GSG2) MSA 30 HER2(ERBB2) MSA 30 HER4(ERBB4) MSA 30 HIPK1 MSA 30 HIPK1 MSA 30 HIPK2 MSA 31 HIPK2 MSA 31 HIPK3 MSA 31 HIPK4 MSA 31 HIPK4 MSA 31 HIPK1(MAP4K1) MSA 31 IKK6(CHUK) IMAP 32 IKK6(IKBKE) MSA 33 IKK6(IKBKE) MSA 32 IRAK1 IMAP 32 IRAK4 MSA 33 JAK1 MSA 34 JAK1 MSA 33 JAK2 M			_
28 FRK MSA 29 FYN[isoform a] MSA 29 FYN[isoform b] MSA 29 GSK3α(GSK3A) MSA 29 GSK3β(GSK3B) MSA 29 HASPINGGSG2) MSA 30 HER2(ERBB2) MSA 30 HER4(ERBB4) MSA 30 HIPK1 MSA 30 HIPK2 MSA 30 HIPK3 MSA 31 HIPK3 MSA 31 HIPK3 MSA 31 HIPK4 MSA 31 HIPK4 MSA 31 HIPK4 MSA 31 IGF1R MSA 31 IKK6(IKBK1) IMAP 32 IKK8(IKBKB) MSA 33 IKK6(IKBKE) MSA 34 IRAK1 IMAP 32 IRAK1 IMAP 33 JAK1 MSA 33 JAK1 MS			
29 FYN[isoform b] MSA 29 GSK3α(GSK3A) MSA 29 GSK3β(GSK3B) MSA 29 HCK MSA 30 HER2(ERBB2) MSA 30 HER4(ERBB4) MSA 30 HGK(MAP4K4) MSA 30 HIPK1 MSA 30 HIPK2 MSA 31 HIPK3 MSA 31 HIPK3 MSA 31 HIPK4 MSA 31 HFK1(MAP4K1) MSA 31 JEFIR MSA 31 JIKK6(CHUK) IMAP 32 IKK8(IKBKB) MSA 33 JIKK6(IKBKB) MSA 34 JIKK6(IKBKE) MSA 33 JIKK6(IKBKE) MSA 33 JAK1 MSA 33 JAK1 MSA 33 JAK1 MSA 34 JNK1(MAPK8) MSA 34 JNK2(MAPK9	28	FRK	MSA
29 GSK3a(GSK3A) MSA 29 GSK3B(GSK3B) MSA 29 Haspin(GSG2) MSA 30 HER2(ERBB2) MSA 30 HER4(ERBB4) MSA 30 HIPK1 MSA 30 HIPK1 MSA 30 HIPK2 MSA 31 HIPK3 MSA 31 HIPK3 MSA 31 HIPK4 MSA 31 IGF1R MSA 31 IKK6(CHUK) IMAP 32 IKK6(IKBKB) MSA 33 IKK6(IKBKE) MSA 34 IKK6(INSRR) MSA 33 IRAK1 IMAP 33 JAK1 MSA 33 JAK1 MSA 33 JAK3 MSA <td></td> <td></td> <td></td>			
29 Haspin(GSG2) MSA 30 HER2(ERBB2) MSA 30 HER4(ERBB4) MSA 30 HER4(ERBB4) MSA 30 HIPK1 MSA 30 HIPK2 MSA 31 HIPK2 MSA 31 HIPK3 MSA 31 HIPK4 MSA 31 HIPK4 MSA 31 HIPK1(MAP4K1) MSA 31 IKK6(CHUK) IMAP 32 IKK8(IKBKB) MSA 33 IKK6(CHUK) IMAP 32 IKK8(IKBKE) MSA 33 IKK6(IKBKE) MSA 34 IKKE (IKBKE) MSA 33 IRAK1 IMAP 32 IRAK4 MSA 33 IRR(INSRR) MSA 33 JAK1 MSA 33 JAK2 MSA 33 JAK3 MSA 34 JNK1(MAPK8)			
29 HCK MSA 30 HER2(ERBB2) MSA 30 HER4(ERBB4) MSA 30 HGK(MAP4K4) MSA 30 HIPK1 MSA 30 HIPK2 MSA 31 HIPK3 MSA 31 HPK1(MAP4K1) MSA 31 HPK1(MAP4K1) MSA 31 IGF1R MSA 31 IKKα(CHUK) IMAP 32 IKKβ(IKBKB) MSA 32 IKKβ(IKBKB) MSA 32 IKKKE(IKBKE) MSA 32 IRAK1 IMAP 32 IRAK1 IMAP 33 IRR(INSRR) MSA 33 JAK1 MSA 33 JAK1 MSA 33 JAK2 MSA 33 JAK3 MSA 34 JNK1(MAPK8) MSA 34 JNK2(MAPK10) MSA 34 KIT M	29	GSK3β(GSK3B)	
HER2(ERBB2) MSA HER4(ERBB4) MSA HER4(ERBB4) MSA HIPK1 MSA HIPK1 MSA HIPK2 MSA HIPK3 MSA HIPK4 MSA HIPK4 MSA HIPK4 MSA HIPK1 MSA HIPK4 MSA HIPK1 MSA HIPK4 MSA HIPK1 MSA HIPK3 MSA HIPK4 M			
30 HGK(MAP4K4) MSA 30 HIPK1 MSA 31 HIPK2 MSA 31 HIPK4 MSA 31 HPK1(MAP4K1) MSA 31 IGF1R MSA 31 IKKa(CHUK) IMAP 32 IKKβ(IKBKB) MSA 32 IKKε(IKBKE) MSA 32 IRAK1 IMAP 32 IRAK1 IMAP 32 IRAK1 IMAP 33 IR(INSRR) MSA 33 IRK(INSRR) MSA 33 JAK1 MSA 33 JAK2 MSA 33 JAK3 MSA 34 JNK1(MAPK8) MSA 34 JNK2(MAPK9) MSA 34 JNK3(MAPK10) MSA 34 KIT[D816E] MSA 35 KIT[D816F] MSA 35 KIT[D816F] MSA 35 KIT[D816F]			
HIPK1			
HIPK2			
HIPK4			
HPK1 (MAP4K1) MSA	-		
31 IGF1R MSA 31 IKKα(CHUK) IMAP 32 IKKβ(IKBKB) MSA 32 IKKβ(IKBKE) MSA 32 INSR MSA 32 IRAK1 IMAP 32 IRAK1 IMAP 32 IRAK4 MSA 33 IRR(INSRR) MSA 33 JAK1 MSA 33 JAK1 MSA 33 JAK2 MSA 33 JAK3 MSA 34 JNK1(MAPK8) MSA 34 JNK2(MAPK9) MSA 34 JNK3(MAPK10) MSA 34 KIT MSA 34 KIT MSA 35 KIT[D816E] MSA 35 KIT[D816V] MSA </td <td>-</td> <td></td> <td></td>	-		
32	31	IGF1R	MSA
32 IKKɛ(IKBKE) MSA 32 INSR MSA 32 INSR MSA 33 IRAK1 IMAP 33 IRR(INSRR) MSA 33 IRR(INSRR) MSA 33 JAK1 MSA 33 JAK2 MSA 34 JAK3 MSA 34 JAK3 MSA 34 JAK3 MSA 35 JAK3 MSA 36 JAK3 MSA 37 KITT MSA 38 JAK3 MSA 39 JAK3 MSA 30 JAK3 MSA 31 JAK3 MSA 32 JAK3 MSA 33 JAK3 MSA 34 JAK3 MSA 35 JAK3 MSA 36 JAK3 MSA 37 LYNBA 38 JAK3 MSA 39 JAK3 MSA 39 JAK3 MSA 30 JAK3 MSA 31 JAK3 MSA 32 JAK3 MSA 33 JAK3 MSA 34 JAK3 MSA 35 JAK3 MSA 36 JAK3 MSA 37 JAK3 MSA 38 KIT[D816E] MSA 38 KIT[D816F] MSA 39 KIT[D816Y] MSA 31 JAK3 MSA 32 JAK3 MSA 33 JAK1 MSA 34 JAK3 MSA 35 KIT[D816F] MSA 36 KIT[D816F] MSA 37 KIT[D816F] MSA 38 KIT[D816F] MSA 38 KIT[D816F] MSA 39 KIT[D816F] MSA 31 JAK1 MSA 32 KIT[D816F] MSA 33 JAK1 MSA 34 JAK1 MSA 35 KIT[D816F] MSA 36 KIT[D816F] MSA 37 KIT[D816F] MSA 38 KIT[D816F] MSA 38 KIT[D816F] MSA 39 KIT[D816F] MSA 31 KIT[D816F] MSA 32 KIT[D816F] MSA 33 JAK1 MSA 34 JAK1 MSA 35 KIT[D816F] MSA 36 KIT[D816F] MSA 37 KIT[D816F] MSA 38 KIT[D816F] MSA 38 KIT[D816F] MSA 39 KIT[D816F] MSA 31 KIT[D816F] MSA 31 KIT[D816F] MSA 32 KIT[D816F] MSA 33 MSA 34 KIT MSA 35 KIT[D816F] MSA 36 KIT[D816F] MSA 37 KIT MSA 37 MSA 38 MSA 38 MSA 39 MSA			
32 INSR MSA 32 IRAK1 IMAP 32 IRAK4 MSA 33 IRR(INSRR) MSA 33 JIK1 MSA 33 JAK1 MSA 33 JAK2 MSA 34 JNK1(MAPK8) MSA 34 JNK2(MAPK9) MSA 34 JNK3(MAPK10) MSA 34 KIT MSA 34 KIT MSA 35 KIT[D816E] MSA 35 KIT[D816V] MSA 36 KIT[V856G]			
32 IRAK4 MSA 33 IRR(INSRR) MSA 33 ITK MSA 33 JAK1 MSA 33 JAK2 MSA 33 JAK3 MSA 34 JNK1(MAPK8) MSA 34 JNK2(MAPK9) MSA 34 JNK3(MAPK10) MSA 34 KIT MSA 34 KIT MSA 35 KIT[D816E] MSA 35 KIT[D816V] MSA 36 KIT[V550G] MSA 36 LATS2 MSA 36 LCK MSA 36 LCK MSA 37 LYNa MSA <td></td> <td>INSR</td> <td></td>		INSR	
33 IRR(INSRR) MSA 33 ITK MSA 33 JAK1 MSA 33 JAK2 MSA 34 JNK1(MAPK8) MSA 34 JNK2(MAPK10) MSA 34 KDR MSA 34 KIT MSA 34 KIT MSA 35 KIT[D816E] MSA 35 KIT[D816V] MSA 35 KIT[T670I] MSA 35 KIT[V560G] MSA 36 LATS2 MSA 36 LCK MSA 36 LCK MSA 36 LOK(STK10) MSA 37 LYNa MSA 37 MAP4K2 MSA 37 MAPKAPK2 MSA			
33 JAK1 MSA 33 JAK2 MSA 34 JNK1(MAPK8) MSA 34 JNK2(MAPK9) MSA 34 JNK3(MAPK10) MSA 34 KDR MSA 34 KIT MSA 35 KIT[D816E] MSA 35 KIT[D816V] MSA 35 KIT[D816V] MSA 35 KIT[T670I] MSA 35 KIT[V560G] MSA 36 KIT[V5654A] MSA 36 LATS2 MSA 36 LCK MSA 36 LOK(STK10) MSA 36 LTK MSA 37 LYNa MSA 37 MAP4K2 MSA 37 MAPKAPK2 MSA	33	IRR(INSRR)	MSA
33 JAK2 MSA 33 JAK3 MSA 34 JNK1(MAPK9) MSA 34 JNK3(MAPK10) MSA 34 KDR MSA 34 KIT MSA 35 KIT[D816E] MSA 35 KIT[D816V] MSA 35 KIT[D816V] MSA 35 KIT[T670I] MSA 35 KIT[V560G] MSA 36 KIT[V564A] MSA 36 LATS2 MSA 36 LCK MSA 36 LOK(STK10) MSA 36 LTK MSA 37 LYNa MSA 37 MAP4K2 MSA 37 MAPKAPK2 MSA			
33 JAK3 MSA 34 JNK1(MAPK8) MSA 34 JNK2(MAPK9) MSA 34 JNK3(MAPK10) MSA 34 KDR MSA 34 KIT MSA 35 KIT[D816E] MSA 35 KIT[D816V] MSA 35 KIT[T670I] MSA 35 KIT[V560G] MSA 36 KIT[V560G] MSA 36 LATS2 MSA 36 LCK MSA 36 LCK MSA 37 LYNA MSA 37 LYNA MSA 37 MAP4K2 MSA 37 MAP4K2 MSA			
34 JNK2(MAPK9) MSA 34 JNK3(MAPK10) MSA 34 KDR MSA 34 KIT MSA 35 KIT[D816E] MSA 35 KIT[D816V] MSA 35 KIT[D816V] MSA 35 KIT[T670I] MSA 35 KIT[V560G] MSA 36 KIT[V564A] MSA 36 LATS2 MSA 36 LATS2 MSA 36 LOK(STK10) MSA 37 LYNB MSA 37 LYNB MSA 37 MAP4K2 MSA 37 MSA 37 MAP4K2 MSA	33	JAK3	MSA
34 JNK3(MAPK10) MSA 34 KDR MSA 34 KIT MSA 35 KIT[D816E] MSA 35 KIT[D816V] MSA 35 KIT[D816V] MSA 35 KIT[T670I] MSA 35 KIT[V560G] MSA 36 KIT[V560G] MSA 36 LATS2 MSA 36 LCK MSA 36 LOK(STK10) MSA 36 LTK MSA 37 LYNa MSA 37 LYNb MSA 37 MAP4K2 MSA 37 MAPKAPK2 MSA		, ,	
34 KDR MSA 34 KIT MSA 35 KIT[D816E] MSA 35 KIT[D816V] MSA 35 KIT[D816Y] MSA 35 KIT[T670I] MSA 36 KIT[V560G] MSA 36 LATS2 MSA 36 LCK MSA 36 LOK(STK10) MSA 36 LTK MSA 37 LYNa MSA 37 LYNb MSA 37 MAP4K2 MSA 37 MAPKAPK2 MSA			
35 KIT[D816E] MSA 35 KIT[D816V] MSA 35 KIT[D816V] MSA 35 KIT[D816Y] MSA 35 KIT[T670I] MSA 36 KIT[V560G] MSA 36 LATS2 MSA 36 LATS2 MSA 36 LCK MSA 36 LCK MSA 37 LYNA MSA 37 LYNA MSA 37 MAP4K2 MSA 37 MAP4K2 MSA			
35 KIT[D816V] MSA 35 KIT[D816Y] MSA 35 KIT[T670I] MSA 35 KIT[T670I] MSA 36 KIT[V560G] MSA 36 LATS2 MSA 36 LATS2 MSA 36 LCK MSA 36 LCK MSA 37 LYNa MSA 37 LYNa MSA 37 LYNb MSA 37 MAP4K2 MSA 37 MAP4K2 MSA			
35 KIT[T670I] MSA 35 KIT[V560G] MSA 36 KIT[V564A] MSA 36 LATS2 MSA 36 LCK MSA 36 LOK(STK10) MSA 36 LYNa MSA 37 LYNa MSA 37 MAP4K2 MSA 37 MAPKAPK2 MSA	35	KIT[D816V]	MSA
35 KIT[V560G] MSA 36 KIT[V654A] MSA 36 LATS2 MSA 36 LCK MSA 36 LCK MSA 36 LOK(STK10) MSA 37 LYNa MSA 37 LYNb MSA 37 MAP4K2 MSA 37 MAPKAPK2 MSA			
36 KITTV654AT MSA 36 LATS2 MSA 36 LCK MSA 36 LOK(STK10) MSA 36 LTK MSA 37 LYNa MSA 37 LYNb MSA 37 MAP4K2 MSA 37 MAPKAPK2 MSA			
36 LCK MSA 36 LOK(STK10) MSA 36 LTK MSA 37 LYNa MSA 37 LYNb MSA 37 MAP4K2 MSA 37 MAPKAPK2 MSA	36	KITĪV654AĪ	MSA
36 LOK(STK10) MSA 36 LTK MSA 37 LYNa MSA 37 LYNb MSA 37 MAP4K2 MSA 37 MAPKAPK2 MSA			
36 LTK MSA 37 LYNa MSA 37 LYNb MSA 37 MAP4K2 MSA 37 MAPKAPK2 MSA			
37 LYNb MSA 37 MAP4K2 MSA 37 MAPKAPK2 MSA	36	ĹТК	MSA
37 MAP4K2 MSA 37 MAPKAPK2 MSA			
37 MAPKAPK2 MSA	37		

Page	Kinase Name	Assay Platform
38	MAPKAPK5	MSA
38 38	MARK1 MARK2	MSA MSA
38	MARK3	MSA
38 39	MARK4 MELK	MSA MSA
39	MER(MERTK)	MSA
39 39	MET MET[D1228H]	MSA MSA
39	MET[D1226H] MET[M1250T]	MSA MSA
40	MET[Y1235D]	MSA
40 40	MINK(MINK1) MNK1(MKNK1)	MSA MSA
40	MNK2(MKNK2)	MSA
40 41	MRCKα(CDC42BPA) MRCKβ(CDC42BPB)	MSA MSA
41	MSK1(RPS6KA5)	MSA
41 41	MSK2(RPS6KA4)	MSA MSA
41	MSSK1(STK23) MST1(STK4)	MSA MSA
42	MST2(STK3)	MSA
42 42	MST3(STK24) MST4	MSA MSA
42	MUSK	MSA
42 43	NDR1(STK38) NDR2(STK38L)	MSA MSA
43	NDRZ(STR38L) NEK1	MSA MSA
43	NEK2	MSA
43 43	NEK4 NEK6	MSA MSA
44	NEK7	MSA
44 44	NEK9 NIM1K(MGC42105)	MSA MSA
44	NuaK1	MSA
44 45	NuaK2 p38a(MAPK14)	MSA MSA
45	p38G(MAPK14) p38β(MAPK11)	MSA MSA
45	p38y(MAPK12)	MSA
45 45	p38δ(MAPK13) p70S6K(RPS6KB1)	MSA MSA
46	p70S6Kβ(RPS6KB2)	MSA
46 46	PAK1 PAK2	MSA MSA
46	PAK4	MSA
46	PAK5(PAK7)	MSA
47 47	PAK6 PASK	MSA MSA
47	PBK	MSA
47 47	PDGFRa(PDGFRA) PDGFRa(PDGFRA)[D842V]	MSA MSA
48	PDGFRa(PDGFRA)[T674I]	MSA
48 48	PDGFRa(PDGFRA)[V561D] PDGFRβ(PDGFRB)	MSA MSA
48	PDHK2(PDK2)	MSA
48 49	PDHK4(PDK4) PDK1(PDPK1)	MSA MSA
49	PEK(EIF2AK3)	IMAP
49 49	PGK(PRKG1)	MSA MSA
49	PHKG1 PHKG2	MSA MSA
50	PIK3CA/PIK3R1	ADP-Glo
50 50	PIK3CB/PIK3R1 PIK3CD/PIK3R1	ADP-Glo ADP-Glo
50	PIKFYVĖ(PIP5K3)	ADP-Glo
50 51	PIM1 PIM2	MSA MSA
51	PIM3	MSA
51 51	PIP4K2A PIP4K2B	ADP-Glo ADP-Glo
51	PIP5K1A	ADP-Glo
52 52	PIP5K1B PIP5K1C	ADP-Glo ADP-Glo
52	PIP5KIC PIP5KL1	ADP-Glo
52 52	PKACQ(PRKACA)	MSA MSA
52	PKACβ(PRKACB) PKACγ(PRKACG)	MSA MSA
53	PKCa(PRKCA)	MSA
53 53	PKCβ1(PRKCB1) PKCβ2(PRKCB2)	MSA MSA
53	PKĊy(PRKCG) ´	MSA
54 54	PKCδ(PRKCD) PKCε(PRKCE)	MSA MSA
54	PKCζ(PRKCZ)	MSA
54 54	PKCŋ(PRKCH) PKCθ(PRKCO)	MSA MSA
55	PKCi(PRKCI)	MSA
55 55	PKD1(PRKD1) PKD2(PRKD2)	MSA MSA
55	PKD2(PKND2) PKD3(PRKD3)	MSA MSA
55	PKN1	IMAP

Page	Kinase Name	Assay Platform
56	PKR(EIF2AK2)	IMAP
56	PLK1	MSA
56 56	PLK2 PLK3	IMAP MSA
56	PRKX	MSA MSA
57	PYK2(PTK2B)	MSA
57	QIK(SNF1LK2)	MSA
57	RET	MSA
57 57	RET[G691S]	MSA MSA
58	RET[M918T] RET[S891A]	MSA MSA
58	RET[Y791F]	MSA
58	ROCK1	MSA
58	ROCK2	MSA
58	RON(MST1R)	MSA
59 59	ROS(ROS1) RSK1(RPS6KA1)	MSA MSA
59	RSK2(RPS6KA3)	MSA
59	RSK3(RPS6KA2)	MSA
59	RSK4(RPS6KA6)	MSA
60	SGK	MSA
60 60	SGK2 SGK3(SGKL)	MSA MSA
60	SIK(SNF1LK)	MSA MSA
60	skMLCK(MYLK2)	MSA
61	SLK	MSA
61	SPHK1	MSA
61 61	SPHK2 SRC	MSA MSA
61	SRM(SRMS)	MSA MSA
62	SRPK1	IMAP
62	SRPK2	MSA
62	SYK	MSA
62 62	TAOK2 TBK1	MSA MSA
63	TEC	MSA
63	TIE2(TEK)	MSA
63	TNIK	MSA
63	TNK1	MSA
63 64	TRKA(NTRK1) TRKB(NTRK2)	MSA MSA
64	TRKC(NTRK3)	MSA
64	TSSK1	MSA
64	TSSK2	MSA
64	TSSK3	MSA
65 65	TXK TYK2	MSA MSA
65	TYRO3	MSA
65	WNK1	MSA
65	WNK2	MSA
66	WNK3	MSA
66 66	YES(YES1) YES(YES1)[T348I]	MSA MSA
66	ZAP70	MSA MSA
	- W / V	1 13/1

<< Cascade Assay >>

Page	Kinase Name	Assay Platform
67	BRAF	MSA
67	BRAF[V600E]	MSA
67	COT(MAP3K8)	MSA
67	DLK(MAP3K12)	MSA
67	MAP2K1	MSA
68	MAP2K2	MSA
68	MAP2K3	MSA
68	MAP2K4	MSA
68	MAP2K5	MSA
68	MAP2K6	MSA
69	MAP2K7	MSA
69	MAP3K1	MSA
69	MAP3K2	MSA
69	MAP3K3	MSA
69	MAP3K4	MSA
70	MAP3K5	MSA
70	MLK1(MAP3K9)	MSA
70	MLK2(MAP3K10)	MSA
70	MLK3(MAP3K11)	MSA
70	MOS	MSA
71	RAF1	MSA
71	TAK1-TAB1(MAP3K7)	MSA

- The Kinase Company - Carna Biosciences, Inc.

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www.carnabio.com



ABL(ABL1)

Product code 08-001

Full-length human ABL [2-1130(end) amino acids of accession number NP_005148.2] was expressed as N-terminal His-tagged protein (126 kDa) using baculovirus expression system. His-tagged ABL was purified by using Ni-NTA affinity chromatography.

Assay platform : Mobility Shift Assay

Substrate : ABLtide

ATP (μ M) Km app / Bin : 16 / 25 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 75 IC50 at 1 mM ATP (nM) : 1300

ABL(ABL1)[E255K]

Product code 08-094

Full-length human ABL [2-1130(end) amino acids and E255K of accession number NP_005148.2] was expressed as N-terminal Histagged protein (126 kDa) using baculovirus expression system. Histagged ABL[E255K] was purified by using Ni-NTA affinity chromatography.

Assay platform : Mobility Shift Assay

Substrate : ABLtide

ATP (μ M) Km app / Bin : 17 / 25 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 140 IC50 at 1 mM ATP (nM) : 4500

ABL(ABL1)[T315I]

Product code 08-093

Full-length human ABL [2-1130(end) amino acids and T315I of accession number NP_005148.2] was expressed as N-terminal Histagged protein (126 kDa) using baculovirus expression system. Histagged ABL[T315I] was purified by using Ni-NTA affinity chromatography.

Assay platform : Mobility Shift Assay

Substrate : ABLtide

ATP (μ M) Km app / Bin : 4 / 5 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 6.4 IC50 at 1 mM ATP (nM) : 890

ACK(TNK2)

Product code 08-196

Human ACK, catalytic domain [110-476 amino acids of accession number NP_005772.3] was expressed as N-terminal GST-fusion protein (69 kDa) using baculovirus expression system. GST-ACK was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : WASP peptide

ATP (μ M) Km app / Bin : 97 / 100

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 3.2 IC50 at 1 mM ATP (nM) : 3.8

AKT1

Product code 01-101

Human AKT1, catalytic domain [104-480(end) amino acids of accession number NP_005154.1] was co-expressed as N-terminal GST-fusion protein (70 kDa) with His-tagged PDK1 [1-556(end) amino acids of accession number NP_002604.1] using baculovirus expression system. GST-AKT1 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Crosstide

ATP (µM) Km app / Bin : 31 / 50

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 2.7 IC50 at 1 mM ATP (nM) : 22



AKT2

Product code 01-102

Human AKT2, catalytic domain [120-481(end) amino acids of accession number NP_001617.1] was co-expressed as N-terminal GST-fusion protein (69 kDa) with His-tagged PDK1 [1-556(end) amino acids of accession number NP_002604.1] using baculovirus expression system. GST-AKT2 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Crosstide

ATP (μ M) Km app / Bin : 110 / 100

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 5.2 IC50 at 1 mM ATP (nM) : n.a.

AKT3

Product code 01-103

Human AKT3, catalytic domain [108-479(end) amino acids of accession number NP_005456.1] was co-expressed as N-terminal GST-fusion protein (70 kDa) with His-tagged PDK1 [1-556(end) amino acids of accession numberNP_002604.1] using baculovirus expression system. GST-AKT3 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Crosstide

ATP (μM) Km app / Bin : 54 / 50

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 3.2 IC50 at 1 mM ATP (nM) : n.a.

ALK

Product code 08-518

Human ALK, cytoplasmic domain [1058-1620(end) amino acids of accession number NP_004295.2] was expressed as N-terminal GST-fusion protein (90 kDa) using baculovirus expression system. GST-ALK was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 57 / 50 Metal : Ma

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 2.5 IC50 at 1 mM ATP (nM) : 15

ALK[C1156Y]

Product code 08-530

Human ALK, cytoplasmic domain [1058-1620(end) amino acids and C1156Y of accession number NP_004295.2] was expressed as N-terminal GST-fusion protein (90 kDa) using baculovirus expression system. GST-ALK[C1156Y] was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 64 / 75 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 1.9 IC50 at 1 mM ATP (nM) : 11

ALK[F1174L]

Product code 08-519

Human ALK, cytoplasmic domain [1058-1620(end) amino acids and F1174L of accession number NP_004295.2] was expressed as N-terminal GST-fusion protein (90 kDa) using baculovirus expression system. GST-ALK[F1174L] was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (µM) Km app / Bin : 49 / 50

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 2.4 IC50 at 1 mM ATP (nM) : 21



ALK[G1202R]

Product code 08-544

Human ALK, cytoplasmic domain [1058-1620(end) amino acids and G1202R of accession number NP_004295.2] was expressed as N-terminal GST-fusion protein (90 kDa) using baculovirus expression system. GST-ALK[G1202R] was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 31 / 50 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 7.3 IC50 at 1 mM ATP (nM) : 69

ALK[G1269A]

Product code 08-537

Human ALK , cytoplasmic domain [1058-1620(end) amino acids and G1269A of accession number NP_004295.2] was expressed as N-terminal GST-fusion protein (90 kDa) using baculovirus expression system. GST-ALK[G1269A] was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 27 / 25 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.36 IC50 at 1 mM ATP (nM) : 1.6

ALK[L1196M]

Product code 08-529

Human ALK, cytoplasmic domain [1058-1620(end) amino acids and L1196M of accession number NP_004295.2] was expressed as N-terminal GST-fusion protein (90 kDa) using baculovirus expression system. GST-ALK[L1196M] was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 57 / 75 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.66 IC50 at 1 mM ATP (nM) : 4.3

ALK[R1275Q]

Product code 08-520

Human ALK, cytoplasmic domain [1058-1620(end) amino acids and R1275Q of accession number NP_004295.2] was expressed as N-terminal GST-fusion protein (90 kDa) using baculovirus expression system. GST-ALK[R1275Q] was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 84 / 100

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 3.3 IC50 at 1 mM ATP (nM) : 16

ALK[T1151_L1152insT]

Product code 08-539

Human ALK, cytoplasmic domain [1058-1620(end) amino acids and T1151_L1152insT of accession number NP_004295.2] was expressed as N-terminal GST-fusion protein (90 kDa) using baculovirus expression system. GST-ALK[T1151_L1152insT] was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 110 / 100

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 6.5 IC50 at 1 mM ATP (nM) : 16



EML4-ALK

Product code 08-516

Fused gene of human fusion EML4-ALK [1-1059 amino acids of accession number BAF73611.1] was expressed as N-terminal GST-fusion protein (145 kDa) using baculovirus expression system. GST-EML4-ALK was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 43 / 50 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 1.9 IC50 at 1 mM ATP (nM) : 16

NPM1-ALK

Product code 08-517

Fused gene of human fusion NPM1-ALK [1-680 amino acids of accession number BAA08343.1] was expressed as N-terminal GST-fusion protein (103kDa) using baculovirus expression system. GST-NPM1-ALK was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 57 / 50 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 2.4 IC50 at 1 mM ATP (nM) : 14

 $AMPK\alpha1/\beta1/\gamma1(PRKAA1/B1/G1)$

Product code 02-113

Full-length human AMPK α 1 [1-550(end) amino acids of accession number NP_006242.4] was co-expressed as N-terminal GST-fusion protein (90 kDa) with GST-PRKAB1 [1-270(end) amino acids of accession number NP_006244.2] and PRKAG1 [1-331(end) amino acids of accession number NP_002724.1] using baculovirus expression system. GST-AMPK α 1/ β 1/ γ 1 was purified by using glutathione sepharose chromatography and activated with Histagged CaMKK1. Activated GST-AMPK α 1/ β 1/ γ 1 was purified using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay
Substrate : SAMS peptide

ATP (μ M) Km app / Bin : 130 / 150

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.41 IC50 at 1 mM ATP (nM) : 0.87

 $AMPK\alpha 2/\beta 1/\gamma 1(PRKAA2/B1/G1)$

Product code 02-114

Full-length human AMPK α 2 [1-552(end) amino acids of accession number NP_006243.2] was co-expressed as N-terminal GST-fusion protein (89 kDa) with GST-PRKAB1 [1-270(end) amino acids of accession number NP_006244.2] and PRKAG1 [1-331(end) amino acids of accession number NP_002724.1] using baculovirus expression system. GST-AMPK α 2/ β 1/ γ 1 was purified by using glutathione sepharose chromatography and activated with Histagged CaMKK1. Activated GST-AMPK α 2/ β 1/ γ 1 was purified using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : SAMS peptide

ATP (µM) Km app / Bin : 100 / 100

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.79 IC50 at 1 mM ATP (nM) : n.a.

ARG(ABL2)

Product code 08-102

Truncated human ARG [2-52, 74-1182(end) amino acids of accession number NP_009298.1] was expressed as N-terminal GST-fusion protein (153 kDa) using baculovirus expression system. GST-ARG was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : ABLtide

ATP (μ M) Km app / Bin : 24 / 25 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 27 IC50 at 1 mM ATP (nM) : 400



AurA(AURKA)

Product code 05-101

Full-length human AurA [1-403(end) amino acids of accession number NP_940835.1] was expressed as N-terminal GST-fusion protein (73 kDa) using baculovirus expression system. GST-AurA was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Kemptide

ATP (μ M) Km app / Bin : 27 / 25 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 1.8 IC50 at 1 mM ATP (nM) : 17

AurA(AURKA)/TPX2

Product code 05-186

Full-length human AurA [1-403(end) amino acids of accession number NP_940835.1] was expressed as N-terminal GST-fusion protein (73 kDa) using baculovirus expression system. GST-AurA was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Kemptide

ATP (μ M) Km app / Bin : 1.7 / 2 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 6.1 IC50 at 1 mM ATP (nM) : n.a.

AurB(AURKB)/INCENP

Product code 05-102

Full-length human AurB [1-344(end) amino acids of accession number NP_004208.2] was co-expressed as N-terminal GST-fusion protein (66 kDa) with His-tagged INCENP(INBOX) [803-918(end) amino acids of accession number NP_001035784.1] using baculovirus expression system. GST-AurB was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Kemptide

ATP (μ M) Km app / Bin : 16 / 25 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 7.1 IC50 at 1 mM ATP (nM) : 62

AurC(AURKC)

Product code 05-103

Full-length human AurC [1-275(end) amino acids of accession number NP_003151.2] was expressed as N-terminal GST-fusion protein (59 kDa) using baculovirus expression system. GST-AurC was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Kemptide

ATP (μ M) Km app / Bin : 24 / 25 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 3.1 IC50 at 1 mM ATP (nM) : 18

AXL

Product code 08-107

Human AXL, cytoplasmic domain [464-885(end) amino acids of accession number NP_001690.2] was expressed as N-terminal GST-fusion protein (74 kDa) using baculovirus expression system. GST-AXL was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CSKtide

ATP (μ M) Km app / Bin : 32 / 50 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 1.1 IC50 at 1 mM ATP (nM) : 7.9



BLK

Product code 08-164

Full-length human BLK [1-505(end) amino acids of accession number NP_001706.2] was expressed as N-terminal GST-fusion protein (85 kDa) using baculovirus expression system. GST-BLK was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 62 / 75 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 2.6 IC50 at 1 mM ATP (nM) : 17

BMX

Product code 08-179

Full-length human BMX [1-675(end) amino acids of accession number NP_001712.1] was expressed as N-terminal GST-fusion protein (105 kDa) using baculovirus expression system. GST-BMX was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 75 / 75 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 10 IC50 at 1 mM ATP (nM) : 45

BRK(PTK6)

Product code 08-165

Full-length human BRK [2-451(end) amino acids of accession number NP_005966.1] was expressed as N-terminal GST-fusion protein (79 kDa) using baculovirus expression system. GST-BRK was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Blk/Lyntide

ATP (μ M) Km app / Bin : 250 / 250

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 260 IC50 at 1 mM ATP (nM) : 390

BRSK1

Product code 02-115

Full-length human BRSK1 [1-778(end) amino acids of accession number NP_115806.1] was expressed as N-terminal GST-fusion protein (112 kDa) using baculovirus expression system. GST-BRSK1 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CHKtide

ATP (μ M) Km app / Bin : 30 / 25 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.27 IC50 at 1 mM ATP (nM) : 0.57

BRSK2

Product code 02-116

Full-length human BRSK2 [1-674(end) amino acids of accession number ABA17261.1] was expressed as N-terminal GST-fusion protein (102 kDa) using baculovirus expression system. GST-BRSK2 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CHKtide

ATP (μ M) Km app / Bin : 31 / 50 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.31 IC50 at 1 mM ATP (nM) : n.a.



BTK

Product code 08-180

Full-length human BTK [2-659(end) amino acids of accession number NP_000052] was expressed as N-terminal GST-fusion protein (103 kDa) using baculovirus expression system. GST-BTK was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 22 / 25 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 24 IC50 at 1 mM ATP (nM) : 93

BTK[C481S]

Product code 08-547

Full-length human BTK [2-659(end) amino acids and C481S of accession number NP_000052] was expressed as N-terminal GST-fusion protein (103 kDa) using baculovirus expression system. GST-BTK[C481S] was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 27 / 25 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 37 IC50 at 1 mM ATP (nM) : 170

BUB1/BUB3

Product code 05-187

Full-length human BUB1 [1-1085 (end) amino acids of accession number NP_004327] was co-expressed as N-terminal GST-fusion protein (149 kDa) with DYKDDDDK tagged BUB3 [1-328 (end) amino acids of accession number NP_004716] using baculovirus expression system. GST-BUB1 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : H2A peptide

ATP (μ M) Km app / Bin : 2.9 / 5 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 300 IC50 at 1 mM ATP (nM) : n.a.

CaMK1α(CAMK1)

Product code 02-104

Full-length human CaMK1 α [1-370(end) amino acids of accession number NP_003647.1] was expressed as N-terminal GST-fusion protein (68 kDa) using baculovirus expression system. GST-CaMK1 α was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : GS peptide

ATP (µM) Km app / Bin : 750 / 1000

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 16 IC50 at 1 mM ATP (nM) : 16

CaMK1δ(CAMK1D)

Product code 02-106

Full-length human CaMK1 δ [1-357(end) amino acid of accession number NP_065130.1] was expressed as N-terminal GST-fusion protein (67 kDa) using baculovirus expression system. GST-CaMK1 δ was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay
Substrate : Synapsin peptide

ATP (μ M) Km app / Bin : 11 / 10 Metal : Mq

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 2.5 IC50 at 1 mM ATP (nM) : n.a.



CaMK2α(CAMK2A)

Product code 02-109

Full-length human CaMK2 α [1-478(end) amino acids of accession number NP_741960.1] was expressed as N-terminal GST-fusion protein (81 kDa) using baculovirus expression system. GST-CaMK2 α was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : GS peptide

ATP (μ M) Km app / Bin : 33 / 50 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.75 IC50 at 1 mM ATP (nM) : n.a.

CaMK2β(CAMK2B)

Product code 02-110

Full-length human CaMK2 β [1-503 amino acids of accession number NP_742078.1] was expressed as N-terminal GST-fusion protein (83 kDa) using baculovirus expression system. GST-CaMK2 β was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : GS peptide

ATP (μ M) Km app / Bin : 19 / 25 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.54 IC50 at 1 mM ATP (nM) : n.a.

CaMK2y(CAMK2G)

Product code 02-112

Full-length human CaMK2γ[1-518(end) amino acids of accession number NP_751910.1] was expressed as N-terminal GST-fusion protein (85 kDa) using baculovirus expression system. GST-CaMK2γ was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : GS peptide

ATP (μ M) Km app / Bin : 23 / 25 Metal : Ma

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.39 IC50 at 1 mM ATP (nM) : n.a.

CaMK2δ(CAMK2D)

Product code 02-111

Full-length human CaMK2δ [1-478 amino acids of accession number NP_742113.1] was expressed as N-terminal GST-fusion protein (81 kDa) using baculovirus expression system. GST-CaMK2δ was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : GS peptide

ATP (μ M) Km app / Bin : 6.3 / 5 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.26 IC50 at 1 mM ATP (nM) : n.a.

CaMK4

Product code 02-108

Full-length human CaMK4 [1-473(end) amino acids of accession number NP_001735.1] was expressed as N-terminal GST-fusion protein (79 kDa) using baculovirus expression system. GST-CaMK4 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : GS peptide

ATP (μ M) Km app / Bin : 20 / 25 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 140 IC50 at 1 mM ATP (nM) : 1000



CDC2/CycB1

Product code 04-102

Full-length human CDC2 [1-297(end) amino acids of accession number NP_001777.1] was co-expressed as N-terminal GST-fusion protein (61 kDa) with CyclinB1 [1-433(end) amino acids of accession number NP_114172.1] using baculovirus expression system. GST-CDC2 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Modified Histone H1

ATP (μ M) Km app / Bin : 34 / 50 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 3.3 IC50 at 1 mM ATP (nM) : 32

CDC7/ASK

Product code 05-109

Full-length human CDC7 [1-574(end) amino acids of accession number NP_003494.1] was co-expressed as N-terminal GST-fusion protein (92 kDa) with Dbf4(ASK) [1-674(end) amino acids of accession number NP_006707.1] using baculovirus expression system. GST-CDC7 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : MCM2 peptide

ATP (μ M) Km app / Bin : 2.8 / 5 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 16 IC50 at 1 mM ATP (nM) : 1600

CDK2/CycA2

Product code 04-103

Full-length human CDK2 [1-298(end) amino acids of accession number NP_001789.2] was co-expressed as N-terminal GST-tagged protein (61 kDa) with GST-CyclinA2 [1-432(end) amino acids of accession number NP_001228.1] using baculovirus expression system. GST-CDK2 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay
Substrate : Modified Histone H1

ATP (μ M) Km app / Bin : 27 / 25 Metal : Ma

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 1.0 IC50 at 1 mM ATP (nM) : 7.1

CDK2/CycE1

Product code 04-165

Full-length human CDK2 [1-298(end) amino acids of accession number NP_001789.2] was co-expressed as N-terminal GST-tagged protein (61 kDa) with CyclinE1 [1-410(end) amino acids of accession number NP_001229.1] using baculovirus expression system. GST-CDK2 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay
Substrate : Modified Histone H1

ATP (µM) Km app / Bin : 130 / 150

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 2.8 IC50 at 1 mM ATP (nM) : 10

CDK3/CycE1

Product code 04-104

Full-length human CDK3 [1-305(end) amino acids of accession number NP_001249.1] was co-expressed as N-terminal GST-fusion protein (62kDa) with CyclinE1 [1-410(end) amino acids of accession number NP_001229.1] using baculovirus expression system. GST-CDK3 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay
Substrate : Modified Histone H1

ATP (μM) Km app / Bin : 1000 / 1000

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 3.4 IC50 at 1 mM ATP (nM) : 3.4



CDK4/CycD3

Product code 04-105

Full-length human CDK4 [1-303(end) amino acids of accession number NP_000066.1] was co-expressed as N-terminal GST-fusion protein (61 kDa) with human GST-CyclinD3 [1-292(end) amino acids of accession number AAA51927.1] using baculovirus expression system. GST-CDK4/CycD3 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : DYRKtide-F

ATP (µM) Km app / Bin : 200 / 200

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 13 IC50 at 1 mM ATP (nM) : 52

CDK5/p25

Product code 04-106

Full-length human CDK5 [1-292(end) amino acids of accession number NP_004926.1] was co-expressed as N-terminal GST-fusion protein (60 kDa) with p25 [99-307(end) amino acids of accession number NP_003876.1] using baculovirus expression system. GST-CDK5 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay
Substrate : Modified Histone H1

: Mg

ATP (μ M) Km app / Bin : 10 / 10

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 2.5 IC50 at 1 mM ATP (nM) : 86

Metal

CDK6/CycD3

Product code 04-107

Full-length human CDK6 [1-326(end) amino acids of accession number NP_001250.1] was co-expressed as N-terminal GST-fusion protein (64 kDa) with human GST-CyclinD3 [1-292(end) amino acids of accession number AAA51927.1] using baculovirus expression system. GST-CDK6/CycD3 was purified by using glutathione sepharose chromatography and activated with Histagged CDK7. Activated GST-CDK6/CycD3 was purified using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : DYRKtide-F

ATP (μ M) Km app / Bin : 330 / 300

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 58 IC50 at 1 mM ATP (nM) : 110

CDK7/CycH/MAT1

Product code 04-108

Full-length human CDK7 [1-346(end) amino acids of accession number NP_001790.1] was co-expressed as N-terminal GST-fusion protein (66 kDa) with CyclinH [1-323(end) amino acids of accession number NP_001230.1] and MAT1 [1-309(end) amino acids of accession number NP_002422.1] using baculovirus expression system. GST-CDK7 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CTD3 peptide

ATP (μ M) Km app / Bin : 32 / 50 Metal : Mg

Metal : Mg
Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 17 IC50 at 1 mM ATP (nM) : 120

CDK9/CycT1

Product code 04-110

Full-length human CDK9 [1-372(end) amino acids of accession number NP_001252.1] was co-expressed as N-terminal GST-fusion protein (70 kDa) with His-CyclinT1 [1-726(end) amino acids of accession number NP_001231.2] using baculovirus expression system. GST-CDK9 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay
Substrate : CDK9 substrate

ATP (µM) Km app / Bin : 9.4 / 10

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 5.2 IC50 at 1 mM ATP (nM) : 130



CGK2(PRKG2)

Product code 01-143

Full-length human CGK2 [1-762(end) amino acids of accession number NP_006250.1] was expressed as N-terminal GST-fusion protein (114 kDa) using baculovirus expression system. GST-CGK2 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Kemptide

ATP (μ M) Km app / Bin : 24 / 25 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.88 IC50 at 1 mM ATP (nM) : n.a.

CHK1(CHEK1)

Product code 02-117

Full-length human CHK1 [1-476(end) amino acids of accession number NP_001265.1] was expressed as N-terminal GST-fusion protein (81 kDa) using baculovirus expression system. GST-CHK1 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CHKtide

ATP (μ M) Km app / Bin : 50 / 50 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.24 IC50 at 1 mM ATP (nM) : 1.1

CHK2(CHEK2)

Product code 02-162

Full-length human CHK2 [1-543(end) amino acids of accession number NP_009125.1] was expressed as N-terminal GST-fusion protein (88 kDa) using baculovirus expression system. GST-CHK2 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CHKtide

ATP (μ M) Km app / Bin : 51 / 50 Metal : Ma

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 11 IC50 at 1 mM ATP (nM) : 25

CK1α(CSNK1A1)

Product code 03-101

Full-length human CK1 α [1-337(end) amino acids of accession number NP_001883.4] was expressed as N-terminal GST-fusion protein (66 kDa) using baculovirus expression system. GST-CK1 α was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CKtide

ATP (μ M) Km app / Bin : 4.1 / 5 Metal : Mg

Reference compound : 5-lodotubercidin

IC50 at ATP Bin (nM) : 150 IC50 at 1 mM ATP (nM) : >10000

CK1γ1(CSNK1G1)

Product code 03-105

Full-length human CK1γ1 [1-422(end) amino acids of accession number NP_071331.2] was expressed as N-terminal GST-fusion protein (76 kDa) using baculovirus expression system. GST-CK1γ1 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CKtide

ATP (μ M) Km app / Bin : 6.3 / 5 Metal : Mg

Reference compound : 5-lodotubercidin

IC50 at ATP Bin (nM) : 1300 IC50 at 1 mM ATP (nM) : n.a.



CK1y2(CSNK1G2)

Product code 03-106

Full-length human CK1γ2 [1-415(end) amino acids of accession number NP_001310.3] was expressed as N-terminal GST-fusion protein (75 kDa) using baculovirus expression system. GST-CK1γ2 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CKtide

ATP (μ M) Km app / Bin : 10 / 10

Metal : Mg

Reference compound : 5-lodotubercidin

IC50 at ATP Bin (nM) : 510 IC50 at 1 mM ATP (nM) : n.a.

CK1y3(CSNK1G3)

Product code 03-107

Full-length human CK1γ3 [1-447(end) amino acids of accession number NP_004375.2] was expressed as N-terminal GST-fusion protein (78 kDa) using baculovirus expression system. GST-CK1γ3 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CKtide

ATP (μ M) Km app / Bin : 3.2 / 5

Metal : Mg

Reference compound : 5-lodotubercidin

IC50 at ATP Bin (nM) : 920 IC50 at 1 mM ATP (nM) : n.a.

CK1δ(CSNK1D)

Product code 03-103

Human CK1δ, catalytic domain [1-294 amino acids of accession number NP_001884.2] was expressed as N-terminal GST-fusion protein (61 kDa) using E. coli expression system. GST-CK1δ was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CKtide

ATP (μM) Km app / Bin : 7.7 / 10

Metal : Mg

Reference compound : 5-lodotubercidin

IC50 at ATP Bin (nM) : 25 IC50 at 1 mM ATP (nM) : n.a.

CK1ε(CSNK1E)

Product code 03-104

Human CK1 ϵ , catalytic domain [1-348 amino acids of accession number NP_001885.1] was expressed as N-terminal GST-fusion protein (68 kDa) using baculovirus expression system. GST-CK1 ϵ was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CKtide

ATP (μ M) Km app / Bin : 16 / 25 Metal : Mg

Reference compound : 5-lodotubercidin

IC50 at ATP Bin (nM) : 300 IC50 at 1 mM ATP (nM) : 5800

$CK2\alpha1/\beta(CSNK2A1/B)$

Product code 05-184

Full-length human CK2 α 1 [1-391(end) amino acids of accession number NP_001886.1] was co-expressed as N-terminal GST-fusion protein (72 kDa) with human His-tagged CK2 β [1-215 amino acids of accession number NP_001311.3] using baculovirus expression system. GST-CK2 α 1 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CK2tide

ATP (μ M) Km app / Bin : 2.9 / 5 Metal : Mg

Reference compound : TBB IC50 at ATP Bin (nM) : 60 IC50 at 1 mM ATP (nM) : 4800



CK2α2/β(CSNK2A2/B)

Product code 05-185

Full-length human CK2 α 2 [1-350(end) amino acids of accession number NP_001887.1] was co-expressed as N-terminal GST-fusion protein (68 kDa) with human His-tagged CK2 β [1-215 amino acids of accession number NP_001311.3] using baculovirus expression system. GST-CK2 α 2 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CK2tide

ATP (μ M) Km app / Bin : 2.1 / 5

Metal : Mg

Reference compound : TBB IC50 at ATP Bin (nM) : 50

IC50 at 1 mM ATP (nM): n.a.

CLK₁

Product code 04-126

Human CLK1, catalytic domain [129-484(end) amino acids of accession number NP_004062.2] was expressed as N-terminal GST-fusion protein (69 kDa) using baculovirus expression system. GST-CLK1 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : DYRKtide-F

ATP (μM) Km app / Bin : 11 / 10

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 9.6 IC50 at 1 mM ATP (nM) : 60

CLK₂

Product code 04-127

Full-length human CLK2 [1-499(end) amino acids of accession number AAH53603.1] was expressed as N-terminal GST-fusion protein (87 kDa) using baculovirus expression system. GST-CLK2 was purified by using glutathione sepharose chromatography. Assay platform : Mobility Shift Assay

Substrate : DYRKtide-F

ATP (μ M) Km app / Bin : 140 / 150

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 4.6 IC50 at 1 mM ATP (nM) : 28

CLK3

Product code 04-128

Full-length human CLK3 [1-490(end) amino acids of accession number AAH02555.1] was expressed as N-terminal GST-fusion protein (86 kDa) using baculovirus expression system. GST-CLK3 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : DYRKtide-F

ATP (μM) Km app / Bin : 75 / 75

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 820 IC50 at 1 mM ATP (nM) : n.a.

CRIK(CIT)

Product code 01-104

Human citron kinase (CRIK), catalytic domain [1-449 amino acids of accession number NP_009105.1] was expressed as N-terminal GST fusion protein (77 kDa) using baculovirus expression system. GST-CRIK was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay
Substrate : Histone H3 peptide

ATP (μ M) Km app / Bin : 7.8 / 10 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 31 IC50 at 1 mM ATP (nM) : n.a.



CSK

Product code 08-111

Full-length human CSK [1-450(end) amino acids of accession number NP_004374.1] was expressed as N-terminal GST-fusion protein (78 kDa) using baculovirus expression system. GST-CSK was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 4.8 / 5 Metal : Mg+Mn

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 53 IC50 at 1 mM ATP (nM) : 1500

DAPK1

Product code 02-134

Human DAPK1, catalytic domain [1-289 amino acids of accession number NP_004929.1] was expressed as N-terminal GST-fusion protein (60 kDa) using baculovirus expression system. GST-DAPK1 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : DAPK1tide

ATP (μ M) Km app / Bin : 1.1 / 1 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 2.9 IC50 at 1 mM ATP (nM) : 490

DCAMKL2

Product code 02-140

Truncated human DCAMKL2 [1-691 amino acids and Q353 deletion of accession number NP_001035350.2] was expressed as N-terminal GST-fusion protein (103 kDa) using baculovirus expression system. GST-DCAMKL2 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : GS peptide

ATP (μ M) Km app / Bin : 120 / 150

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 22 IC50 at 1 mM ATP (nM) : n.a.

DDR1

Product code 08-113

Human DDR1, cytoplasmic domain [444-876(end) amino acids of accession number NP_001945.3] was expressed as N-terminal GST-fusion protein (75 kDa) using baculovirus expression system. GST-DDR1 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : IRS1

ATP (µM) Km app / Bin : 94 / 100

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 4.0 IC50 at 1 mM ATP (nM) : 3.1

DDR2

Product code 08-114

Human DDR2, cytoplasmic domain [422-855(end) amino acids of accession number NP_006173.2] was expressed as N-terminal GST-fusion protein (77 kDa) using baculovirus expression system. GST-DDR2 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : IRS1

ATP (μ M) Km app / Bin : 38 / 50 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 2.2 IC50 at 1 mM ATP (nM) : 0.77



DYRK1A

Product code 04-130

Full-length human DYRK1A [1-763(end) amino acids of accession number NP_001387.2] was expressed as N-terminal GST-fusion protein (112 kDa) using baculovirus expression system. GST-DYRK1A was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : DYRKtide-F

ATP (μ M) Km app / Bin : 16 / 25 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 7.8 IC50 at 1 mM ATP (nM) : 120

DYRK1B

Product code 04-131

Full-length human DYRK1B [1-629(end) amino acids of accession number NP_004705.1] was expressed as N-terminal GST-fusion protein (96 kDa) using baculovirus expression system. GST-DYRK1B was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : DYRKtide-F

ATP (μ M) Km app / Bin : 59 / 50 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 2.2 IC50 at 1 mM ATP (nM) : 32

DYRK2

Product code 04-132

Full-length human DYRK2 [1-528(end) amino acids of accession number NP_003574.1] was expressed as N-terminal GST-fusion protein (87 kDa) using baculovirus expression system. GST-DYRK2 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : DYRKtide-F

ATP (µM) Km app / Bin : 7.7 / 10

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 130 IC50 at 1 mM ATP (nM) : n.a.

DYRK3

Product code 04-133

Full-length human DYRK3 [1-588(end) amino acids of accession number NP_003573.2] was expressed as N-terminal GST-fusion protein (93 kDa) using baculovirus expression system. GST-DYRK3 was purified by using glutathione sepharose chromatography and anion exchange chromatography.

Assay platform : Mobility Shift Assay

Substrate : DYRKtide-F

ATP (μ M) Km app / Bin : 6.8 / 5 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 17 IC50 at 1 mM ATP (nM) : n.a.

EEF2K

Product code 10-113

Full-length human EEF2K [1-725(end) amino acids of accession number NP_037434.1] was expressed as N-terminal GST-fusion protein (109 kDa) using E. coli expression system. GST-EEF2K was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : EEF2Ktide

ATP (µM) Km app / Bin : 12 / 10

Metal : Mg Reference compound : A-484954

IC50 at ATP Bin (nM) : 330 IC50 at 1 mM ATP (nM) : n.a.



EGFR

Product code 08-115

Human EGFR, cytoplasmic domain [669-1210(end) amino acids of accession number NP_005219.2] was expressed as N-terminal GST-fusion protein (89 kDa) using baculovirus expression system. GST-EGFR was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 53 IC50 at 1 mM ATP (nM) : 7700

EGFR[C797S/L858R]

Product code 08-563

Human EGFR, cytoplasmic domain [669-1210(end) amino acids and C797S/L858R of accession number NP_005219.2] was expressed as N-terminal GST-fusion protein (89kDa) using baculovirus expression system. GST-EGFR[C797S/L858R] was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 4.1 / 5 Metal : Mg+Mn

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 8.8 IC50 at 1 mM ATP (nM) : 270

EGFR[d746-750]

Product code 08-527

Human EGFR, cytoplasmic domain [669-745, 751-1210(end) amino acids of accession number NP_005219.2] was expressed as N-terminal GST-fusion protein (88 kDa) using baculovirus expression system. GST-EGFR[d746-750aa] was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 19 / 25 Metal : Mg+Mn

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 13 IC50 at 1 mM ATP (nM) : 93

EGFR[d746-750/C797S]

Product code 08-564

Human EGFR, cytoplasmic domain [669-745, 751-1210(end) amino acids and C797S of accession number NP_005219.2] was expressed as N-terminal GST-fusion protein (88 kDa) using baculovirus expression system. GST-EGFR[d746-750aa/C797S] was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μM) Km app / Bin : 8.2 / 10 Metal : Mg+Mn

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 8.0 IC50 at 1 mM ATP (nM) : 130

EGFR[d746-750/T790M]

Product code 08-528

Human EGFR, cytoplasmic domain [669-745, 751-1210(end) amino acids and T790M of accession number NP_005219.2] was expressed as N-terminal GST-fusion protein (89 kDa) using baculovirus expression system. GST-EGFR [d746-750aa/T790M] was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 5.4 / 5 Metal : Mg+Mn Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.52

IC50 at 1 mM ATP (nM): 9.7

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EGFR[d746-750/T790M/C797S]

Product code 08-565

Human EGFR, cytoplasmic domain [669-745, 751-1210(end) amino acids and T790M/C797S of accession number NP_005219.2] was expressed as N-terminal GST-fusion protein (88 kDa) using baculovirus expression system. GST-EGFR[d746-750aa/T790M/C797S] was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 1.8 / 2 Metal : Mg+Mn

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.84 IC50 at 1 mM ATP (nM) : 14

EGFR[D770_N771insNPG]

Product code 08-553

Human EGFR, cytoplasmic domain [669-1210(end) amino acids and D770_N771insNPG of accession number NP_005219.2] was expressed as N-terminal GST-fusion protein (89 kDa) using baculovirus expression system. GST-EGFR[D770_N771insNPG] was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 2.3 / 5 Metal : Mg+Mn

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 18 IC50 at 1 mM ATP (nM) : 930

EGFR[L858R]

Product code 08-502

Human EGFR, cytoplasmic domain [669-1210(end) amino acids and L858R of accession number NP_005219.2] was expressed as N-terminal GST-fusion protein (89 kDa) using baculovirus expression system. GST-EGFR[L858R] was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 9.8 / 10 Metal : Mg+Mn

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 11 IC50 at 1 mM ATP (nM) : 360

EGFR[L861Q]

Product code 08-513

Human EGFR, cytoplasmic domain [669-1210(end) amino acids and L861Q of accession number NP_005219.2] was expressed as N-terminal GST-fusion protein (89 kDa) using baculovirus expression system. GST-EGFR[L861Q] was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 7.5 / 10 Metal : Mg+Mn

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 68 IC50 at 1 mM ATP (nM) : 2200

EGFR[T790M]

Product code 08-194

Human EGFR, cytoplasmic domain [669-1210(end) amino acids and T790M of accession number NP_005219.2] was expressed as N-terminal GST-fusion protein (89 kDa) using baculovirus expression system. GST-EGFR[T790M] was purified by using glutathione sepharose chromatography and anion exchange chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 0.9 / 1 Metal : Mg+Mn Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 1.8 IC50 at 1 mM ATP (nM) : 190



EGFR[T790M/C797S/L858R]

Product code 08-559

Human EGFR, cytoplasmic domain [669-1210(end) amino acids and T790M/C797S/L858R of accession number NP_005219.2] was expressed as N-terminal GST-fusion protein (89 kDa) using baculovirus expression system. GST-EGFR[T790M/C797S/L858R] was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 0.85 / n.a. Metal : Mg+Mn

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : n.a. IC50 at 1 mM ATP (nM) : 37

EGFR[T790M/L858R]

Product code 08-510

Human EGFR, cytoplasmic domain [669-1210(end) amino acids and T790M/L858R of accession number NP_005219.2] was expressed as N-terminal GST-fusion protein (89 kDa) using baculovirus expression system. GST-EGFR[T790M/L858R] was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 1.9 / 2 Metal : Mg+Mn

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 1.0 IC50 at 1 mM ATP (nM) : 56

EPHA1

Product code 08-119

Human EPHA1, cytoplasmic domain [586-976(end) amino acids of accession number NP_005223.3] was expressed as N-terminal GST-fusion protein (71 kDa) using baculovirus expression system. GST-EPHA1 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Blk/Lyntide

ATP (μ M) Km app / Bin : 22 / 25 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 20 IC50 at 1 mM ATP (nM) : 340

EPHA2

Product code 08-121

Human EPHA2, cytoplasmic domain [572-976(end) amino acids of accession number NP_004422.2] was expressed as N-terminal GST-fusion protein (73 kDa) using baculovirus expression system. GST-EPHA2 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Blk/Lyntide

ATP (μ M) Km app / Bin : 67 / 75 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 160 IC50 at 1 mM ATP (nM) : 530

EPHA3

Product code 08-122

Human EPHA3, cytoplasmic domain [579-983(end) amino acids of accession number NP_005224.2] was expressed as N-terminal GST-fusion protein (72 kDa) using baculovirus expression system. GST-EPHA3 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Blk/Lyntide

ATP (µM) Km app / Bin : 170 / 150

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 37 IC50 at 1 mM ATP (nM) : 76



EPHA4

Product code 08-123

Human EPHA4, cytoplasmic domain [586-986(end) amino acids of accession number NP_004429.1] was expressed as N-terminal GST-fusion protein (72 kDa) using baculovirus expression system. GST-EPHA4 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Blk/Lyntide

ATP (μ M) Km app / Bin : 52 / 50 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 50 IC50 at 1 mM ATP (nM) : 330

EPHA5

Product code 08-124

Human EPHA5, catalytic domain [662-948 amino acids of accession number NP_004430.3] was expressed as N-terminal GST-fusion protein (59 kDa) using baculovirus expression system. GST-EPHA5 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Blk/Lyntide

ATP (μ M) Km app / Bin : 56 / 50 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 34 IC50 at 1 mM ATP (nM) : 220

EPHA6

Product code 08-125

Human EPHA6, cytoplasmic domain [683-1130(end) amino acids of accession number NP_001073917.2] was expressed as N-terminal GST-fusion protein (77 kDa) using baculovirus expression system. GST-EPHA6 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Blk/Lyntide

ATP (μM) Km app / Bin : 27 / 25 Metal : Mα

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 17 IC50 at 1 mM ATP (nM) : 60

EPHA7

Product code 08-126

Human EPHA7, cytoplasmic domain [595-998(end) amino acids of accession number NP_004431.1] was expressed as N-terminal GST-fusion protein (73 kDa) using baculovirus expression system. GST-EPHA7 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Blk/Lyntide

ATP (μ M) Km app / Bin : 58 / 50 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 48 IC50 at 1 mM ATP (nM) : 480

EPHA8

Product code 08-127

Human EPHA8, catalytic domain [571-924 amino acids of accession number NP_065387.1] was expressed as N-terminal GST-fusion protein (67 kDa) using baculovirus expression system. GST-EPHA8 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Blk/Lyntide

ATP (μ M) Km app / Bin : 69 / 75 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 61 IC50 at 1 mM ATP (nM) : 240



EPHB1

Product code 08-128

Human EPHB1, cytoplasmic domain [578-984(end) amino acids of accession number NP_004432.1] was expressed as N-terminal GST-fusion protein (73 kDa) using baculovirus expression system. GST-EPHB1 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Blk/Lyntide

ATP (μM) Km app / Bin : 29 / 25 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 53 IC50 at 1 mM ATP (nM) : 760

EPHB2

Product code 08-129

Human EPHB2, cytoplasmic domain [581-987(end) amino acids of accession number NP_004433.2] was expressed as N-terminal GST-fusion protein (73 kDa) using baculovirus expression system. GST-EPHB2 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Blk/Lyntide

ATP (μ M) Km app / Bin : 86 / 100

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 73 IC50 at 1 mM ATP (nM) : 400

EPHB3

Product code 08-130

Human EPHB3, cytoplasmic domain [596-998(end) amino acids of accession number NP_004434.2] was expressed as N-terminal GST-fusion protein (73 kDa) using baculovirus expression system. GST-EPHB3 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Blk/Lyntide

ATP (μ M) Km app / Bin : 49 / 50 Metal : Mg

Metal : Mg
Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 2000 IC50 at 1 mM ATP (nM) : >10000

EPHB4

Product code 08-131

Human EPHB4, cytoplasmic domain [577-987(end) amino acids of accession number NP_004435.3] was expressed as N-terminal GST-protein (73 kDa) using baculovirus expression system. GST-EPHB4 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Blk/Lyntide

ATP (μ M) Km app / Bin : 56 / 50 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 230 IC50 at 1 mM ATP (nM) : 1500

Erk1(MAPK3)

Product code 04-142

Full-length human Erk1 [1-379(end) amino acids of accession number NP_002737.1] was expressed as N-terminal GST-fusion protein (70 kDa) using E.coli expression system. GST-Erk1 was purified by using glutathione sepharose chromatography and activated with His-tagged MAP2K1. Activated GST-Erk1 was purified using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay
Substrate : Modified Erktide

ATP (μM) Km app / Bin : 34 / 50

Metal : Mg

Reference compound : K252a

IC50 at ATP Bin (nM) : 37

IC50 at 1 mM ATP (nM) : 400

23



Erk2(MAPK1)

Product code 04-143

Full-length human Erk2 [1-360(end) amino acids of accession number NP_002736.3] was expressed as N-terminal GST-fusion protein (69 kDa) using E.coli expression system. GST-Erk2 was purified by using glutathione sepharose chromatography and activated with His-tagged MAP2K1. Activated GST-Erk2 was purified using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Modified Erktide

ATP (μM) Km app / Bin : 33 / 50

Metal : Mg

Reference compound : K252a IC50 at ATP Bin (nM) : 21

IC50 at 1 mM ATP (nM): 180

Erk5(MAPK7)

Product code 04-146

Human Erk5, catalytic domain [1-398 amino acids of accession number NP_002740.2] was expressed as N-terminal GST-fusion protein (72 kDa) using E. coli expression system. GST-Erk5 was purified by using glutathione sepharose chromatography and activated with His-tagged MAP2K5. Activated GST-Erk5 was purified using Ni-NTA affinity chromatography.

Assay platform : Mobility Shift Assay
Substrate : EGFR-derived peptide

ATP (µM) Km app / Bin : 450 / 1000

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 280 IC50 at 1 mM ATP (nM) : 280

FAK(PTK2)

Product code 08-137

Truncated human FAK[376-1052(end) amino acids of accession number NP_722560.1] was expressed as N-terminal GST-fusion protein (103 kDa) using baculovirus expression system. GST-FAK was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Blk/Lyntide

ATP (μ M) Km app / Bin : 25 / 25 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 47 IC50 at 1 mM ATP (nM) : 230

FER

Product code 08-139

Full-length human FER [1-822(end) amino acids of accession number NP_005237.1] was expressed as N-terminal GST-fusion protein (122 kDa) using baculovirus expression system. GST-FER was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 26 / 25 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 1.0 IC50 at 1 mM ATP (nM) : 12

FES

Product code 08-140

Full-length human FES [1-413, 416-822(end) amino acids of accession number NP_001996.1] was expressed as N-terminal GST-fusion protein (120 kDa) using baculovirus expression system. GST-FES was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μM) Km app / Bin : 43 / 50

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 2.9 IC50 at 1 mM ATP (nM) : 25



FGFR1

Product code 08-133

Human FGFR1, cytoplasmic domain [398-822(end) amino acids of accession number NP_075598.2] was expressed as N-terminal GST-fusion protein (75 kDa) using baculovirus expression system. GST-FGFR1 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CSKtide

ATP (μM) Km app / Bin : 89 / 100

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 2.3 IC50 at 1 mM ATP (nM) : 12

FGFR1[V561M]

Product code 08-536

Human FGFR1, cytoplasmic domain [398-822(end) amino acids and V561M of accession number NP_075598.2] was expressed as N-terminal GST-fusion protein (75 kDa) using baculovirus expression system. GST-FGFR1[V561M] was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CSKtide

ATP (μ M) Km app / Bin : 33 / 50 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.14 IC50 at 1 mM ATP (nM) : 1.3

FGFR2

Product code 08-134

Human FGFR2, cytoplasmic domain [399-821(end) amino acids of accession number NP_000132.1] was expressed as N-terminal GST-fusion protein (75 kDa) using baculovirus expression system. GST-FGFR2 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CSKtide

ATP (μ M) Km app / Bin : 66 / 75 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 1.4 IC50 at 1 mM ATP (nM) : 5.4

FGFR2[V564I]

Product code 08-546

Human FGFR2, cytoplasmic domain [399-821(end) amino acids and V564I of accession number NP_000132] was expressed as N-terminal GST-fusion protein (75 kDa) using baculovirus expression system. GST-FGFR2[V564I] was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CSKtide

ATP (μ M) Km app / Bin : 21 / 25 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 2.0 IC50 at 1 mM ATP (nM) : 47

FGFR3

Product code 08-135

Human FGFR3, cytoplasmic domain [436-806(end) amino acids of accession number NP_000133.1] was expressed as N-terminal GST-fusion protein (68 kDa) using baculovirus expression system. GST-FGFR3 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CSKtide

ATP (μ M) Km app / Bin : 43 / 50 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 2.6 IC50 at 1 mM ATP (nM) : 15



FGFR3[K650E]

Product code 08-501

Human FGFR3, cytoplasmic domain [436-806(end) amino acids and K650E of accession number NP_000133.1] was expressed as N-terminal GST-fusion protein (68 kDa) using baculovirus expression system. GST-FGFR3[K650E] was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CSKtide

ATP (μ M) Km app / Bin : 41 / 50 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 1.2 IC50 at 1 mM ATP (nM) : 14

FGFR3[K650M]

Product code 08-199

Human FGFR3, cytoplasmic domain [436-806(end) amino acids and K650M of accession number NP_000133.1] was expressed as N-terminal GST-fusion protein (68 kDa) using baculovirus expression system. GST-FGFR3[K650M] was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CSKtide

ATP (μ M) Km app / Bin : 17 / 25 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.68 IC50 at 1 mM ATP (nM) : 17

FGFR3[V555L]

Product code 08-548

Human FGFR3, cytoplasmic domain [436-806(end) amino acids and V555L of accession number NP_000133.1] was expressed as N-terminal GST-fusion protein (68 kDa) using baculovirus expression system. GST-FGFR3[V555L] was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CSKtide

ATP (μ M) Km app / Bin : 29 / 25 Metal : Ma

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.49 IC50 at 1 mM ATP (nM) : 9.4

FGFR3[V555M]

Product code 08-543

Human FGFR3, cytoplasmic domain [436-806(end) amino acids and V555M of accession number NP_000133.1] was expressed as N-terminal GST-fusion protein (68 kDa) using baculovirus expression system. GST-FGFR3[V555M] was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CSKtide

ATP (μ M) Km app / Bin : 37 / 50 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.21 IC50 at 1 mM ATP (nM) : 1.8

FGFR4

Product code 08-136

Human FGFR4, cytoplasmic domain [460-802(end) amino acids of accession number NP_002002.3] was expressed as N-terminal GST-fusion protein (65 kDa) using baculovirus expression system. GST-FGFR4 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CSKtide

ATP (µM) Km app / Bin : 230 / 250

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 43 IC50 at 1 mM ATP (nM) : 120



FGFR4[N535K]

Product code 08-524

Human FGFR4, cytoplasmic domain [460-802(end) amino acids and N535K of accession number NP_002002.3] was expressed as N-terminal GST-fusion protein (65 kDa) using baculovirus expression system. GST-FGFR4[N535K] was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CSKtide

ATP (μ M) Km app / Bin : 30 / 25 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 160 IC50 at 1 mM ATP (nM) : 1200

FGFR4[V550E]

Product code 08-525

Human FGFR4, cytoplasmic domain [460-802(end) amino acids and V550E of accession number NP_002002.3] was expressed as N-terminal GST-fusion protein (65 kDa) using baculovirus expression system. GST-FGFR4[V550E] was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CSKtide

ATP (μM) Km app / Bin : 210 / 200

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 370 IC50 at 1 mM ATP (nM) : 1300

FGFR4[V550L]

Product code 08-526

Human FGFR4, cytoplasmic domain [460-802(end) amino acids and V550L of accession number NP_002002.3] was expressed as N-terminal GST-fusion protein (65 kDa) using baculovirus expression system. GST-FGFR4[V550L] was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CSKtide

ATP (µM) Km app / Bin : 160 / 150

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 10 IC50 at 1 mM ATP (nM) : 44

FGR

Product code 08-166

Full-length human FGR [1-529(end) amino acids of accession number NP_005239.1] was expressed as N-terminal GST-fusion protein (86 kDa) using baculovirus expression system. GST-FGR was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 34 / 50 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 1.3 IC50 at 1 mM ATP (nM) : 16

FLT1

Product code 08-189

Human FLT1, cytoplasmic domain [781-1338(end) amino acids of accession number NP_002010.1] was expressed as N-terminal GST-fusion protein (90 kDa) using baculovirus expression system. GST-FLT1 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CSKtide

ATP (μ M) Km app / Bin : 140 / 150

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 3.2 IC50 at 1 mM ATP (nM) : 6.8



FLT3

Product code 08-154

Human FLT3, cytoplasmic domain [564-993(end) amino acids of accession number NP_004110.2] was expressed as N-terminal GST-fusion protein (77 kDa) using baculovirus expression system. GST-FLT3 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μM) Km app / Bin : 94 / 100

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.20 IC50 at 1 mM ATP (nM) : 0.34

FLT4

Product code 08-190

Human FLT4, cytoplasmic domain [798-1298(end) amino acids of accession number NP_002011.1] was expressed as N-terminal GST-fusion protein (83 kDa) using baculovirus expression system. GST-FLT4 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CSKtide

ATP (μ M) Km app / Bin : 72 / 75 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.66 IC50 at 1 mM ATP (nM) : 2.4

FMS(CSF1R)

Product code 08-155

Human FMS, cytoplasmic domain [538-972(end) amino acids of accession number NP_005202.2] was expressed as N-terminal GST-fusion protein (76 kDa) using baculovirus expression system. GST-FMS was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μM) Km app / Bin : 26 / 25 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.26 IC50 at 1 mM ATP (nM) : 0.70

FRK

Product code 08-167

Human FRK, catalytic domain [223-505(end) amino acids of accession number NP_002022.1] was expressed as N-terminal GST-fusion protein (60 kDa) using baculovirus expression system. GST-FRK was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 62 / 75 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 3.4 IC50 at 1 mM ATP (nM) : 40

FYN[isoform a]

Product code 08-168

Full-length human FYN [isoform a] [1-537(end) amino acids of accession number NP_002028.1] was expressed as N-terminal GST-fusion protein (88 kDa) using baculovirus expression system. GST-FYN [isoform a] was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 36 / 50 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 3.9 IC50 at 1 mM ATP (nM) : 24



FYN[isoform b]

Product code 08-531

Full-length human FYN [isoform b] [1-534(end) amino acids of accession number NP_694592.1] was expressed as N-terminal GST-fusion protein (87 kDa) using baculovirus expression system. GST-FYN [isoform b] was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 20 / 25 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 2.8 IC50 at 1 mM ATP (nM) : 42

GSK3α(GSK3A)

Product code 04-140

Full-length human GSK3α [1-483(end) amino acids of accession number NP_063937.2] was expressed as N-terminal GST-fusion protein (78 kDa) using baculovirus expression system. GST-GSK3α was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CREBtide-p

ATP (μ M) Km app / Bin : 12 / 10 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 15 IC50 at 1 mM ATP (nM) : 180

GSK3B(GSK3B)

Product code 04-141

Full-length human GSK3 β [1-420(end) amino acids of accession number NP_001139628.1] was expressed as N-terminal GST-fusion protein (74 kDa) using baculovirus expression system. GST-GSK3 β was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CREBtide-p

ATP (μM) Km app / Bin : 9.1 / 10

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 9.2 IC50 at 1 mM ATP (nM) : 240

Haspin(GSG2)

Product code 05-111

Full-length human Haspin [1-798(end) amino acids of accession number NP_114171.2] was expressed as N-terminal GST-fusion protein (116 kDa) using baculovirus expression system. GST-Haspin was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay Substrate : Histone H3 peptide

ATP (µM) Km app / Bin : 140 / 150

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 5.8 IC50 at 1 mM ATP (nM) : n.a.

HCK

Product code 08-169

Truncated human HCK [25-526(end) amino acids of accession number NP_002101.2] was expressed as N-terminal GST-fusion protein (84 kDa) using baculovirus expression system. GST-HCK was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 11 / 10 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 1.1 IC50 at 1 mM ATP (nM) : 22



HER2(ERBB2)

Product code 08-016

Human HER2, cytoplasmic domain [676-1255(end) amino acids of accession number NP_004439.1] was expressed as N-terminal Histagged protein (67 kDa) using baculovirus expression system. Histagged HER2 was purified by using Ni-NTA affinity chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 3.5 / 5 Metal : Mn

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 90 IC50 at 1 mM ATP (nM) : >10000

HER4(ERBB4)

Product code 08-118

Human HER4, cytoplasmic domain [676-1308(end) amino acids of accession number NP_005226.1] was expressed as N-terminal GST-fusion protein (99 kDa) using baculovirus expression system. GST-HER4 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 27 / 25 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 34 IC50 at 1 mM ATP (nM) : 1000

HGK(MAP4K4)

Product code 07-137

Human HGK, catalytic domain [1-328 amino acids of accession number NP_004825.2] was expressed as N-terminal GST-fusion protein (64 kDa) using baculovirus expression system. GST-HGK was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay
Substrate : Moesin-derived peptide

ATP (μ M) Km app / Bin : 9.4 / 10 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 1.0 IC50 at 1 mM ATP (nM) : 23

HIPK1

Product code 04-135

Human HIPK1, catalytic domain [158-555 amino acids of accession number NP_689909.2] was expressed as N-terminal GST-fusion protein (73 kDa) using baculovirus expression system. GST-HIPK1 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : DYRKtide-F

ATP (μ M) Km app / Bin : 4.4 / 5 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 570 IC50 at 1 mM ATP (nM) : n.a.

HIPK2

Product code 04-136

Full-length human HIPK2 [1-1198(end) amino acids of accession number NP_073577.3] was expressed as N-terminal GST-fusion protein (158 kDa) using baculovirus expression system. GST-HIPK2 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : DYRKtide-F

ATP (μ M) Km app / Bin : 5.9 / 5 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 170 IC50 at 1 mM ATP (nM) : n.a.



HIPK3

Product code 04-137

Human HIPK3, catalytic domain [161-562 amino acids of accession number NP_005725.3] was expressed as N-terminal GST-fusion protein (73 kDa) using baculovirus expression system. GST-HIPK3 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : DYRKtide-F

ATP (μ M) Km app / Bin : 7.3 / 5 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 120 IC50 at 1 mM ATP (nM) : n.a.

HIPK4

Product code 04-138

Full-length human HIPK4 [1-616(end) amino acids of accession number NP_653286.2] was expressed as N-terminal GST-fusion protein (96 kDa) using baculovirus expression system. GST-HIPK4 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : DYRKtide-F

 $\begin{array}{lll} \text{ATP (μM$) Km app / Bin} & : ~7~/~5 \\ \text{Metal} & : ~\text{Mg} \end{array}$

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 71 IC50 at 1 mM ATP (nM) : >10000

HPK1(MAP4K1)

Product code 07-410

Human HPK1, catalytic domain [1-346 amino acids of accession number NP_009112.1] was expressed as N-terminal DYKDDDDK tagged protein (41 kDa) using baculovirus expression system. The protein was purified by using DYKDDDDK tag antibody agarose.

Assay platform : Mobility Shift Assay

Substrate : S6K2 peptide

ATP (μM) Km app / Bin : 22 / 25

Metal : Mg
Reference compound : K252a
IC50 at ATP Bin (nM) : 6.9
IC50 at 1 mM ATP (nM) : n.a.

IGF1R

Product code 08-141

Human IGF1R, cytoplasmic domain [959-1367(end) amino acids of accession number NP_000866.1] was expressed as N-terminal GST-fusion protein (73 kDa) using baculovirus expression system. GST-IGF1R was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : IRS1

ATP (μ M) Km app / Bin : 63 / 75 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 40 IC50 at 1 mM ATP (nM) : 150

IKKα(CHUK)

Product code 05-112

Full-length human IKKα [1-745(end) amino acids of accession number NP_001269.3] was expressed as N-terminal GST-fusion protein (111 kDa) using baculovirus expression system. GST-IKKα was purified by using glutathione sepharose chromatography.

Assay platform : IMAP

Substrate : IκBα peptide

ATP (µM) Km app / Bin : 41 / 40

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 310 IC50 at 1 mM ATP (nM) : n.a.



IKKB(IKBKB)

Product code 05-084

Truncated human IKK β [1-662 amino acids of accession number NP_001547.1] was expressed as N-terminal His-tagged protein (77 kDa) using baculovirus expression system. His-tagged IKK β was purified by using Ni-NTA affinity chromatography and anion exchange chromatography.

Assay platform : Mobility Shift Assay

Substrate : Modified IκBα-derived

peptide

ATP (μ M) Km app / Bin : 16 / 25 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 410 IC50 at 1 mM ATP (nM) : >10000

IKKε(IKBKE)

Product code 05-114

Full-length human IKKε [1-716(end) amino acids of accession number NP_054721.1] was expressed as N-terminal GST-fusion protein (108 kDa) using baculovirus expression system. GST-IKKε was purified by using glutathione sepharose chromatography and gel filtration chromatography.

Assay platform : Mobility Shift Assay

Substrate : IκBα peptide

ATP (μ M) Km app / Bin : 9.5 / 10

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 2.7 IC50 at 1 mM ATP (nM) : n.a.

INSR

Product code 08-142

Human INSR, catalytic domain [1005-1310 amino acids of accession number NP_000199.1] was expressed as N-terminal GST-fusion protein (62 kDa) using baculovirus expression system. GST-INSR was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : IRS1

ATP (μ M) Km app / Bin : 58 / 50 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 12 IC50 at 1 mM ATP (nM) : 70

IRAK1

Product code 09-101

Truncated human IRAK1 [194-712(end) amino acids of accession number NP_001560.2] was expressed as N-terminal GST-fusion protein (83 kDa) using baculovirus expression system. GST-IRAK1 was purified by using glutathione sepharose chromatography.

Assay platform : IMAP

Substrate : SRPKtide

ATP (μM) Km app / Bin : 27 / 25

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 54 IC50 at 1 mM ATP (nM) : n.a.

IRAK4

Product code 09-145

Full-length human IRAK4 [1-460(end) amino acids of accession number NP_057207.2] was expressed as N-terminal GST-fusion protein (79 kDa) using baculovirus expression system. GST-IRAK4 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : IRAK1 peptide

ATP (µM) Km app / Bin : 920 / 1000

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 11 IC50 at 1 mM ATP (nM) : 11



IRR(INSRR)

Product code 08-143

Human IRR, cytoplasmic domain [953-1297(end) amino acids of accession number NP_055030.1] was expressed as N-terminal GST-fusion protein (66 kDa) using baculovirus expression system. GST-IRR was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : IRS1

ATP (μ M) Km app / Bin : 64 / 75 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 15 IC50 at 1 mM ATP (nM) : 98

ITK

Product code 08-181

Full-length human ITK [2-620(end) amino acids of accession number NP_005537.3] was expressed as N-terminal GST-fusion protein (99 kDa) using baculovirus expression system. GST-ITK was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 6.1 / 10

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 3.2 IC50 at 1 mM ATP (nM) : 200

JAK1

Product code 08-144

Human JAK1, catalytic domain [850-1154(end) amino acids of accession number NP_002218.2] was expressed as N-terminal GST-fusion protein (62 kDa) using baculovirus expression system. GST-JAK1 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay
Substrate : JAK1 substrate peptide

ATP (μ M) Km app / Bin : 68 / 75 Metal : Ma

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.71 IC50 at 1 mM ATP (nM) : 10

JAK2

Product code 08-045

Human JAK2, catalytic domain [826-1132(end) amino acids of accession number NP_004963.1] was expressed as N-terminal Histagged protein (39 kDa) using baculovirus expression system. Histagged JAK2 was purified by using Ni-NTA affinity chromatography and gel filtration chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 13 / 10 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.34 IC50 at 1 mM ATP (nM) : 6.0

JAK3

Product code 08-046

Human JAK3, catalytic domain [795-1124(end) amino acids of accession number NP_000206.2] was expressed as N-terminal Histagged protein (41 kDa) using baculovirus expression system. Histagged JAK3 was purified by using Ni-NTA affinity chromatography and gel filtration chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 3.5 / 5 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.20 IC50 at 1 mM ATP (nM) : 12



JNK1(MAPK8)

Product code 04-163

Human JNK1, catalytic domain [2-364 amino acids of accession number NP_620634.1] was expressed as N-terminal GST-fusion protein (69 kDa) using E. coli expression system. GST-JNK1 was purified by using glutathione sepharose chromatography and activated with His-tagged MAP2K4 and MAP2K7. Activated GST-JNK1 was purified using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Modified Erktide

ATP (μ M) Km app / Bin : 29 / 100

Metal : Mg
Reference compound : K252a
IC50 at ATP Bin (nM) : 99
IC50 at 1 mM ATP (nM) : 770

JNK2(MAPK9)

Product code 04-164

Human JNK2, catalytic domain [1-364 amino acids of accession number NP_002743.3] was expressed as N-terminal GST-fusion protein (69 kDa) using E. coli expression system. GST-JNK2 was purified by using glutathione sepharose chromatography and activated with His-tagged MAP2K4 and MAP2K7. Activated GST-JNK2 was purified using Ni-NTA chromatography.

Assay platform : Mobility Shift Assay
Substrate : Modified Erktide

ATP (μ M) Km app / Bin : 21 / 50

Metal : Mg
Reference compound : K252a
IC50 at ATP Bin (nM) : 110
IC50 at 1 mM ATP (nM) : 1600

JNK3(MAPK10)

Product code 04-150

Full-length human JNK3 [1-426(end) amino acids of accession number NP_620446.1] was expressed as N-terminal GST-fusion protein (75 kDa) using E.coli expression system. GST-JNK3 was purified by using glutathione sepharose chromatography and activated with His-tagged MAP2K4 and MAP2K7. Activated GST-JNK3 was purified using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay
Substrate : Modified Erktide

ATP (μM) Km app / Bin : 6 / 25 Metal : Mg Reference compound : K252a IC50 at ATP Bin (nM) : 26 IC50 at 1 mM ATP (nM) : 730

KDR

Product code 08-191

Human KDR, cytoplasmic domain [790-1356(end) amino acids of accession number NP_002244.1] was expressed as N-terminal GST-fusion protein (90 kDa) using baculovirus expression system. GST-KDR was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CSKtide

ATP (μ M) Km app / Bin : 74 / 75 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 3.1 IC50 at 1 mM ATP (nM) : 13

KIT

Product code 08-156

Human KIT, cytoplasmic domain [544-976(end) amino acids of accession number NP_000213.1] was expressed as N-terminal GST-fusion protein (76 kDa) using baculovirus expression system. GST-KIT was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 370 / 400

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 1.2 IC50 at 1 mM ATP (nM) : 2.0



KIT[D816E]

Product code 08-541

Human KIT, cytoplasmic domain [544-976(end) amino acids and D816E of accession number NP_000213.1] was expressed as N-terminal GST-fusion protein (76 kDa) using baculovirus expression system. GST-KIT[D816E] was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 40 / 50 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 1.7 IC50 at 1 mM ATP (nM) : 1.3

KIT[D816V]

Product code 08-505

Human KIT, cytoplasmic domain [544-976(end) amino acids and D816V of accession number NP_000213.1] was expressed as N-terminal GST-fusion protein (76 kDa) using baculovirus expression system. GST-KIT[D816V] was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 14 / 10 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.18 IC50 at 1 mM ATP (nM) : 2.8

KIT[D816Y]

Product code 08-534

Human KIT, cytoplasmic domain [544-976(end) amino acids and D816Y of accession number NP_000213.1] was expressed as N-terminal GST-fusion protein (76 kDa) using baculovirus expression system. GST-KIT[D816Y] was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 22 / 25 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.27 IC50 at 1 mM ATP (nM) : 2.1

KIT[T670I]

Product code 08-195

Human KIT, cytoplasmic domain [544-976(end) amino acids and T670I of accession number NP_000213.1] was expressed as N-terminal GST-fusion protein (76 kDa) using baculovirus expression system. GST-KIT[T670I] was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (µM) Km app / Bin : 100 / 100

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.80 IC50 at 1 mM ATP (nM) : 3.4

KIT[V560G]

Product code 08-504

Human KIT, cytoplasmic domain [544-976(end) amino acids and V560G of accession number NP_000213.1] was expressed as N-terminal GST-fusion protein (76 kDa) using baculovirus expression system. GST-KIT[V560G] was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 110 / 250

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 1.2 IC50 at 1 mM ATP (nM) : 1.6



KIT[V654A]

Product code 08-511

Human KIT, cytoplasmic domain [544-976(end) amino acids and V654A of accession number NP_000213.1] was expressed as N-terminal GST-fusion protein (76 kDa) using baculovirus expression system. GST-KIT[V654A] was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 220 / 250

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 4.5 IC50 at 1 mM ATP (nM) : 8.2

LATS2

Product code 01-124

Human LATS2, catalytic domain [553-1088(end) amino acids of accession number NP_055387.2] was co-expressed as N-terminal GST-fusion protein (89 kDa) with human His-tagged MOBKL1A [1-216(end) amino acids of accession number NP_775739.1] using baculovirus expression system. GST-LATS2 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : SGKtide

ATP (µM) Km app / Bin : 380 / 400

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 1.6 IC50 at 1 mM ATP (nM) : n.a.

LCK

Product code 08-170

Full-length human LCK [1-509(end) amino acids of accession number NP_005347.2] was expressed as N-terminal GST-fusion protein (85 kDa) using baculovirus expression system. GST-LCK was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 14 / 10 Metal : Ma

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 1.5 IC50 at 1 mM ATP (nM) : 14

LOK(STK10)

Product code 07-315

Full-length human LOK [1-968(end) amino acids of accession number BAA35073.1] was expressed as N-terminal GST-fusion protein using baculovirus expression system. GST-LOK was purified by using glutathione sepharose chromatography. GST-LOK was cleaved by PreScission protease and GST-free LOK (114 kDa) was collected as flow-through fraction from glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay
Substrate : Moesin-derived peptide

ATP (µM) Km app / Bin : 100 / 100

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.49 IC50 at 1 mM ATP (nM) : n.a.

LTK

Product code 08-106

Human LTK, catalytic domain [498-796 amino acids of accession number NP_002335.2] was expressed as N-terminal GST-fusion protein (60 kDa) using baculovirus expression system. GST-LTK was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 49 / 50 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 2.0 IC50 at 1 mM ATP (nM) : 7.1



LYNa

Product code 08-171

Full-length human LYNa [1-512(end) amino acids of accession number NP_002341.1] was expressed as N-terminal GST-fusion protein (86 kDa) using baculovirus expression system. GST-LYNa was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 14 / 10 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 2.0 IC50 at 1 mM ATP (nM) : 22

LYNb

Product code 08-172

Full-length human LYNb [1-491(end) amino acids of accession number AAB50019.1] was expressed as N-terminal GST-fusion protein (83 kDa) using baculovirus expression system. GST-LYNb was purified by using glutathione sepharose chromatography and anion exchange chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 18 / 25 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 2.0 IC50 at 1 mM ATP (nM) : 21

MAP4K2

Product code 07-111

Full-length human MAP4K2 [1-820(end) amino acid of accession number NP_004570.2] was expressed as N-terminal GST-fusion protein (119 kDa) using baculovirus expression system. GST-MAP4K2 was purified by using glutathione sepharose chromatography

Assay platform : Mobility Shift Assay

Substrate : S6K2 peptide

ATP (μM) Km app / Bin : 93 / 100

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 1.3 IC50 at 1 mM ATP (nM) : n.a.

MAPKAPK2

Product code 02-142

Full-length human MAPKAPK2 [1-400(end) amino acids of accession number NP_116584.2] was co-expressed as N-terminal GST-fusion protein (73 kDa) with human His-tagged p38 β [1-364(end) amino acids of accession number NP_002742.3] and human His-tagged MAP2K6 [1-334(end) amino acids of accession number NP_002749.2] using baculovirus expression system. GST-MAPKAPK2 was purified by using glutathione sepharose chromatography and Ni-NTA chromatography.

Assay platform : Mobility Shift Assay

Substrate : GS peptide

ATP (μ M) Km app / Bin : 3.6 / 5 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 80 IC50 at 1 mM ATP (nM) : 9300

MAPKAPK3

Product code 02-143

Full-length human MAPKAPK3 [1-382(end) amino acids of accession number NP_004626.1] was co-expressed as N-terminal GST-fusion protein (70 kDa) with human His-tagged p38 β [1-364(end) amino acids of accession number NP_002742.3] and human His-tagged MAP2K6 [1-334(end) amino acids of accession number NP_002749.2] using baculovirus expression system. GST-MAPKAPK3 was purified by using glutathione sepharose chromatography and Ni-NTA chromatography.

Assay platform : Mobility Shift Assay

Substrate : GS peptide

ATP (μM) Km app / Bin : 13 / 10

Metal : Mg
Reference compound : K252a
IC50 at ATP Bin (nM) : 410
IC50 at 1 mM ATP (nM) : n.a.



MAPKAPK5

Product code 02-144

Full-length human MAPKAPK5 [1-471(end) amino acids of accession number NP_003659.2] was co-expressed as N-terminal GST-fusion protein (81 kDa) with human His-tagged p38 β [1-364(end) amino acids of accession number NP_002742.3] and human His-tagged MAP2K6 [1-334(end) amino acids of accession number NP_002749] using baculovirus expression system. GST-MAPKAPK5 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : GS peptide

ATP (μ M) Km app / Bin : 12 / 10 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 320 IC50 at 1 mM ATP (nM) : n.a.

MARK1

Product code 02-120

Full-length human MARK1 [1-795(end) amino acids of accession number AAF72103.1] was expressed as N-terminal GST-fusion protein (116 kDa) using baculovirus expression system. GST-MARK1 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CHKtide

ATP (μ M) Km app / Bin : 8 / 10 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.17 IC50 at 1 mM ATP (nM) : n.a.

MARK2

Product code 02-121

Full-length human MARK2 [1-745(end) amino acids of accession number NP_059672.2] was expressed as N-terminal GST-fusion protein (110 kDa) using baculovirus expression system. GST-MARK2 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CHKtide

ATP (μM) Km app / Bin : 8.8 / 10 Metal : Mα

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.12 IC50 at 1 mM ATP (nM) : n.a.

MARK3

Product code 02-122

Full-length human MARK3 [1-729(end) amino acids of accession number NP_002367.4] was expressed as N-terminal GST-fusion protein (108 kDa) using baculovirus expression system. GST-MARK3 was purified by using glutathione sepharose chromatography and anion exchange chromatography.

Assay platform : Mobility Shift Assay

Substrate : CHKtide

ATP (μ M) Km app / Bin : 5 / 5 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.20 IC50 at 1 mM ATP (nM) : n.a.

MARK4

Product code 02-123

Full-length human MARK4 [1-688(end) amino acids of accession number NP_113605.2] was expressed as N-terminal GST-fusion protein (103 kDa) using baculovirus expression system. GST-MARK4 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CHKtide

ATP (μ M) Km app / Bin : 12 / 10 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.12 IC50 at 1 mM ATP (nM) : n.a.



MELK

Product code 02-124

Truncated human MELK [1-493 amino acids of accession number NP_055606.1] was expressed as N-terminal GST-fusion protein (83 kDa) using E. coli expression system. GST-MELK was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : GS peptide

ATP (μ M) Km app / Bin : 38 / 50 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.81 IC50 at 1 mM ATP (nM) : n.a.

MER(MERTK)

Product code 08-108

Human MER, cytoplasmic domain [528-999(end) amino acids of accession number NP_006334.2] was expressed as N-terminal GST-fusion protein (80 kDa) using baculovirus expression system. GST-MER was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CSKtide

ATP (μ M) Km app / Bin : 36 / 50 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.61 IC50 at 1 mM ATP (nM) : 5.3

MET

Product code 08-151

Human MET, cytoplasmic domain [956-1390(end) amino acids of accession number NP_000236.2] was expressed as N-terminal GST-fusion protein (76 kDa) using baculovirus expression system. GST-MET was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 27 / 25 Metal : Ma

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 67 IC50 at 1 mM ATP (nM) : 730

MET[D1228H]

Product code 08-540

Human MET, cytoplasmic domain [956-1390(end) amino acids and D1228H of accession number NP_000236.2] was expressed as N-terminal GST-fusion protein (76 kDa) using baculovirus expression system. GST-MET[D1228H] was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 25 / 25 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 59 IC50 at 1 mM ATP (nM) : 1200

MET[M1250T]

Product code 08-545

Human MET, cytoplasmic domain [956-1390(end) amino acids and M1250T of accession number NP_000236.2] was expressed as N-terminal GST-fusion protein (76 kDa) using baculovirus expression system. GST-MET[M1250T] was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 17 / 25 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 86 IC50 at 1 mM ATP (nM) : 1900



MET[Y1235D]

Product code 08-198

Human MET, cytoplasmic domain [956-1390(end) amino acids and Y1235D of accession number NP_000236.2] was expressed as N-terminal GST-fusion protein (76 kDa) using baculovirus expression system. GST-MET[Y1235D] was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 71 / 75 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 79 IC50 at 1 mM ATP (nM) : 390

MINK(MINK1)

Product code 07-139

Human MINK, catalytic domain [1-314 amino acids of accession number NP_056531.1] was expressed as N-terminal GST-fusion protein (63 kDa) using baculovirus expression system. GST-MINK was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay
Substrate : Modified Erktide

ATP (μ M) Km app / Bin : 16 / 50 Metal : Mg

Reference compound : K252a IC50 at ATP Bin (nM) : 5.5 IC50 at 1 mM ATP (nM) : 4.7

MNK1(MKNK1)

Product code 02-145

Full-length human MNK1 [1-424(end) amino acids and T344D of accession number BAA19885.1] was expressed as N-terminal GST-fusion protein (74 kDa) using baculovirus expression system. GST-MNK1 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : RS peptide

ATP (µM) Km app / Bin : 460 / 450

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 21 IC50 at 1 mM ATP (nM) : n.a.

MNK2(MKNK2)

Product code 02-146

Full-length human MNK2 [1-465(end) amino acids and T379D of accession number NP_951009.1] was expressed as N-terminal GST-fusion protein (79 kDa) using baculovirus expression system. GST-MNK2 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : RS peptide

ATP (µM) Km app / Bin : 110 / 100

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 7.5 IC50 at 1 mM ATP (nM) : n.a.

MRCKα(CDC42BPA)

Product code 01-107

Truncated human MRCK α [1-574 amino acids of accession number NP_003598.2] was expressed as N-terminal GST-fusion protein (93 kDa) using baculovirus expression system. GST-MRCK α was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : DAPK1tide

ATP (μ M) Km app / Bin : 0.45 / 1

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 2.2 IC50 at 1 mM ATP (nM) : n.a.



MRCKβ(CDC42BPB)

Product code 01-108

Truncated human MRCKß [1-473 amino acids of accession number NP_006026.3] was expressed as N-terminal GST-fusion protein (82 kDa) using baculovirus expression system. GST-MRCKβ was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : DAPK1tide

ATP (μ M) Km app / Bin : 0.67 / 1

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) IC50 at 1 mM ATP (nM): n.a.

MSK1(RPS6KA5)

Product code 01-147

Full-length human MSK1 [1-802(end) amino acids of accession number NP_004746.2] was co-expressed as N-terminal GST-fusion protein (117 kDa) with human His-tagged Erk2 [1-360 amino acids of accession number NP 002736.3] using baculovirus expression system. GST-MSK1 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

: Mg

Substrate : Crosstide

ATP (µM) Km app / Bin : 13 / 10 Metal

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 1.6 IC50 at 1 mM ATP (nM): n.a.

MSK2(RPS6KA4)

Product code 01-148

Full-length human MSK2 [1-772(end) amino acids of accession number NP_003933.1] was co-expressed as N-terminal GST-fusion protein (114 kDa) with human His-tagged Erk2 [1-360 amino acids of accession number NP_002736.3] using baculovirus expression system. GST-MSK2 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Crosstide

ATP (µM) Km app / Bin : 40 / 50 Metal : Ma

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 4.3 IC50 at 1 mM ATP (nM): n.a.

MSSK1(STK23)

Product code 04-159

Full-length human MSSK1 [1-567(end) amino acids of accession number NP_055185.2] was expressed as N-terminal GST-fusion protein (89 kDa) using baculovirus expression system, GST-MSSK1 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : DYRKtide-F

ATP (μM) Km app / Bin : 56 / 50

Metal : Mg Reference compound : K252a

IC50 at ATP Bin (nM) : 220 IC50 at 1 mM ATP (nM): n.a.

MST1(STK4)

Product code 07-116

Full-length human MST1 [1-487(end) amino acids of accession number NP_006273.1] was expressed as N-terminal GST-fusion protein (83 kDa) using baculovirus expression system. GST-MST1 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : IRS1

ATP (µM) Km app / Bin : 50 / 50

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 1.0 IC50 at 1 mM ATP (nM): 0.55



MST2(STK3)

Product code 07-117

Full-length human MST2 [1-491(end) amino acids of accession number NP_006272.2] was expressed as N-terminal GST-fusion protein (83 kDa) using baculovirus expression system. GST-MST2 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : IRS1

ATP (μ M) Km app / Bin : 69 / 75 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 6.7 IC50 at 1 mM ATP (nM) : n.a.

MST3(STK24)

Product code 07-118

Full-length human MST3 [1-431(end) amino acids of accession number NP_001027467.2] was expressed as N-terminal GST-fusion protein (75 kDa) using baculovirus expression system. GST-MST3 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Moesin-derived peptide

ATP (μ M) Km app / Bin : 66 / 75 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 1.9 IC50 at 1 mM ATP (nM) : n.a.

MST4

Product code 07-119

Full-length human MST4 [1-416(end) amino acids of accession number NP_057626.2] was expressed as N-terminal GST-fusion protein (74 kDa) using baculovirus expression system. GST-MST4 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Moesin-derived peptide

ATP (μM) Km app / Bin : 76 / 75 Metal : Mα

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 6.3 IC50 at 1 mM ATP (nM) : n.a.

MUSK

Product code 08-153

Human MUSK, catalytic domain [527-869(end) amino acids of accession number NP_005583.1] was expressed as N-terminal GST fusion protein (66 kDa) using baculovirus expression system. GST-MUSK was purified by using glutathione sepharose chromatography and anion exchange chromatography.

Assay platform : Mobility Shift Assay

Substrate : CSKtide

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 2.1 IC50 at 1 mM ATP (nM) : 2.6

NDR1(STK38)

Product code 01-125

Full-length human NDR1[1-465(end) amino acids of accession number NP_009202.1] was co-expressed as N-terminal GST-fusion protein (81kDa) with human His-tagged MOBKL1A [1-216(end) amino acids of accession number NP_775739.1] using baculovirus expression system. GST-NDR1 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : SGKtide

ATP (µM) Km app / Bin : 12 / 10

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 1.8 IC50 at 1 mM ATP (nM) : n.a.



NDR2(STK38L)

Product code 01-126

Full-length human NDR2 [1-464(end) amino acids of accession number NP_055815.1] was co-expressed as N-terminal GST-fusion protein (81 kDa) with human His-tagged MOBKL1A [1-216(end) amino acids of accession number NP_775739.1] using baculovirus expression system. GST-NDR2 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : SGKtide

ATP (μM) Km app / Bin : 7.6 / 10

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 2.1 IC50 at 1 mM ATP (nM) : n.a.

NEK1

Product code 05-123

Human NEK1, catalytic domain [1-505 amino acids of accession number NP_036356.1] was expressed as N-terminal GST-fusion protein (85 kDa) using baculovirus expression system. GST-NEK1 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CDK7 peptide

ATP (μ M) Km app / Bin : 64 / 75 Metal : Ma

Metal : Mg
Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 51 IC50 at 1 mM ATP (nM) : 650

NEK2

Product code 05-226

Full-length human NEK2 [1-445(end) amino acids of accession number NP_002488.1] was expressed as N-terminal His-tagged protein (55 kDa) using baculovirus expression system. His-tagged NEK2 was purified by using Ni-NTA affinity chromatography. Purified His-NEK2 was digested by recombinant His-TEV protease, and His-tag free NEK2 (ca. 52 kDa) was collected as flow-through fraction from Ni-NTA affinity chromatography.

Assay platform : Mobility Shift Assay

Substrate : CDK7 peptide

ATP (μ M) Km app / Bin : 65 / 75 Metal : Ma

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 3700 IC50 at 1 mM ATP (nM) : >10000

NEK4

Product code 05-128

Full-length human NEK4 [1-841(end) amino acids of accession number NP_003148.2] was expressed as N-terminal GST-fusion protein (122 kDa) using baculovirus expression system. GST-NEK4 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : GS peptide

ATP (μ M) Km app / Bin : 51 / 50 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 120 IC50 at 1 mM ATP (nM) : n.a.

NEK6

Product code 05-130

Full-length human NEK6 [1-313(end) amino acids of accession number NP_055212.2] was expressed as N-terminal GST-fusion protein (63 kDa) using baculovirus expression system. GST-NEK6 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CDK7 peptide

ATP (μ M) Km app / Bin : 69 / 75 Metal : Mg

Reference compound : PKR Inhibitor

IC50 at ATP Bin (nM) : 19000 IC50 at 1 mM ATP (nM) : >10000



NEK7

Product code 05-131

Full-length human NEK7 [1-302(end) amino acids of accession number NP_598001.1] was expressed as N-terminal GST-fusion protein (62 kDa) using baculovirus expression system. GST-NEK7 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CDK7 peptide

ATP (μ M) Km app / Bin : 40 / 50 Metal : Mg

Reference compound : PKR Inhibitor

IC50 at ATP Bin (nM) : 8500 IC50 at 1 mM ATP (nM) : >10000

NEK9

Product code 05-133

Truncated human NEK9 [1-346, 733-979(end) amino acids of accession number NP_149107.4] was expressed as N-terminal GST-fusion protein (93 kDa) using baculovirus expression system. GST-NEK9 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CDK7 peptide

ATP (μM) Km app / Bin : 190 / 200

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 150 IC50 at 1 mM ATP (nM) : 400

NIM1K(MGC42105)

Product code 02-175

Full-length human NIM1K [1-436(end) amino acids of accession number NP_699192.1] was expressed as N-terminal GST-fusion protein (76 kDa) using baculovirus expression system. GST-NIM1K was purified by using glutathione sepharose chromatography and anion exchange chromatography.

Assay platform : Mobility Shift Assay

Substrate : CHKtide

ATP (μ M) Km app / Bin : 21 / 25 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 300 IC50 at 1 mM ATP (nM) : n.a.

Nuak1

Product code 02-126

Full-length human NuaK1 [1-661(end) amino acids of accession number NP_055655.1] was expressed as N-terminal GST-fusion protein (102 kDa) using baculovirus expression system. GST-NuaK1 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CHKtide

ATP (μ M) Km app / Bin : 59 / 50 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 1.0 IC50 at 1 mM ATP (nM) : n.a.

NuaK2

Product code 02-127

Full-length human NuaK2 [1-628(end) amino acids of accession number NP_112214.1] was expressed as N-terminal GST-fusion protein (98kDa) using baculovirus expression system. GST-NuaK2 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CHKtide

ATP (μ M) Km app / Bin : 26 / 25 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 2.2 IC50 at 1 mM ATP (nM) : n.a.



p38α(MAPK14)

Product code 04-152

Truncated human p38 α [9-352 amino acids of accession number NP_620581.1] was expressed as N-terminal GST-fusion protein (66 kDa) using E. coli expression system. GST-p38 α was purified by using glutathione sepharose chromatography and activated with His-tagged MAP2K6. Activated GST-p38 α was purified using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Modified Erktide

ATP (μ M) Km app / Bin : 150 / 150

Metal : Mg

Reference compound : SB202190

IC50 at ATP Bin (nM) : 6.3 IC50 at 1 mM ATP (nM) : 22

p38β(MAPK11)

Product code 04-153

Full-length human p38 β [1-364(end) amino acids of accession number NP_002742.3] was expressed as N-terminal GST-fusion protein (69 kDa) using E. coli expression system. GST-p38 β was purified by using glutathione chromatography.

Assay platform : Mobility Shift Assay
Substrate : Modified Erktide

ATP (μ M) Km app / Bin : 63 / 75 Metal : Mg

Reference compound : SB202190

IC50 at ATP Bin (nM) : 16 IC50 at 1 mM ATP (nM) : 110

p38y(MAPK12)

Product code 04-155

Full-length human p38 γ [1-367(end) amino acids of accession number NP_002960.2] was expressed as N-terminal GST-fusion protein (69 kDa) using E. coli expression system. GST-p38 γ was purified by using glutathione sepharose chromatography and activated with His-tagged MAP2K6. Activated GST-p38 γ was purified using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay
Substrate : Modified Erktide

ATP (μ M) Km app / Bin : 13 / 10 Metal : Ma

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 88 IC50 at 1 mM ATP (nM) : 2800

p38δ(MAPK13)

Product code 04-154

Full-length human p385 [1-365(end) amino acids of accession number NP_002745.1] was expressed as N-terminal GST-fusion protein (69 kDa) using E. coli expression system. GST-p385 was purified by using glutathione sepharose chromatography and activated with His-tagged MAP2K6. Activated GST-p385 was purified using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay Substrate : Modified Erktide

ATP (μ M) Km app / Bin : 5.8 / 5 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 220 IC50 at 1 mM ATP (nM) : >10000

p70S6K(RPS6KB1)

Product code 01-154

Human p70S6K, catalytic domain [1-421 amino acids and T412E of accession number NP_003152.1] was expressed as N-terminal GST-fusion protein (75 kDa) using baculovirus expression system. GST-p70S6K was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : S6K2 peptide

ATP (μ M) Km app / Bin : 14 / 10 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 2.1 IC50 at 1 mM ATP (nM) : 9.8



p70S6Kβ(RPS6KB2)

Product code 01-155

Full-length human p70S6Kβ [1-482(end) amino acids of accession number NP_003943.2] was expressed as N-terminal GST-fusion protein (81 kDa) using baculovirus expression system. GST-p70S6Kβ was purified by using glutathione sepharose chromatography

Assay platform : Mobility Shift Assay

Substrate : S6K2 peptide

ATP (μ M) Km app / Bin : 3.3 / 5 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 2.7 IC50 at 1 mM ATP (nM) : n.a.

PAK1

Product code 07-123

Full-length human PAK1 [1-545(end) amino acids of accession number NP_002567.3] was expressed as N-terminal GST-fusion protein (88 kDa) using baculovirus expression system. GST-PAK1 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : LIMKtide

ATP (µM) Km app / Bin : 300 / 300

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 4.0 IC50 at 1 mM ATP (nM) : n.a.

PAK2

Product code 07-124

Full-length human PAK2 [1-524(end) amino acids of accession number NP_002568.2] was expressed as N-terminal GST-fusion protein (85 kDa) using baculovirus expression system. GST-PAK2 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : DAPK1tide

ATP (μM) Km app / Bin : 81 / 100

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 4.5 IC50 at 1 mM ATP (nM) : 22

PAK4

Product code 07-126

Full-length human PAK4 [1-591(end) amino acids of accession number NP_005875.1] was expressed as N-terminal GST-fusion protein (91 kDa) using baculovirus expression system. GST-PAK4 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : SGKtide

ATP (μ M) Km app / Bin : 2.5 / 5 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 12 IC50 at 1 mM ATP (nM) : n.a.

PAK5(PAK7)

Product code 07-127

Human PAK5, catalytic domain [425-719(end) amino acids of accession number NP_065074.1] was expressed as N-terminal GST-fusion protein (60 kDa) using baculovirus expression system. GST-PAK5 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : DAPK1tide

ATP (µM) Km app / Bin : 1.9 / 1

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 2.5 IC50 at 1 mM ATP (nM) : n.a.



PAK6

Product code 07-128

Full-length human PAK6 [1-681(end) amino acids of accession number NP_064553.1] was expressed as N-terminal GST-fusion protein (102 kDa) using baculovirus expression system. GST-PAK6 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : SGKtide

ATP (μ M) Km app / Bin : 3.7 / 5 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 1.2 IC50 at 1 mM ATP (nM) : n.a.

PASK

Product code 02-128

Human PASK, catalytic domain [949-1323(end) amino acids of accession number NP_055963.2] was expressed as N-terminal GST-fusion protein (69 kDa) using baculovirus expression system. GST-PASK was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : GS peptide

ATP (μM) Km app / Bin : 9.7 / 10

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 13 IC50 at 1 mM ATP (nM) : n.a.

PBK

Product code 05-168

Full-length human PBK [1-322(end) amino acids of accession number NP_060962.2] was expressed as N-terminal GST-fusion protein (63 kDa) using baculovirus expression system. GST-PBK was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Histone H3 peptide

ATP (μ M) Km app / Bin : 33 / 50 Metal : Ma

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 69 IC50 at 1 mM ATP (nM) : 720

PDGFRα(PDGFRA)

Product code 08-157

Human PDGFR α , cytoplasmic domain [550-1089(end) amino acids of accession number NP_006197.1] was expressed as N-terminal GST-fusion protein(89 kDa) using baculovirus expression system. GST-PDGFR α was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CSKtide

ATP (μ M) Km app / Bin : 28 / 25 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.30 IC50 at 1 mM ATP (nM) : 1.4

PDGFRα(PDGFRA)[D842V]

Product code 08-506

Human PDGFR α , cytoplasmic domain [550-1089(end) amino acids and D842V of accession number NP_006197.1] was expressed as N-terminal GST-fusion protein (89 kDa) using baculovirus expression system. GST-PDGFR α [D842V] was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CSKtide

ATP (μ M) Km app / Bin : 21 / 25 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.25 IC50 at 1 mM ATP (nM) : 1.9



PDGFRα(PDGFRA)[T674I]

Product code 08-503

Human PDGFR α , cytoplasmic domain [550-1089(end) amino acids and T674I of accession number NP_006197.1] was expressed as N-terminal GST-fusion protein (89 kDa) using baculovirus expression system. GST-PDGFR α [T674I] was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CSKtide

ATP (μ M) Km app / Bin : 11 / 10 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.12 IC50 at 1 mM ATP (nM) : 1.1

PDGFRα(PDGFRA)[V561D]

Product code 08-507

Human PDGFR α , cytoplasmic domain [550-1089(end) amino acids and V561D of accession number NP_006197.1] was expressed as N-terminal GST-fusion protein (89 kDa) using baculovirus expression system. GST-PDGFR α [V561D] was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CSKtide

ATP (μ M) Km app / Bin : 35 / 50 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.32 IC50 at 1 mM ATP (nM) : 1.6

PDGFRβ(PDGFRB)

Product code 08-158

Human PDGFR β , cytoplasmic domain [557-1106(end) amino acids of accession number NP_002600.1] was expressed as N-terminal GST-fusion protein (88 kDa) using baculovirus expression system. GST-PDGFR β was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CSKtide

ATP (μ M) Km app / Bin : 23 / 25 Metal : Ma

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.27 IC50 at 1 mM ATP (nM) : 0.62

PDHK2(PDK2)

Product code 10-140

Full-length human PDHK2 [1-407(end) amino acids of accession number NP_002602.2] was expressed as N-terminal GST-fusion protein (74 kDa) using baculovirus expression system. GST-PDHK2 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : PDHKtide

ATP (µM) Km app / Bin : 28 / 25

Metal : Mg+K

Reference compound : DCA

IC50 at ATP Bin (nM) : 610000

IC50 at 1 mM ATP (nM) : n.a.

PDHK4(PDK4)

Product code 10-125

Full-length human PDHK4 [1-411(end) amino acids of accession number NP_002603.1] was expressed as N-terminal GST-fusion protein (73 kDa) using E.coli expression system. GST-PDHK4 was purified by using glutathione affinity chromatography.

Assay platform : Mobility Shift Assay

Substrate : PDHKtide



PDK1(PDPK1)

Product code 01-132

Full-length human PDK1 [1-556(end) amino acids of accession number NP_002604.1] was expressed as N-terminal GST-fusion protein (91 kDa) using baculovirus expression system. GST-PDK1 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : T308tide

ATP (μ M) Km app / Bin : 9.6 / 10

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 9.2 IC50 at 1 mM ATP (nM) : 12

PEK(EIF2AK3)

Product code 05-155

Human PEK, cytoplasmic domain [536-1116(end) amino acids of accession number NP_004827.3] was expressed as N-terminal GST-fusion protein (94 kDa) using E.coli expression system. GST-PEK was purified by using glutathione sepharose chromatography and anion exchange chromatography.

Assay platform : IMAP

Substrate : SRPKtide

ATP (μM) Km app / Bin : 13 / 10

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 3600 IC50 at 1 mM ATP (nM) : n.a.

Metal

PGK(PRKG1)

Product code 01-142

Full-length human PGK [1-686(end) amino acids of accession number NP_006249.1] was expressed as N-terminal GST-fusion protein (105 kDa) using baculovirus expression system. GST-PGK was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

: Mg

Substrate : Kemptide

ATP (μM) Km app / Bin : 8.2 / 10

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 2.1 IC50 at 1 mM ATP (nM) : n.a.

PHKG1

Product code 02-152

Full-length human PHKG1 [1-387(end) amino acids of accession number NP_006204.1] was expressed as N-terminal GST-fusion protein (72 kDa) using baculovirus expression system. GST-PHKG1 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : GS peptide

ATP (μM) Km app / Bin : 71 / 75

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.22 IC50 at 1 mM ATP (nM) : n.a.

PHKG2

Product code 02-153

Full-length human PHKG2 [1-406(end) amino acids of accession number NP_000285.1] was expressed as N-terminal GST-fusion protein (74 kDa) using baculovirus expression system. GST-PHKG2 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : GS peptide

ATP (µM) Km app / Bin : 8.1 / 10

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.74 IC50 at 1 mM ATP (nM) : n.a.



PIK3CA/PIK3R1

Product code 11-401-20N

Full-length human PIK3CA[1-1068(end) amino acids of accession number NP_006209.2] was co-expressed as N-terminal DYKDDDDK tagged, biotinylated protein (128 kDa) with PIK3R1[1-724(end) amino acids of accession number NP 852664.1] using baculovirus expression system. The protein was purified by using DYKDDDDK tag antibody agarose.

Assay platform : ADP-Glo

Substrate : PI(4,5)P2

ATP (µM) Km app / Bin : 89 / 100

Metal : Mg Reference compound : PI-103 IC50 at ATP Bin (nM)

IC50 at 1 mM ATP (nM): n.a.

PIK3CB/PIK3R1

Product code 11-402-20N

Full-length human PIK3CB[1-1070(end) amino acids of accession number NP_006210.1] was co-expressed as N-terminal DYKDDDDK tagged, biotinylated protein (126 kDa) with PIK3R1[1-724(end) amino acids of accession number NP 852664.1] (84kDa) using baculovirus expression system. The protein was purified by using DYKDDDDK tag antibody agarose.

Assay platform : ADP-Glo

: PI(4,5)P2

Substrate

ATP (µM) Km app / Bin : 88 / 100

Metal : Mg Reference compound : PI-103 IC50 at ATP Bin (nM) : 22 IC50 at 1 mM ATP (nM): n.a.

PIK3CD/PIK3R1

Product code 11-403-20N

Full-length human PIK3CD[1-1044(end) amino acids of accession number NP_005017.3] was co-expressed as N-terminal DYKDDDDK tagged, biotinylated protein (123 kDa) with PIK3R1[1-724(end) amino acids of accession number NP_852664.1] (84kDa) using baculovirus expression system. The protein was purified by using DYKDDDDK tag antibody agarose.

Assay platform : ADP-Glo Substrate : PI(4,5)P2

ATP (µM) Km app / Bin : 37 / 50 Metal : Ma Reference compound : PI-103 IC50 at ATP Bin (nM) : 24

IC50 at 1 mM ATP (nM): n.a.

PIKFYVE(PIP5K3)

Product code 11-118

Full-length human PIKFYVE [1-2098(end) amino acids and S696N, L932S, Q995L, T998S, S1033A and Q1183K of accession number NP 055855.21 was expressed as N-terminal GST-fusion protein (265 kDa) using baculovirus expression system. GST-PIKFYVE was purified by using glutathione sepharose chromatography.

Assay platform : ADP-Glo Substrate : PI(3)P

ATP (µM) Km app / Bin : 36 / 50

Metal : Mg Reference compound : AG-183

IC50 at ATP Bin (nM) : 3900 IC50 at 1 mM ATP (nM): n.a.

PIM1

Product code 02-054

Full-length human PIM1 [1-313(end) amino acids of accession number NP_002639.1] was expressed as N-terminal His-tagged protein (39 kDa) using baculovirus expression system. His-tagged PIM1 was purified by using Ni-NTA affinity chromatography.

Assay platform : Mobility Shift Assay

Substrate : S6K2 peptide

ATP (µM) Km app / Bin : 640 / 500

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 10 IC50 at 1 mM ATP (nM): 20



PIM₂

Product code 02-155

Full-length human PIM2 [1-311(end) amino acids of accession number NP_006866.2] was expressed as N-terminal GST-fusion protein (61 kDa) using baculovirus expression system. GST-PIM2 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : S6K2 peptide

ATP (μ M) Km app / Bin : 4 / 5 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 14 IC50 at 1 mM ATP (nM) : 480

PIM₃

Product code 02-156

Full-length human PIM3 [1-326(end) amino acids of accession number NP_001001852.1] was expressed as N-terminal GST-fusion protein (63 kDa) using baculovirus expression system. GST-PIM3 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : S6K2 peptide

ATP (μM) Km app / Bin : 130 / 150

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.36 IC50 at 1 mM ATP (nM) : n.a.

PIP4K2A

Product code 11-115

Full-length human PIP4K2A [1-406(end) amino acids of accession number NP_005019] was expressed as N-terminal GST-fusion protein (73 kDa) using baculovirus expression system. GST-PIP4K2A was purified by using glutathione sepharose chromatography.

Assay platform : ADP-Glo Substrate : PI(5)P

ATP (μ M) Km app / Bin : 20 / 25 Metal : Mg

Reference compound : AG-183 IC50 at ATP Bin (nM) : 7600 IC50 at 1 mM ATP (nM) : n.a.

PIP4K2B

Product code 11-116

Full-length human PIP4K2B [1-416(end) amino acids of accession number NP_003550] was expressed as N-terminal GST-fusion protein (74 kDa) using baculovirus expression system. GST-PIP4K2B was purified by using glutathione sepharose chromatography.

Assay platform : ADP-Glo Substrate : PI(5)P

Cubstrate : 11(0)1

ATP (μ M) Km app / Bin : 18 / 25

Metal : Mn

Reference compound : AG-183 IC50 at ATP Bin (nM) : 48000 IC50 at 1 mM ATP (nM) : n.a.

PIP5K1A

Product code 11-111

Full-length human PIP5K1A [1-549(end) amino acids of accession number NP_003548.1] was expressed as N-terminal GST-fusion protein (88 kDa) using baculovirus expression system. GST-PIP5K1A was purified by using glutathione sepharose chromatography.

Assay platform : ADP-Glo Substrate : PI(4)P

ATP (μM) Km app / Bin : 28 / 25

Metal : Mg
Reference compound : AG-183
IC50 at ATP Bin (nM) : 10000
IC50 at 1 mM ATP (nM) : n.a.

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PIP5K1B

Product code 11-112

Full-length human PIP5K1B [1-540(end) amino acids of accession number NP_003549.1] was expressed as N-terminal GST-fusion protein (88 kDa) using baculovirus expression system. GST-PIP5K1B was purified by using glutathione sepharose chromatography.

Assay platform : ADP-Glo

Substrate : PI(4)P

ATP (μ M) Km app / Bin : 95 / 100

Metal : Mg

Reference compound : AG-183 IC50 at ATP Bin (nM) : 4300 IC50 at 1 mM ATP (nM) : n.a.

PIP5K1C

Product code 11-113

Full-length human PIP5K1C [1-668(end) amino acids of accession number NP_036530] was expressed as N-terminal GST-fusion protein (101 kDa) using baculovirus expression system. GST-PIP5K1C was purified by using glutathione sepharose chromatography.

Assay platform : ADP-Glo

Substrate

: PI(4)P

ATP (μ M) Km app / Bin : 33 / 50 Metal : Mg Reference compound : AG-183 IC50 at ATP Bin (nM) : 1900

IC50 at 1 mM ATP (nM): n.a.

PIP5KL1

Product code 11-114

Full-length human PIP5KL1 [1-394(end) amino acids of accession number NP_001128691.1] was expressed as N-terminal GST-fusion protein (72 kDa) using baculovirus expression system. GST-PIP5KL1 was purified by using glutathione sepharose chromatography.

Assay platform : ADP-Glo

Substrate : PI(4)P

ATP (μ M) Km app / Bin : 1 / 1

Metal : Mg

Reference compound : AG-183 IC50 at ATP Bin (nM) : 2200 IC50 at 1 mM ATP (nM) : n.a.

PKACα(PRKACA)

Product code 01-127

Full-length human PKAC α [1-351(end) amino acids of accession number NP_002721.1] was expressed as N-terminal GST-fusion protein (68 kDa) using baculovirus expression system. GST-PKAC α was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Kemptide

ATP (µM) Km app / Bin : 2.6 / 5

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.80 IC50 at 1 mM ATP (nM) : 86

PKACβ(PRKACB)

Product code 01-128

Full-length human PKAC β [1-351(end) amino acids of accession number NP_002722.1] was expressed as N-terminal GST-fusion protein (68 kDa) using baculovirus expression system. GST-PKAC β was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Kemptide

ATP (μ M) Km app / Bin : 4.7 / 5

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 1.0 IC50 at 1 mM ATP (nM) : n.a.



PKACγ(PRKACG)

Product code 01-129

Full-length human PKACγ [1-351(end) amino acids of accession number NP_002723.2] was expressed as N-terminal GST-fusion protein (68 kDa) using baculovirus expression system. GST-PKACγ was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Kemptide

ATP (μ M) Km app / Bin : 4.5 / 5 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 3.1 IC50 at 1 mM ATP (nM) : n.a.

PKCα(PRKCA)

Product code 01-133

Full-length human PKC α [1-672(end) amino acids of accession number NP_002728.1] was expressed as N-terminal GST-fusion protein (104 kDa) using baculovirus expression system. GST-PKC α was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : PKC peptide

ATP (μ M) Km app / Bin : 36 / 50 Metal : Mg+Ca

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.33 IC50 at 1 mM ATP (nM) : 3.6

PKCβ1(PRKCB1)

Product code 01-134

Full-length human PKCβ1 [1-671(end) amino acids of accession number NP_997700.1] was expressed as N-terminal GST-fusion protein (104 kDa) using baculovirus expression system. GST-PKCβ1 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : PKC peptide

 $\begin{array}{lll} \text{ATP (μM$) Km app / Bin} & : & 79 \ / & 75 \\ \text{Metal} & : & \text{Mg+Ca} \end{array}$

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.71 IC50 at 1 mM ATP (nM) : n.a.

PKCβ2(PRKCB2)

Product code 01-165

Full-length human PKC β 2 [1-673(end) amino acids of accession number NP_002729.2] was expressed as N-terminal GST-fusion protein (104 kDa) using baculovirus expression system. GST-PKC β 2 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : PKC peptide

ATP (μ M) Km app / Bin : 41 / 50 Metal : Mg+Ca

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.43 IC50 at 1 mM ATP (nM) : n.a.

PKCγ(PRKCG)

Product code 01-137

Full-length human PKCy [1-697(end) amino acids of accession number NP_002730.1] was expressed as N-terminal GST-fusion protein (106 kDa) using baculovirus expression system. GST-PKCy was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : PKC peptide

ATP (μ M) Km app / Bin : 74 / 75 Metal : Mg+Ca

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 1.1 IC50 at 1 mM ATP (nM) : n.a.



PKCδ(PRKCD)

Product code 01-135

Full-length human PKCδ [1-676(end) amino acids of accession number NP_006245.2] was expressed as N-terminal GST-fusion protein (105 kDa) using baculovirus expression system. GST-PKCδ was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : PKC peptide

ATP (μ M) Km app / Bin : 26 / 25 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.22 IC50 at 1 mM ATP (nM) : n.a.

PKCε(PRKCE)

Product code 01-136

Full-length human PKCε [1-737(end) amino acids of accession number NP_005391.1] was expressed as N-terminal GST-fusion protein (111 kDa) using baculovirus expression system. GST-PKCε was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : PKC peptide

ATP (μ M) Km app / Bin : 16 / 25 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.56 IC50 at 1 mM ATP (nM) : 5.6

PKCζ(PRKCZ)

Product code 01-141

Full-length human PKC ζ [1-592(end) amino acids of accession number NP_002735.3] was expressed as N-terminal GST-fusion protein (94kDa) using baculovirus expression system. GST-PKC ζ was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : PKC peptide

ATP (μ M) Km app / Bin : 11 / 10 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 55 IC50 at 1 mM ATP (nM) : n.a.

PKCη(PRKCH)

Product code 01-138

Full-length human PKCη [1-683(end) amino acids of accession number NP_006246.2] was expressed as N-terminal GST-fusion protein (105 kDa) using baculovirus expression system. GST-PKCη was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : PKC peptide

ATP (μ M) Km app / Bin : 36 / 50 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.92 IC50 at 1 mM ATP (nM) : n.a.

PKCθ(PRKCQ)

Product code 01-140

Full-length human PKC0 [1-706(end) amino acids of accession number NP_006248.1] was expressed as N-terminal GST-fusion protein (109 kDa) using baculovirus expression system. GST-PKC0 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : PKC peptide

ATP (μ M) Km app / Bin : 18 / 25 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.72 IC50 at 1 mM ATP (nM) : n.a.



PKC_I(PRKCI)

Product code 01-139

Full-length human PKCI[1-587(end) amino acids of accession number NP_002731.3] was expressed as N-terminal GST-fusion protein (94 kDa) using baculovirus expression system. GST-PKCI was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : PKC peptide

ATP (μ M) Km app / Bin : 27 / 25 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 13 IC50 at 1 mM ATP (nM) : n.a.

PKD1(PRKD1)

Product code 02-157

Full-length human PKD1 [1-912(end) amino acids of accession number NP_002733.1] was expressed as N-terminal GST-fusion protein (129 kDa) using baculovirus expression system. GST-PKD1 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : GS peptide

ATP (μ M) Km app / Bin : 25 / 25 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 1.1 IC50 at 1 mM ATP (nM) : n.a.

PKD2(PRKD2)

Product code 02-158

Full-length human PKD2 [1-878(end) amino acids of accession number NP_057541.2] was expressed as N-terminal GST-fusion protein (124 kDa) using baculovirus expression system. GST-PKD2 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : GS peptide

ATP (μ M) Km app / Bin : 26 / 25 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 1.1 IC50 at 1 mM ATP (nM) : 16

PKD3(PRKD3)

Product code 02-159

Full-length human PKD3 [1-890(end) amino acids of accession number NP_005804.1] was expressed as N-terminal GST-fusion protein (127 kDa) using baculovirus expression system. GST-PKD3 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : GS peptide

ATP (μ M) Km app / Bin : 34 / 50 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.80 IC50 at 1 mM ATP (nM) : n.a.

PKN1

Product code 01-144

Full-length human PKN1 [1-942(end) amino acids of accession number NP_002732.3] was expressed as N-terminal GST-fusion protein (132 kDa) using baculovirus expression system. GST-PKN1 was purified by using glutathione sepharose chromatography.

Assay platform : IMAP

Substrate : S6K peptide

ATP (μ M) Km app / Bin : 19 / 25 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.15 IC50 at 1 mM ATP (nM) : n.a.



PKR(EIF2AK2)

Product code 05-156

Human PKR, catalytic domain [252-551(end) amino acids of accession number NP_002750.1] was expressed as N-terminal GST-fusion protein (62 kDa) using baculovirus expression system. GST-PKR was purified by using glutathione sepharose chromatography and anion exchange cromatography.

Assay platform : IMAP

Substrate : SRPKtide

ATP (μ M) Km app / Bin : 13 / 10 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 87 IC50 at 1 mM ATP (nM) : n.a.

PLK1

Product code 05-157

Full-length human PLK1 [1-603(end) amino acids of accession number NP_005021.2] was expressed as N-terminal GST-fusion protein (95 kDa) using baculovirus expression system. GST-PLK1 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CDC25ctide

ATP (μ M) Km app / Bin : 5.6 / 5 Metal : Mg

Reference compound : GW843682X

IC50 at ATP Bin (nM) : 3.6 IC50 at 1 mM ATP (nM) : 47

PLK₂

Product code 05-158

Full-length human PLK2 [1-685(end) amino acids of accession number NP_006613.2] was expressed as N-terminal GST-fusion protein (105 kDa) using baculovirus expression system. GST-PLK2 was purified by using glutathione sepharose chromatography.

Assay platform : IMAP

Substrate : CHK2 peptide

ATP (μ M) Km app / Bin : 30 / 30 Metal : Mg

Reference compound : GW843682X

IC50 at ATP Bin (nM) : 4.8 IC50 at 1 mM ATP (nM) : n.a.

PLK3

Product code 05-159

Human PLK3, catalytic domain [58-340 amino acids of accession number NP_004064.2] was expressed as N-terminal GST-fusion protein (59 kDa) using baculovirus expression system. GST-PLK3 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CDC25ctide

ATP (μ M) Km app / Bin : 6.8 / 5 Metal : Mg

Reference compound : GW843682X

IC50 at ATP Bin (nM) : 33 IC50 at 1 mM ATP (nM) : 450

PRKX

Product code 01-130

Full-length human PRKX [1-358(end) amino acids of accession number NP_005035.1] was expressed as N-terminal GST-fusion protein (68 kDa) using baculovirus expression system. GST-PRKX was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Kemptide

ATP (μ M) Km app / Bin : 20 / 25 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.59 IC50 at 1 mM ATP (nM) : n.a.



PYK2(PTK2B)

Product code 08-138

Full-length human PYK2 [1-967(end) amino acids of accession number NP_775267.1] was expressed as N-terminal GST-fusion protein (138 kDa) using baculovirus expression system. GST-PYK2 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Blk/Lyntide

ATP (μM) Km app / Bin : 56 / 50 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 2.2 IC50 at 1 mM ATP (nM) : 4.9

QIK(SNF1LK2)

Product code 02-129

Full-length human QIK(SNF1LK2) [1-926(end) amino acids of accession number NP_056006.1] was expressed as N-terminal GST-fusion protein (132 kDa) using baculovirus expression system. GST-QIK was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

: AMARA peptide

ATP (μ M) Km app / Bin : 42 / 50 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 2.4 IC50 at 1 mM ATP (nM) : 2.9

Substrate

RET

Product code 08-159

Human RET, cytoplasmic domain [658-1114(end) amino acids of accession number NP_066124.1] was expressed as N-terminal GST-fusion protein(79 kDa) using baculovirus expression system. GST-RET was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CSKtide

ATP (μ M) Km app / Bin : 7.5 / 10 Metal : Ma

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 1.3 IC50 at 1 mM ATP (nM) : 20

RET[G691S]

Product code 08-522

Human RET, cytoplasmic domain [658-1114(end) amino acids and G691S of accession number NP_066124.1] was expressed as N-terminal GST-fusion protein (79 kDa) using baculovirus expression system. GST-RET[G691S] was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CSKtide

ATP (μ M) Km app / Bin : 13 / 10 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 1.1 IC50 at 1 mM ATP (nM) : 24

RET[M918T]

Product code 08-508

Human RET, cytoplasmic domain [658-1114(end) amino acids and M918T of accession number NP_066124.1] was expressed as N-terminal GST-fusion protein (79 kDa) using baculovirus expression system. GST-RET[M918T] was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CSKtide

ATP (μ M) Km app / Bin : 4.2 / 5 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 1.4 IC50 at 1 mM ATP (nM) : 81



RET[S891A]

Product code 08-523

Human RET, cytoplasmic domain [658-1114(end) amino acids and S891A of accession number NP_066124.1] was expressed as N-terminal GST-fusion protein (79 kDa) using baculovirus expression system. GST-RET[S891A] was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CSKtide

ATP (μ M) Km app / Bin : 11 / 10 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.44 IC50 at 1 mM ATP (nM) : 9.6

RET[Y791F]

Product code 08-521

Human RET, cytoplasmic domain [658-1114(end) amino acids and Y791F of accession number NP_066124.1] was expressed as N-terminal GST-fusion protein (79 kDa) using baculovirus expression system. GST-RET[Y791F] was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CSKtide

ATP (μ M) Km app / Bin : 29 / 25 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 1.5 IC50 at 1 mM ATP (nM) : 26

ROCK1

Product code 01-109

Human ROCK1, catalytic domain [1-477 amino acids of accession number NP_005397.1] was expressed as N-terminal GST-fusion protein (82 kDa) using baculovirus expression system. GST-ROCK1 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : LIMKtide

ATP (μ M) Km app / Bin : 3.1 / 5 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 1.6 IC50 at 1 mM ATP (nM) : 73

ROCK2

Product code 01-110

Human ROCK2, catalytic domain [1-553 amino acids of accession number NP_004841.2] was expressed as N-terminal GST-fusion protein (91 kDa) using baculovirus expression system. GST-ROCK2 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : LIMKtide

ATP (μ M) Km app / Bin : 7.4 / 5 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.92 IC50 at 1 mM ATP (nM) : n.a.

RON(MST1R)

Product code 08-152

Human RON, cytoplasmic domain [979-1400(end) amino acids of accession number NP_002438.1] was expressed as N-terminal GST-fusion protein (75kDa) using baculovirus expression system. GST-RON was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 27 / 25 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 37 IC50 at 1 mM ATP (nM) : 550



ROS(ROS1)

Product code 08-163

Human ROS, cytoplasmic domain [1883-2347(end) amino acids of accession number NP_002935.2] was expressed as N-terminal GST-fusion protein (79 kDa) using baculovirus expression system. GST-ROS was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : IRS1

ATP (µM) Km app / Bin : 37 / 50 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) IC50 at 1 mM ATP (nM): 1.0

RSK1(RPS6KA1)

Product code 01-149

Full-length human RSK1 [1-735(end) amino acids of accession number NP_002944.2] was expressed as N-terminal GST-fusion protein (110 kDa) using baculovirus expression system. GST-RSK1 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : S6K peptide(N-FL)

ATP (µM) Km app / Bin : 21 / 25 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.14 IC50 at 1 mM ATP (nM): 2.5

RSK2(RPS6KA3)

Product code 01-150

Full-length human RSK2 [1-740(end) amino acids of accession number NP_004577.1] was expressed as N-terminal GST-fusion protein (111 kDa) using baculovirus expression system. GST-RSK2 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : S6K peptide(N-FL)

ATP (μ M) Km app / Bin : 14 / 10 Metal : Ma

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.10 IC50 at 1 mM ATP (nM): n.a.

RSK3(RPS6KA2)

Product code 01-151

Full-length human RSK3 [1-733(end) amino acids of accession number NP_066958.2] was expressed as N-terminal GST-fusion protein (111 kDa) using baculovirus expression system. GST-RSK3 was purified by using glutathione sepharose chromatography.

: Mobility Shift Assay Assay platform Substrate : S6K peptide(N-FL)

ATP (µM) Km app / Bin : 9.9 / 10

: Mg Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.11 IC50 at 1 mM ATP (nM): 1.7

Metal

RSK4(RPS6KA6)

Product code 01-152

Full-length human RSK4 [1-745(end) amino acids of accession number NP_055311.1] was expressed as N-terminal GST-fusion protein (111 kDa) using baculovirus expression system. GST-RSK4 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay Substrate : S6K peptide(N-FL)

ATP (µM) Km app / Bin : 20 / 25 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.051 IC50 at 1 mM ATP (nM): 0.56



SGK

Product code 01-158

Truncated human SGK [61-431(end) amino acids and S422D of accession number NP_005618.2] was co-expressed as N-terminal GST-fusion protein (68 kDa) with His-tagged PDK1 [1-556(end) amino acids of accession number NP_002604.1] using baculovirus expression system. GST-SGK was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : SGKtide

ATP (μ M) Km app / Bin : 52 / 50 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 10 IC50 at 1 mM ATP (nM) : 99

SGK2

Product code 01-159

Full-length human SGK2 [1-367(end) amino acids and S356D of accession number NP_733794.1] was co-expressed as N-terminal GST-fusion protein (68 kDa) with His-tagged PDK1 [1-556(end) amino acids of accession number NP_002604.1] using baculovirus expression system. GST-SGK2 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : SGKtide

ATP (μ M) Km app / Bin : 58 / 50 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 30 IC50 at 1 mM ATP (nM) : n.a.

SGK3(SGKL)

Product code 01-160

Truncated human SGK3 [119-496(end) amino acids and S486D of accession number NP_037389.4] was co-expressed as N-terminal GST-fusion protein (68 kDa) with His-tagged PDK1 [1-556(end) amino acids of accession number NP_002604.1] using baculovirus expression system. GST-SGK3 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : SGKtide

ATP (μ M) Km app / Bin : 17 / 25 Metal : Ma

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 42 IC50 at 1 mM ATP (nM) : n.a.

SIK(SNF1LK)

Product code 02-131

Full-length human SIK [1-783(end) amino acids of accession number NP_775490.2] was expressed as N-terminal GST-fusion protein (112 kDa) using baculovirus expression system. GST-SIK was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay Substrate : AMARA peptide

ATP (μ M) Km app / Bin : 47 / 50 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 1.8 IC50 at 1 mM ATP (nM) : 1.0

skMLCK(MYLK2)

Product code 02-150

Full-length human skMLCK [1-596(end) amino acids of accession number NP_149109.1] was expressed as N-terminal GST-fusion protein (93 kDa) using baculovirus expression system. GST-skMLCK was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : MLCtide

ATP (µM) Km app / Bin : 820 / 1000

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 51 IC50 at 1 mM ATP (nM) : 51



SLK

Product code 07-129

Full-length human SLK [1-1152(end) amino acids and S5N of accession number NP_055535.1] was expressed as N-terminal GST-fusion protein (160 kDa) using baculovirus expression system. GST-SLK was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Moesin-derived peptide

ATP (μ M) Km app / Bin : 36 / 50 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.32 IC50 at 1 mM ATP (nM) : n.a.

SPHK1

Product code 11-105

Full-length human SPHK1 [1-384(end) amino acids of accession number NP_001136074.1] was expressed as N-terminal GST-fusion protein (69 kDa) using baculovirus expression system. GST-SPHK1 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Sphingosine

IC50 at ATP Bin (nM) : 3.9 IC50 at 1 mM ATP (nM) : n.a.

SPHK2

Product code 11-106

Full-length human SPHK2 [1-618(end) amino acids of accession number NP_001191089.1] was expressed as N-terminal GST-fusion protein (92 kDa) using baculovirus expression system. GST-SPHK2 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Sphingosine

ATP (μ M) Km app / Bin : 620 / 600

Metal : Mg
Reference compound : PF-543
IC50 at ATP Bin (nM) : 400
IC50 at 1 mM ATP (nM) : n.a.

SRC

Product code 08-173

Full-length human SRC [1-536(end) amino acids of accession number NP_005408.1] was expressed as N-terminal GST-fusion protein (87 kDa) using baculovirus expression system. GST-SRC was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 31 / 50 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 5.3 IC50 at 1 mM ATP (nM) : 33

SRM(SRMS)

Product code 08-174

Human SRM, catalytic domain [215-488(end) amino acids of accession number NP_543013.1] was expressed as N-terminal GST-fusion protein (58kDa) using baculovirus expression system. GST-SRM was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Blk/Lyntide

ATP (μ M) Km app / Bin : 38 / 50 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 290 IC50 at 1 mM ATP (nM) : 5000



SRPK1

Product code 04-160

Full-length human SRPK1 [1-655(end) amino acids and V564 deletion of accession number NP_003128.3] was expressed as Nterminal GST-fusion protein (101 kDa) using E. coli expression system. GST-SRPK1 was purified by using glutathione sepharose chromatography.

Assay platform : IMAP

Substrate : SRPKtide

ATP (µM) Km app / Bin : 200 / 100

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) IC50 at 1 mM ATP (nM): n.a.

SRPK2

Product code 04-161

Full-length human SRPK2 [1-688(end) amino acids of accession number NP 872633.11 was expressed as N-terminal GST-fusion protein (104 kDa) using baculovirus expression system. GST-SRPK2 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

: Mg

Substrate : DYRKtide-F

ATP (µM) Km app / Bin : 14 / 10 Metal

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 600 IC50 at 1 mM ATP (nM): n.a.

SYK

Product code 08-176

Full-length human SYK [1-635(end) amino acids of accession number NP_003168.2] was expressed as N-terminal GST-fusion protein (99 kDa) using baculovirus expression system. GST-SYK was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

: Blk/Lyntide Substrate

ATP (µM) Km app / Bin : 59 / 50 Metal : Ma

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.29 IC50 at 1 mM ATP (nM): 0.63

TAOK2

Product code 07-133

Human TAOK2, catalytic domain [1-319 amino acid of accession number NP_004774.1] was expressed as N-terminal GST-fusion protein (63 kDa) using baculovirus expression system, GST-TAOK2 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : TAOKtide

ATP (μM) Km app / Bin : 39 / 50

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) IC50 at 1 mM ATP (nM): n.a.

TBK1

Product code 05-115

Full-length human TBK1 [1-729(end) amino acids of accession number NP_037386.1] was expressed as N-terminal GST-fusion protein (111 kDa) using baculovirus expression system. GST-TBK1 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CKtide

ATP (µM) Km app / Bin : 21 / 25 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 1.2 IC50 at 1 mM ATP (nM): n.a.



TEC

Product code 08-182

Human TEC, catalytic domain [359-631 amino acids of accession number AAI01712.1] was expressed as N-terminal GST-fusion protein (59 kDa) using baculovirus expression system. GST-TEC was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 55 / 50 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 20 IC50 at 1 mM ATP (nM) : 220

TIE2(TEK)

Product code 08-185

Human TIE2, cytoplasmic domain [771-1124(end) amino acids of accession number NP_000450.1] was expressed as N-terminal GST-fusion protein (68 kDa) using baculovirus expression system. GST-TIE2 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Blk/Lyntide

ATP (µM) Km app / Bin : 94 / 100

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 100 IC50 at 1 mM ATP (nM) : 190

TNIK

Product code 07-138

Human TNIK, catalytic domain [1-314 amino acids of accession number NP_055843.1] was expressed as N-terminal GST-fusion protein (62 kDa) using baculovirus expression system. GST-TNIK was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay
Substrate : Moesin-derived peptide

ATP (μM) Km app / Bin : 16 / 25 Metal : Mα

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 1.0 IC50 at 1 mM ATP (nM) : 11

TNK₁

Product code 08-104

Human TNK1, catalytic domain [106-390 amino acids of accession number Q13470-2] was expressed as N-terminal GST-fusion protein (58 kDa) using baculovirus expression system. GST-TNK1 was purified by using glutathione sepharose chromatography and gel filtration chromatography.

Assay platform : Mobility Shift Assay

Substrate : CSKtide

ATP (μ M) Km app / Bin : 71 / 75 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.55 IC50 at 1 mM ATP (nM) : 1.7

TRKA(NTRK1)

Product code 08-186

Human TRKA, cytoplasmic domain [436-790(end) amino acids of accession number NP_001012331.1] was expressed as N- terminal GST-fusion protein (67 kDa) using baculovirus expression system. GST-TRKA was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CSKtide

ATP (μ M) Km app / Bin : 65 / 75 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.34 IC50 at 1 mM ATP (nM) : 0.64



TRKB(NTRK2)

Product code 08-187

Human TRKB, cytoplasmic domain [456-822(end) amino acids of accession number NP_001018074.1] was expressed as N- terminal GST-fusion protein (69 kDa) using baculovirus expression system. GST-TRKB was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 80 / 75 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.29 IC50 at 1 mM ATP (nM) : 0.55

TRKC(NTRK3)

Product code 08-197

Human TRKC, cytoplasmic domain [456-825(end) amino acids of accession number NP_002521.2] was expressed as N-terminal GST-fusion protein (69 kDa) using baculovirus expression system. GST-TRKC was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 47 / 50 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.32 IC50 at 1 mM ATP (nM) : 1.0

TSSK1

Product code 02-364

Full-length human TSSK1 [1-367(end) amino acids of accession number NP_114417.1] was expressed as N-terminal GST-fusion protein using baculovirus expression system. GST-TSSK1 was purified by using glutathione sepharose chromatography. GST-TSSK1 was cleaved by PreScission protease and GST-free TSSK1 (42 kDa) was collected as flow-through fraction from glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : GS peptide

ATP (μ M) Km app / Bin : 11 / 10 Metal : Ma

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.19 IC50 at 1 mM ATP (nM) : 0.95

TSSK2

Product code 02-165

Full-length human TSSK2 [1-358(end) amino acids of accession number NP_443732.3] was expressed as N-terminal GST-fusion protein (68 kDa) using baculovirus expression system. GST-TSSK2 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : GS peptide

ATP (μM) Km app / Bin : 8.8 / 10

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 4.7 IC50 at 1 mM ATP (nM) : n.a.

TSSK3

Product code 02-166

Full-length human TSSK3 [2-268(end) amino acids of accession number NP_443073.1] was expressed as N-terminal GST-fusion protein (57 kDa) using baculovirus expression system. GST-TSSK3 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : GS peptide

ATP (μ M) Km app / Bin : 45 / 50 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 12 IC50 at 1 mM ATP (nM) : n.a.



TXK

Product code 08-183

Human TXK, catalytic domain [260-527(end) amino acids of accession number NP_003319.1] was expressed as N-terminal GST-fusion protein (58 kDa) using baculovirus expression system. GST-TXK was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 110 / 100

Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 45 IC50 at 1 mM ATP (nM) : 220

TYK2

Product code 08-147

Human TYK2, catalytic domain [871-1187(end) amino acids of accession number NP_003322.3] was expressed as N-terminal GST-fusion protein (63 kDa) using baculovirus expression system. GST-TYK2 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 18 / 25 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 1.0 IC50 at 1 mM ATP (nM) : 7.0

TYRO3

Product code 08-109

Human TYRO3, cytoplasmic domain of [453-890(end) amino acids of accession number NP_006284.2] was expressed as N-terminal GST fusion protein (76 kDa) using baculovirus expression system. GST-TYRO3 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : CSKtide

ATP (μ M) Km app / Bin : 80 / 75 Metal : Ma

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 1.3 IC50 at 1 mM ATP (nM) : 2.9

WNK1

Product code 05-179

Human WNK1, catalytic domain [1-491 amino acids of accession number NP_061852.1] was expressed as N-terminal GST-fusion protein (81 kDa) using baculovirus expression system. GST-WNK1 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : SPAKtide

WNK2

Product code 05-180

Human WNK2, catalytic domain [166-489 amino acids of accession number NP_006639.3] was expressed as N-terminal GST-fusion protein (65 kDa) using baculovirus expression system. GST-WNK2 was purified by using glutathione sepharose chromatography and anion exchange chromatography.

Assay platform : Mobility Shift Assay

Substrate : SPAKtide

IC50 at 1 mM ATP (nM): n.a.

ATP (µM) Km app / Bin : 48 / 50 Metal : Mg+Mn Reference compound : K252a IC50 at ATP Bin (nM) : 2300 IC50 at 1 mM ATP (nM) : n.a.



WNK3

Product code 05-181

Human WNK3, catalytic domain [1-434 amino acids of accession number NP_065973.2] was expressed as N-terminal GST-fusion protein (76 kDa) using baculovirus expression system. GST-WNK3 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : SPAKtide

ATP (µM) Km app / Bin : 48 / 50 Metal : Mg+Mn Reference compound : K252a IC50 at ATP Bin (nM) : 1300

IC50 at 1 mM ATP (nM): n.a.

YES(YES1)

Product code 08-175

Full-length human YES [1-543(end) amino acids of accession number NP_005424.1] was expressed as N-terminal GST-fusion protein (88 kDa) using baculovirus expression system. GST-YES was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Srctide

ATP (μ M) Km app / Bin : 13 / 10 Metal : Mg

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 2.4 IC50 at 1 mM ATP (nM) : 23

YES(YES1)[T348I]

Product code 08-533

Full-length human YES [1-543(end) amino acids and T348I of accession number NP_005424.1] was expressed as N-terminal GST-fusion protein (89 kDa) using baculovirus expression system. GST-YES[T348I] was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

: Staurosporine

Substrate : Srctide

ATP (µM) Km app / Bin : 8.5 / 10

Metal : Mg

IC50 at ATP Bin (nM) : 1.4 IC50 at 1 mM ATP (nM) : 45

Reference compound

ZAP70

Product code 08-177

Full-length human ZAP70 [1-619(end) amino acids of accession number NP_001070] was expressed as N-terminal GST-fusion protein (97 kDa) using baculovirus expression system. GST-ZAP70 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Blk/Lyntide

ATP (μ M) Km app / Bin : 3.3 / 5 Metal : Mg+Mn

Reference compound : Staurosporine

IC50 at ATP Bin (nM) : 0.76 IC50 at 1 mM ATP (nM) : 34



BRAF

Product code 09-122

Human BRAF, catalytic domain [433-726 amino acid of accession number NP_004324.2] was expressed as N-terminal GST-fusion protein (60 kDa) using baculovirus expression system. GST-BRAF was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : MAP2K1

Cascade Assay*

Metal : Mg

Reference compound : ZM336372 IC50 at 1 mM ATP (nM) : >10000

*MAP2K1/Erk2/Modified Erktide

BRAF[V600E]

Product code 09-144

Human BRAF, catalytic domain [433-726 amino acids and V600E of accession number NP_004324.2] was expressed as N-terminal GST-fusion protein (60 kDa) using baculovirus expression system. GST-BRAF[V600E] was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : MAP2K1

Cascade Assay*

Metal : Mg

Reference compound : ZM336372

IC50 at 1 mM ATP (nM): 662

*MAP2K1/Erk2/Modified Erktide

COT(MAP3K8)

Product code 07-301

Human COT, catalytic domain [30-397 amino acids of accession number NP_005195.2] was expressed as N-terminal GST-fusion protein using baculovirus expression system. GST-COT was purified by using glutathione sepharose chromatography. GST-COT was cleaved by PreScission protease and GST-free COT (42 kDa) was collected as flow-through fraction from glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : MAP2K1

Cascade Assay*

Metal : Mg

Reference compound : Staurosporine

IC50 at 1 mM ATP (nM): 120

*MAP2K1/Erk2/Modified Erktide

DLK(MAP3K12)

Product code 09-111

Human DLK, catalytic domain [1-520 amino acid of accession number NP_006292.3] was expressed as N-terminal GST-fusion protein (86 kDa) using baculovirus expression system. GST-DLK was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay
Substrate : MAP2K4/MAP2K7

Cascade Assay*

Metal : Mg

Reference compound : Staurosporine

IC50 at 1 mM ATP (nM): 460

*(MAP2K4/MAP2K7)/JNK2/Modified Erktide

MAP2K1

Product code 07-141

Full-length human MAP2K1 [1-393(end) amino acids of accession number NP_002746.1] was co-expressed as N-terminal GST-fusion protein (71 kDa) with human His-tagged RAF1 [306-648(end) amino acids and Y340D and Y341D of accession number NP_002871.1] using baculovirus expression system. GST-MAP2K1 was purified by using glutathione sepharose chromatography and Ni-NTA affinity chromatography.

Assay platform : Mobility Shift Assay

Substrate : Erk2

Cascade Assay*

Metal : Mg

Reference compound : Staurosporine

IC50 at 1 mM ATP (nM): 58

*Erk2/Modified Erktide

67



MAP2K2

Product code 07-142

Full-length human MAP2K2 [1-400(end) amino acids of accession number NP_109587.1] was co-expressed as N-terminal GST-fusion protein (71 kDa) with human His-tagged RAF1 [306-648(end) amino acids and Y340D and Y341D of accession number NP_002871.1] using baculovirus expression system. GST-MAP2K2 was purified by using glutathione sepharose chromatography and Ni-NTA affinity chromatography.

Assay platform : Mobility Shift Assay

Substrate : Erk2

Cascade Assay*

Metal : Mg

Reference compound : Staurosporine

IC50 at 1 mM ATP (nM): 54

*Erk2/Modified Erktide

MAP2K3

Product code 07-143

Full-length human MAP2K3 [1-347(end) amino acids of accession number NP_659731.1] was co-expressed as N-terminal GST-fusion protein (67 kDa) with human His-tagged MLK3 [99-398 amino acids of accession number NP_002410.1] using baculovirus expression system. GST-MAP2K3 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : p38α

Cascade Assay*

Metal : Mg

Reference compound : Staurosporine

IC50 at 1 mM ATP (nM): 790

MAP2K4

Product code 07-144

Full-length human MAP2K4 [1-399(end) amino acids of accession number NP_003001.1] was co-expressed as N-terminal GST-fusion protein (71 kDa) with human His-tagged MAP3K3 [1-626(end) amino acids of accession number NP_002392.2] using baculovirus expression system. GST-MAP2K4 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : JNK2

Cascade Assay*

Metal : Mg

Reference compound : Staurosporine

IC50 at 1 mM ATP (nM): 4600

MAP2K5

Product code 07-145

Full-length human MAP2K5 [1-448(end) amino acids of accession number NP_660143.1] was co-expressed as N-terminal GST-fusion protein (77 kDa) with human His-tagged MAP3K3[1-626(end) amino acids of accession number NP_002392.2], CDC37 and HSP90 using baculovirus expression system. GST-MAP2K5 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : Erk5

Cascade Assay*

Metal : Mg

Reference compound : Staurosporine

IC50 at 1 mM ATP (nM): 62

MAP2K6

Product code 07-146

Full-length human MAP2K6 [1-334(end) amino acids of accession number NP_002749.2] was co-expressed as N-terminal GST-fusion protein (64 kDa) with human His-tagged MLK3 [99-398 amino acids of accession number NP_002410.1] using baculovirus expression system. GST-MAP2K6 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : $p38\alpha$

Cascade Assay*

Metal : Mg

Reference compound : Staurosporine

IC50 at 1 mM ATP (nM): 140

^{*}p38\alpha/Modified Erktide

^{*}JNK2/Modified Erktide

^{*}Erk5/EGFR-derived peptide

^{*}p38a/Modified Erktide



MAP2K7

Product code 07-148

Full-length human MAP2K7 [1-419(end) amino acids of accession number NP_660186.1] was co-expressed as N-terminal GST-fusion protein (75 kDa) with human His-tagged MAP3K3 [1-626(end) amino acids of accession number NP_002392.2] using baculovirus expression system. GST-MAP2K7 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : JNK2

Cascade Assay*

Metal : Mg

Reference compound : Staurosporine

IC50 at 1 mM ATP (nM): 1100

*JNK2/Modified Erktide

MAP3K1

Product code 07-103

Human MAP3K1, catalytic domain [1327-1646(end) amino acids of accession number XP_042066.8] was expressed as N-terminal GST-fusion protein (62 kDa) using baculovirus expression system. GST-MAP3K1 was purified by using glutathione sepharose chromatography and anion exchange chromatography.

Assay platform : Mobility Shift Assay

Substrate : MAP2K1

Cascade Assay*

Metal : Mg

Reference compound : Staurosporine

IC50 at 1 mM ATP (nM): 160

*MAP2K1/Erk2/Modified Erktide

MAP3K2

Product code 07-104

Human MAP3K2, catalytic domain [337-620(end) amino acids of accession number NP_006600.3] was expressed as N-terminal GST-fusion protein (59 kDa) using baculovirus expression system. GST-MAP3K2 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay
Substrate : MAP2K4/MAP2K7

: MAP2K4/MAP2K7 Cascade Assay*

Metal : Mg

Reference compound : Staurosporine

IC50 at 1 mM ATP (nM): 45

*(MAP2K4/MAP2K7)/JNK2/Modified Erktide

MAP3K3

Product code 07-105

Full-length human MAP3K3 [1-626(end) amino acids of accession number NP_002392.2] was expressed as N-terminal GST-fusion protein (98 kDa) using baculovirus expression system. GST-fusion MAP3K3 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : MAP2K6

Cascade Assay*

Metal : Mg

Reference compound : Staurosporine

IC50 at 1 mM ATP (nM): 72

*MAP2K6/p38a/Modified Erktide

MAP3K4

Product code 07-106

Human MAP3K4, catalytic domain [1312-1608(end) amino acids of accession number NP_005913.2] was expressed as N-terminal GST-fusion protein (61 kDa) using baculovirus expression system. GST-MAP3K4 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : MAP2K6

Cascade Assay*

Metal : Mg

Reference compound : Staurosporine

IC50 at 1 mM ATP (nM): 100

*MAP2K6/p38\alpha/Modified Erktide



MAP3K5

Product code 07-107

Human MAP3K5, catalytic domain [654-971 amino acids of accession number NP_005914.1] was expressed as N-terminal GST-tagged protein (62 kDa) using baculovirus expression system. GST-MAP3K5 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : MAP2K6

Cascade Assay*

Metal : Mg

Reference compound : Staurosporine

IC50 at 1 mM ATP (nM): 14

*MAP2K6/p38\alpha/Modified Erktide

MLK1(MAP3K9)

Product code 09-115

Human MLK1, catalytic domain [110-422 amino acids of accession number NP_149132.2] was expressed as N-terminal GST-fusion protein (62kDa) using baculovirus expression system. GST-MLK1 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : MAP2K1

Cascade Assay*

Metal : Mg

Reference compound : Staurosporine

IC50 at 1 mM ATP (nM): 11

*MAP2K1/Erk2/Modified Erktide

MLK2(MAP3K10)

Product code 09-116

Human MLK2, catalytic domain and leucine-zipper domain [75-462 amino acids of accession number NP_002437.2] was expressed as N-terminal GST-fusion protein (71kDa) using baculovirus expression system. GST-MLK2 was purified by using glutathione sepharose chromatography and gel filtration chromatography.

Assay platform : Mobility Shift Assay

Substrate : MAP2K1

Cascade Assay*

Metal : Mg

Reference compound : Staurosporine

IC50 at 1 mM ATP (nM): 45

*MAP2K1/Erk2/Modified Erktide

MLK3(MAP3K11)

Product code 09-017

Human MLK3, catalytic domain [99-398 amino acids of accession number NP_002410.1] was expressed as N-terminal His-tagged protein (37kDa) using baculovirus expression system. His-tagged MLK3 was purified by using Ni-NTA affinity chromatography.

Assay platform : Mobility Shift Assay

Substrate : MAP2K1

Cascade Assay*

Metal : Mg

Reference compound : Staurosporine

IC50 at 1 mM ATP (nM): 4.8

*MAP2K1/Erk2/Modified Erktide

MOS

Product code 05-118

Full-length, human MOS [1-346(end) amino acids of accession number NP_005363.1] was expressed as N-terminal GST-fusion protein (65 kDa) using baculovirus expression system. GST-MOS was purified by using glutathione sepharose chromatography and anion exchange chromatography.

Assay platform : Mobility Shift Assay

Substrate : MAP2K1

Cascade Assay*

Metal : Mg

Reference compound : Staurosporine

IC50 at 1 mM ATP (nM): 32

*MAP2K1/Erk2/Modified Erktide

70



RAF1

Product code 09-125

Human RAF1, catalytic domain [306-648(end) amino acids and Y340D and Y341D of accession number NP_002871.1] was expressed as N-terminal GST-fusion protein (66 kDa) using baculovirus expression system. GST-RAF1 was purified by using glutathione sepharose chromatography.

Assay platform : Mobility Shift Assay

Substrate : MAP2K1

Cascade Assay*

Metal : Mg

Reference compound : ZM336372

IC50 at 1 mM ATP (nM): 2800

*MAP2K1/Erk2/Modified Erktide

TAK1-TAB1(MAP3K7)

Product code 09-019

Fused gene of human TAK1 [1-303 amino acids of accession number NP_663304.1] and human TAB1 [437-504 amino acids of accession number NP_006107.1] was expressed as N-terminal Histagged protein (45kDa) using baculovirus expression system. Histagged TAK1-TAB1 was purified by using Ni-NTA affinity chromatography.

Assay platform : Mobility Shift Assay Substrate : MAP2K4/MAP2K7

Cascade Assay*

Metal : Mg

Reference compound : Staurosporine

IC50 at 1 mM ATP (nM): 340

*(MAP2K4/MAP2K7)/JNK2/Modified Erktide



Assay conditions

Test compounds

The test compound is dissolved in and diluted with dimethylsulfoxide (DMSO) to achieve 100-fold higher concentration which is specified by the sponsor. Then the solution is further 25-fold diluted with assay buffer to make the final test compound solution. Reference compounds for the assay control are prepared similarly.

Assay reagents and procedures

IMAP™ assay

- 1) 4x Substrate/ATP/Metal solution and 2x kinase solution are prepared with assay buffer (20 mM HEPES, 0.01% Tween-20, 2 mM DTT, pH7.4).
- 2) 5 μ L of 4x compound solution, 5 μ L of 4x Substrate/ATP/Metal solution, and 10 μ L of 2x kinase solution are mixed and incubated in a well of polystyrene 384 well black microplate for 1 hour at room temperature.
- 3) 60 μ L of IMAP binding reagent (IMAPTM Screening Express kit; Molecular Devices) is added to the well, and incubated for over 30 minutes.
- 4) The kinase reaction is evaluated by the fluorescence polarization at 485 nm for excitation and 530 nm for emission of the well.

Off-chip Mobility Shift Assay (MSA)

- 1) 4x Substrate/ATP/Metal solution is prepared with kit buffer (20 mM HEPES, 0.01% Triton X-100, 5 mM DTT, pH7.5), and 2x kinase solution is prepared with assay buffer (20 mM HEPES, 0.01% Triton X-100, 1 mM DTT, pH7.5).
- 2) 5 μ L of 4x compound solution, 5 μ L of 4x Substrate/ATP/Metal solution, and 10 μ L of 2x kinase solution are mixed and incubated in a well of polypropylene 384 well microplate for 1 or 5 hour(s)* at room temperature. (*; depend on kinase)
- 3) 70 μ L of Termination Buffer (QuickScout Screening Assist MSA; Carna Biosciences) is added to the well.
- 4) The reaction mixture is applied to LabChipTM system (PerkinElmer), and the product and substrate peptide peaks are separated and quantitated.
- 5) The kinase reaction is evaluated by the product ratio calculated from peak heights of product(P) and substrate(S) peptides (P/(P+S)).

Off-chip Mobility Shift Assay (MSA) with pre-incubation

- 1) 4x Substrate/ATP/Metal solution is prepared with kit buffer (20 mM HEPES, 0.01% Triton X-100, 5 mM DTT, pH7.5), and 2x kinase solution is prepared with assay buffer (20 mM HEPES, 0.01% Triton X-100, 1 mM DTT, pH7.5).
- 2) 5 μ L of 4x compound solution and 10 μ L of 2x kinase solution are mixed and incubated in a well of polypropylene 384 well microplate for 30 minutes at room temperature.
- 3) 5 μ L of 4x Substrate/ATP/Metal solution is added to the well, and incubated for 1 hour at room temperature.
- 4) 70 μ L of Termination Buffer (QuickScout Screening Assist MSA; Carna Biosciences) is added to the well.
- 5) The reaction mixture is applied to LabChipTM system (PerkinElmer), and the product and substrate peptide peaks are separated and quantitated.
- 6) The kinase reaction is evaluated by the product ratio calculated from peak heights of product(P) and substrate(S) peptides (P/(P+S)).



ADP-Glo™ Kinase Assay

- 1) 4x compound solution and 4x ATP solution are prepared with assay buffer (50 mM MOPS, 1 mM DTT, pH7.2). 4x Substrate solution and 4x kinase/Metal solution are prepared with MOPS based buffer containing individual kinase specific additives.
- 2) 5 μ L of 4x compound solution, 5 μ L of 4x Substrate solution, 5 μ L of 4x ATP solution, and 5 μ L of 4x kinase/Metal solution are mixed and incubated in a well of polystyrene 384 well black microplate for 1 hour at room temperature.
- 3) 20 μL of ADP-GloTM Reagent (Promega) is added to the well, and incubated for over 40 minutes.
- 4) 40 μ L of Kinase Detection Reagent (Promega) is added to the well, and incubated for over 40 minutes.
- 5) The kinase reaction is evaluated by the endpoint luminescence of the well.



Reaction conditions

ATP Km Bin

ATT KIII DIII	701 . C	Substrate		ATP	(μΜ)	Me	etal	D. C.
Kinase	Platform	Name	(nM)	Km	Assay	Name	(mM)	Positive control
ABL	MSA	ABLtide	1000	16	25	Mg	5	Staurosporine
ABL[E255K]	MSA	ABLtide	1000	17	25	Mg	5	Staurosporine
ABL[T315I]	MSA	ABLtide	1000	4.0	5	Mg	5	Staurosporine
ACK ¹⁾	MSA	WASP peptide	1000	97	100	Mg	5	Staurosporine
AKT1	MSA	Crosstide	1000	31	50	Mg	5	Staurosporine
AKT2	MSA	Crosstide	1000	110	100	Mg	5	Staurosporine
AKT3	MSA	Crosstide	1000	54	50	Mg	5	Staurosporine
ALK	MSA	Srctide	1000	57	50	Mg	5	Staurosporine
ALK[C1156Y]	MSA	Srctide	1000	64	75	Mg	5	Staurosporine
ALK[F1174L]	MSA	Srctide	1000	49	50	Mg	5	Staurosporine
ALK[G1202R]	MSA	Srctide	1000	31	50	Mg	5	Staurosporine
ALK[G1269A]	MSA	Srctide	1000	27	25	Mg	5	Staurosporine
ALK[L1196M]	MSA	Srctide	1000	57	75	Mg	5	Staurosporine
ALK[R1275Q]	MSA	Srctide	1000	84	100	Mg	5	Staurosporine
ALK[T1151_L1152insT]	MSA	Srctide	1000	110	100	Mg	5	Staurosporine
EML4-ALK ¹⁾	MSA	Srctide	1000	43	50	Mg	5	Staurosporine
NPM1-ALK	MSA	Srctide	1000	57	50	Mg	5	Staurosporine
ΑΜΡΚα1/β1/γ1	MSA	SAMS peptide	1000	130	150	Mg	5	Staurosporine
ΑΜΡΚα2/β1/γ1	MSA	SAMS peptide	1000	100	100	Mg	5	Staurosporine
ARG	MSA	ABLtide	1000	24	25	Mg	5	Staurosporine
AurA	MSA	Kemptide	1000	27	25	Mg	5	Staurosporine
AurA/TPX2 ⁹⁾	MSA	Kemptide	1000	1.7	2	Mg	5	Staurosporine
AurB/INCENP	MSA	Kemptide	1000	16	25	Mg	5	Staurosporine
AurC	MSA	Kemptide	1000	24	25	Mg	5	Staurosporine
AXL	MSA	CSKtide	1000	32	50	Mg	5	Staurosporine
BLK	MSA	Srctide	1000	62	75	Mg	5	Staurosporine
BMX	MSA	Srctide	1000	75	75	Mg	5	Staurosporine
BRK ¹⁾	MSA	Blk/Lyntide	1000	250	250	Mg	5	Staurosporine
BRSK1	MSA	CHKtide	1000	30	25	Mg	5	Staurosporine
BRSK2	MSA	CHKtide	1000	31	50	Mg	5	Staurosporine
BTK	MSA	Srctide	1000	22	25	Mg	5	Staurosporine
BTK[C481S]	MSA	Srctide	1000	27	25	Mg	5	Staurosporine
BUB1/BUB3	MSA	H2A peptide	1000	2.9	5	Mg	5	Staurosporine
$CaMK1\alpha^{1)3)}$	MSA	GS peptide	1000	750	1000	Mg	5	Staurosporine
CaMK1 $\delta^{1/3}$	MSA	Synapsin peptide	1000	11	10	Mg	5	Staurosporine
CaMK2α ³⁾	MSA	GS peptide	1000	33	50	Mg	5	Staurosporine
CaMK2β ³⁾	MSA	GS peptide	1000	19	25	Mg	5	Staurosporine
CaMK2γ ³⁾	MSA	GS peptide	1000	23	25	Mg	5	Staurosporine
$CaMK2\delta^{3)}$	MSA	GS peptide	1000	6.3	5	Mg	5	Staurosporine
CaMK4 ³⁾	MSA	GS peptide	1000	20	25	Mg	5	Staurosporine
CDC2/CycB1	MSA	Modified Histone H1	1000	34	50	Mg	5	Staurosporine
CDC7/ASK ¹⁾	MSA	MCM2 peptide	1000	2.8	5	Mg	10	Staurosporine
CDK2/CycA2	MSA	Modified Histone H1	1000	27	25	Mg	5	Staurosporine
CDK2/CycE1	MSA	Modified Histone H1	1000	130	150	Mg	5	Staurosporine



Kinase	D1-46	Substrate		ATP	(μΜ)	Me	etal	Danisian annual
Killase	Platform	Name	(nM)	Km	Assay	Name	(mM)	Positive control
CDK3/CycE1	MSA	Modified Histone H1	1000	1000	1000	Mg	5	Staurosporine
CDK4/CycD3 ¹⁾	MSA	DYRKtide-F	1000	200	200	Mg	5	Staurosporine
CDK5/p25	MSA	Modified	1000	10	10	Mg	5	Staurosporine
CDK6/CycD3 ¹⁾	MSA	Histone H1 DYRKtide-F	1000	330	300	Mg	5	Staurosporine
CDK7/CycH/MAT1 ¹⁾	MSA	CTD3 peptide	1000	32	50	Mg	5	Staurosporine
CDK9/CycT1 ¹⁾	MSA	CDK9 substrate	1000	9.4	10	Mg	5	Staurosporine
CGK2 ⁴⁾	MSA	Kemptide	1000	24	25	Mg	5	Staurosporine
CHK1	MSA	CHKtide	1000	50	50	Mg	5	Staurosporine
CHK2	MSA	CHKtide	1000	51	50	Mg	5	Staurosporine
CK1α ¹⁾		CKtide	1000				5	5-Iodotubercidin
	MSA			4.1	5	Mg		
CK171	MSA	CKtide	1000	6.3	5	Mg	5	5-Iodotubercidin
CK1γ2	MSA	CKtide	1000	10	10	Mg	5	5-Iodotubercidin
CK1γ3 CK1δ	MSA MSA	CKtide CKtide	1000	7.7	5 10	Mg Mg	5	5-Iodotubercidin 5-Iodotubercidin
CK1e ¹⁾	MSA	CKtide	1000	16	25	Mg	5	5-Iodotubercidin
CK2α1/β	MSA	CK2tide	1000	2.9	5	Mg	5	TBB
CK2α2/β	MSA	CK2tide CK2tide	1000	2.1	5	Mg	5	ТВВ
CLK1	MSA	DYRKtide-F	1000		10	ŭ	5	Staurosporine
				11		Mg		-
CLK2	MSA	DYRKtide-F	1000	140	150	Mg	5	Staurosporine
CLK3	MSA	DYRKtide-F	1000	75	75	Mg	5	Staurosporine
CRIK ¹⁾	MSA	Histone H3 peptide	1000	7.8	10	Mg	5	Staurosporine
CSK ¹⁾	MSA	Srctide	1000	4.8	5	Mg+Mn	5+1	Staurosporine
DAPK1	MSA	DAPK1tide	1000	1.1	1	Mg	5	Staurosporine
DCAMKL2 ¹⁾	MSA	GS peptide	1000	120	150	Mg	5	Staurosporine
DDR1 ¹⁾	MSA	IRS1	1000	94	100	Mg	5	Staurosporine
DDR2 ¹⁾	MSA	IRS1	1000	38	50	Mg	5	Staurosporine
DYRK1A	MSA	DYRKtide-F	1000	16	25	Mg	5	Staurosporine
DYRK1B	MSA	DYRKtide-F	1000	59	50	Mg	5	Staurosporine
DYRK2	MSA	DYRKtide-F	1000	7.7	10	Mg	5	Staurosporine
DYRK3	MSA	DYRKtide-F	1000	6.8	5	Mg	5	Staurosporine
EEF2K ¹⁾³⁾	MSA	EEF2Ktide	1000	12	10	Mg	5	A-484954
EGFR	MSA	Srctide	1000	2.7	5	Mg+Mn	5+1	Staurosporine
EGFR[C797S/L858R]	MSA	Srctide	1000	4.1	5	Mg+Mn	5+1	Staurosporine
EGFR[d746-750]	MSA	Srctide	1000	19	25	Mg+Mn	5+1	Staurosporine
EGFR[d746-750/C797S]	MSA	Srctide	1000	8.2	10	Mg+Mn	5+1	Staurosporine
EGFR[d746-750/T790M]	MSA	Srctide	1000	5.4	5	Mg+Mn	5+1	Staurosporine
EGFR[d746-750/T790M/C797S]	MSA	Srctide	1000	1.8	2	Mg+Mn	5+1	Staurosporine
EGFR[D770_N771insNPG]	MSA	Srctide	1000	2.3	5	Mg+Mn	5+1	Staurosporine
EGFR[L858R]	MSA	Srctide	1000	9.8	10	Mg+Mn	5+1	Staurosporine
EGFR[L861Q]	MSA	Srctide	1000	7.5	10	Mg+Mn	5+1	Staurosporine
EGFR[T790M]	MSA	Srctide	1000	0.90	1	Mg+Mn	5+1	Staurosporine
EGFR[T790M/L858R]	MSA	Srctide	1000	1.9	2	Mg+Mn	5+1	Staurosporine
EPHA1	MSA	Blk/Lyntide	1000	22	25	Mg	5	Staurosporine
EPHA2	MSA	Blk/Lyntide	1000	67	75	Mg	5	Staurosporine
EPHA3	MSA	Blk/Lyntide	1000	170	150	Mg	5	Staurosporine
EPHA4	MSA	Blk/Lyntide	1000	52	50	Mg	5	Staurosporine
EPHA5	MSA	Blk/Lyntide	1000	56	50	Mg	5	Staurosporine
ЕРНА6	MSA	Blk/Lyntide	1000	27	25	Mg	5	Staurosporine



		Substrate		ATP	(μΜ)	Me	Metal		
Kinase	Platform	Name	(nM)	Km	Assay	Name	(mM)	Positive control	
EPHA7	MSA	Blk/Lyntide	1000	58	50	Mg	5	Staurosporine	
EPHA8	MSA	Blk/Lyntide	1000	69	75	Mg	5	Staurosporine	
EPHB1	MSA	Blk/Lyntide	1000	29	25	Mg	5	Staurosporine	
EPHB2	MSA	Blk/Lyntide	1000	86	100	Mg	5	Staurosporine	
EPHB3	MSA	Blk/Lyntide	1000	49	50	Mg	5	Staurosporine	
EPHB4	MSA	Blk/Lyntide	1000	56	50	Mg	5	Staurosporine	
Erk1	MSA	Modified Erktide	1000	34	50	Mg	5	K252a	
Erk2	MSA	Modified Erktide	1000	33	50	Mg	5	K252a	
		EGFR-derived							
Erk5 ¹⁾	MSA	peptide	1000	450	1000	Mg	5	Staurosporine	
FAK ¹⁾	MSA	Blk/Lyntide	1000	25	25	Mg	5	Staurosporine	
FER	MSA	Srctide	1000	26	25	Mg	5	Staurosporine	
FES	MSA	Srctide	1000	43	50	Mg	5	Staurosporine	
FGFR1	MSA	CSKtide	1000	89	100	Mg	5	Staurosporine	
FGFR1[V561M]	MSA	CSKtide	1000	33	50	Mg	5	Staurosporine	
FGFR2	MSA	CSKtide	1000	66	75	Mg	5	Staurosporine	
FGFR2[V564I]	MSA	CSKtide	1000	21	25	Mg	5	Staurosporine	
FGFR3	MSA	CSKtide	1000	43	50	Mg	5	Staurosporine	
FGFR3[K650E]	MSA	CSKtide	1000	41	50	Mg	5	Staurosporine	
FGFR3[K650M]	MSA	CSKtide	1000	17	25	Mg	5	Staurosporine	
FGFR3[V555L]	MSA	CSKtide	1000	29	25	Mg	5	Staurosporine	
FGFR3[V555M]	MSA	CSKtide	1000	37	50	Mg	5	Staurosporine	
FGFR4	MSA	CSKtide	1000	230	250	Mg	5	Staurosporine	
FGFR4[N535K]	MSA	CSKtide	1000	30	25	Mg	5	Staurosporine	
FGFR4[V550E]	MSA	CSKtide	1000	210	200	Mg	5	Staurosporine	
FGFR4[V550L]	MSA	CSKtide	1000	160	150	Mg	5	Staurosporine	
FGR	MSA	Srctide	1000	34	50	Mg	5	Staurosporine	
FLT1	MSA	CSKtide	1000	140	150	Mg	5	Staurosporine	
FLT3	MSA	Srctide	1000	94	100	Mg	5	Staurosporine	
FLT4	MSA	CSKtide	1000	72	75	Mg	5	Staurosporine	
FMS	MSA	Srctide	1000	26	25	Mg	5	Staurosporine	
FRK	MSA	Srctide	1000	62	75	Mg	5	Staurosporine	
FYN[isoform a]	MSA	Srctide	1000	36	50	Mg	5	Staurosporine	
FYN[isoform b]	MSA	Srctide	1000	20	25	Mg	5	Staurosporine	
GSK3α	MSA	CREBtide-p	1000	12	10	Mg	5	Staurosporine	
GSK3β	MSA	CREBtide-p	1000	9.1	10	Mg	5	Staurosporine	
Haspin	MSA	Histone H3 peptide	1000	140	150	Mg	5	Staurosporine	
HCK	MSA	Srctide	1000	11	10	Mg	5	Staurosporine	
HER2	MSA	Srctide	1000	3.5	5	Mn	5	Staurosporine	
HER4	MSA	Srctide	1000	27	25	Mg	5	Staurosporine	
HGK	MSA	Moesin-derived peptide	1000	9.4	10	Mg	5	Staurosporine	
HIPK1	MSA	DYRKtide-F	1000	4.4	5	Mg	5	Staurosporine	
HIPK2	MSA	DYRKtide-F	1000	5.9	5	Mg	5	Staurosporine	
HIPK3	MSA	DYRKtide-F	1000	7.3	5	Mg	5	Staurosporine	
HIPK4	MSA	DYRKtide-F	1000	7.0	5	Mg	5	Staurosporine	
HPK1	MSA	S6K2 peptide	1000	22	25	Mg	1.25	K252a	
IGF1R	MSA	IRS1	1000	63	75	Mg	5	Staurosporine	
ΙΚΚα	IMAP	ΙκΒα peptide	100	41	40	Mg	10	Staurosporine	
ΙΚΚβ	MSA	Modified IκΒα-derived peptide	1000	16	25	Mg	5	Staurosporine	



		Substrate		ATP	(μΜ)	Me	etal	Positive control
Kinase	Platform	Name	(nM)	Km	Assay	Name	(mM)	Positive control
IKKe ¹⁾	MSA	ΙκΒα peptide	1000	9.5	10	Mg	5	Staurosporine
INSR	MSA	IRS1	1000	58	50	Mg	5	Staurosporine
IRAK1	IMAP	SRPKtide	100	27	25	Mg	2.5	Staurosporine
IRAK4 ¹⁾	MSA	IRAK1 peptide	1000	920	1000	Mg	5	Staurosporine
IRR	MSA	IRS1	1000	64	75	Mg	5	Staurosporine
ITK	MSA	Srctide	1000	6.1	10	Mg	5	Staurosporine
JAK1 ¹⁾⁶⁾	MSA	JAK1 substrate peptide	1000	68	75	Mg	5	Staurosporine
JAK2	MSA	Srctide	1000	13	10	Mg	5	Staurosporine
JAK3	MSA	Srctide	1000	3.5	5	Mg	5	Staurosporine
JNK1	MSA	Modified Erktide	1000	29	100	Mg	5	K252a
JNK2	MSA	Modified Erktide	1000	21	50	Mg	5	K252a
JNK3	MSA	Modified Erktide	1000	6.0	25	Mg	5	K252a
KDR	MSA	CSKtide	1000	74	75	Mg	5	Staurosporine
KIT ⁶⁾	MSA	Srctide	1000	370	400	Mg	5	Staurosporine
KIT[D816E] ⁶⁾	MSA	Srctide	1000	40	50	Mg	5	Staurosporine
KIT[D816V] ⁶⁾	MSA	Srctide	1000	14	10	Mg	5	Staurosporine
KIT[D816Y] ⁶⁾	MSA	Srctide	1000	22	25	Mg	5	Staurosporine
KIT[T670I] ⁶⁾	MSA	Srctide	1000	100	100	Mg	5	Staurosporine
KIT[V560G] ⁶⁾	MSA	Srctide	1000	110	250	Mg	5	Staurosporine
KIT[V654A] ⁶⁾	MSA	Srctide	1000	220	250	Mg	5	Staurosporine
LATS2 ¹⁾	MSA	SGKtide	1000	380	400	Mg	5	Staurosporine
LCK	MSA	Srctide	1000	14	10	Mg	5	Staurosporine
LOK ¹⁾	MSA	Moesin-derived peptide	1000	100	100	Mg	5	Staurosporine
LTK	MSA	Srctide	1000	49	50	Mg	5	Staurosporine
LYNa	MSA	Srctide	1000	14	10	Mg	5	Staurosporine
LYNb	MSA	Srctide	1000	18	25	Mg	5	Staurosporine
MAP4K2	MSA	S6K2 peptide	1000	93	100	Mg	5	Staurosporine
MAPKAPK2	MSA	GS peptide	1000	3.6	5	Mg	5	Staurosporine
MAPKAPK3	MSA	GS peptide	1000	13	10	Mg	5	K252a
MAPKAPK5	MSA	GS peptide	1000	12	10	Mg	5	Staurosporine
MARK1	MSA	CHKtide	1000	8.0	10	Mg	5	Staurosporine
MARK2	MSA	CHKtide	1000	8.8	10	Mg	5	Staurosporine
MARK3	MSA	CHKtide	1000	5.0	5	Mg	5	Staurosporine
MARK4	MSA	CHKtide	1000	12	10	Mg	5	Staurosporine
MELK ¹⁾	MSA	GS peptide	1000	38	50	Mg	5	Staurosporine
MER	MSA	CSKtide	1000	36	50	Mg	5	Staurosporine
MET	MSA	Srctide	1000	27	25	Mg	5	Staurosporine
MET[D1228H]	MSA	Srctide	1000	25	25	Mg	5	Staurosporine
MET[M1250T]	MSA	Srctide	1000	17	25	Mg	5	Staurosporine
MET[Y1235D]	MSA	Srctide	1000	71	75	Mg	5	Staurosporine
MINK ¹⁾	MSA	Modified Erktide	1000	16	50	Mg	5	K252a
MNK1	MSA	RS peptide	1000	460	450	Mg	5	Staurosporine
MNK2	MSA	RS peptide	1000	110	100	Mg	5	Staurosporine
MRCKα ¹⁾	MSA	DAPK1tide	1000	0.45	1	Mg	5	Staurosporine
МRСКβ	MSA	DAPK1tide	1000	0.67	1	Mg	5	Staurosporine
MSK1	MSA	Crosstide	1000	13	10	Mg	5	Staurosporine
MSK2 ¹⁾	MSA	Crosstide	1000	40	50	Mg	5	Staurosporine
MSSK1 ¹⁾	MSA	DYRKtide-F	1000	56	50	Mg	5	K252a



Kinase	Platform	Substrate		ATP	(μΜ)	Me	etal	Positive control
	Flatioilli	Name	(nM)	Km	Assay	Name	(mM)	Positive control
MST1 ¹⁾²⁾	MSA	IRS1	1000	50	50	Mg	5	Staurosporine
MST2 ¹⁾⁷⁾	MSA	IRS1	1000	69	75	Mg	5	Staurosporine
MST3 ¹⁾	MSA	Moesin-derived peptide	1000	66	75	Mg	5	Staurosporine
MST4 ¹⁾	MSA	Moesin-derived peptide	1000	76	75	Mg	5	Staurosporine
MUSK ¹⁾	MSA	CSKtide	1000	14	10	Mg+Mn	5+1	Staurosporine
NDR1 ¹⁾	MSA	SGKtide	1000	12	10	Mg	5	Staurosporine
NDR2 ¹⁾	MSA	SGKtide	1000	7.6	10	Mg	5	Staurosporine
NEK1 ¹⁾	MSA	CDK7 peptide	1000	64	75	Mg	5	Staurosporine
NEK2	MSA	CDK7 peptide	1000	65	75	Mg	5	Staurosporine
NEK4	MSA	GS peptide	1000	51	50	Mg	5	Staurosporine
NEK6 ¹⁾	MSA	CDK7 peptide	1000	69	75	Mg	5	PKR Inhibitor
NEK7 ¹⁾	MSA	CDK7 peptide	1000	40	50	Mg	5	PKR Inhibitor
NEK9 ¹⁾	MSA	CDK7 peptide	1000	190	200	Mg	5	Staurosporine
NIM1K	MSA	CHKtide	1000	21	25	Mg	5	Staurosporine
NuaK1	MSA	CHKtide	1000	59	50	Mg	5	Staurosporine
NuaK2	MSA	CHKtide	1000	26	25	Mg	5	Staurosporine
p38α	MSA	Modified Erktide	1000	150	150	Mg	5	SB202190
p38β	MSA	Modified Erktide	1000	63	75	Mg	5	SB202190
p38γ	MSA	Modified Erktide	1000	13	10	Mg	5	Staurosporine
p38δ	MSA	Modified Erktide	1000	5.8	5	Mg	5	Staurosporine
p70S6K	MSA	S6K2 peptide	1000	14	10	Mg	5	Staurosporine
p70S6Kβ	MSA	S6K2 peptide	1000	3.3	5	Mg	5	Staurosporine
PAK1	MSA	LIMKtide	1000	300	300	Mg	5	Staurosporine
PAK2	MSA	DAPK1tide	1000	81	100	Mg	5	Staurosporine
PAK4 ¹⁾	MSA	SGKtide	1000	2.5	5	Mg	5	Staurosporine
PAK5	MSA	DAPK1tide	1000	1.9	1	Mg	5	Staurosporine
PAK6 ¹⁾	MSA	SGKtide	1000	3.7	5	Mg	5	Staurosporine
PASK ¹⁾	MSA	GS peptide	1000	9.7	10	Mg	5	Staurosporine
PBK ¹⁾	MSA	Histone H3 peptide	1000	33	50	Mg	5	Staurosporine
PDGFRα	MSA	CSKtide	1000	28	25	Mg	5	Staurosporine
PDGFRα[D842V]	MSA	CSKtide	1000	21	25	Mg	5	Staurosporine
PDGFRα[T674I] ¹⁾	MSA	CSKtide	1000	11	10	Mg	5	Staurosporine
PDGFRα[V561D]	MSA	CSKtide	1000	35	50	Mg	5	Staurosporine
PDGFRβ	MSA	CSKtide	1000	23	25	Mg	5	Staurosporine
PDHK2 ¹⁾	MSA	PDHKtide	1000	28	25	Mg+K	5+3	DCA
PDHK4 ¹⁾	MSA	PDHKtide	1000	19	25	Mg+K	5+25	DCA
PDK1 ¹⁾⁸⁾	MSA	T308tide	1000	9.6	10	Mg	5	Staurosporine
PEK	IMAP	SRPKtide	100	13	10	Mg	5	Staurosporine
PGK ¹⁾⁴⁾	MSA	Kemptide	1000	8.2	10	Mg	5	Staurosporine
PHKG1 ¹⁾	MSA	GS peptide	1000	71	75	Mg	5	Staurosporine
PHKG2	MSA	GS peptide	1000	8.1	10	Mg	5	Staurosporine
PIK3CA/PIK3R1	ADP-Glo	PI(4,5)P2	10000	89	100	Mg	5	PI-103
PIK3CB/PIK3R1	ADI-Glo	PI(4,5)P2	10000	88	100	Mg	5	PI-103
PIK3CD/PIK3R1	ADP-Glo	PI(4,5)P2	10000	37	50	Mg	5	PI-103
PIKFYVE	ADP-Glo	PI(3)P	10000	36	50		5	AG-183
PIM1	MSA	S6K2 peptide	10000	640	500	Mg	5	Staurosporine
PIM1 PIM2 ¹⁾					500	Mg		_
	MSA	S6K2 peptide	1000	4.0		Mg	5	Staurosporine
PIM3	MSA	S6K2 peptide	1000	130	150	Mg	5	Staurosporine



		Substrate		ATP	(μΜ)	Me	etal	Positive control	
Kinase	Platform	Name	(nM)	Km	Assay	Name	(mM)	Positive control	
PIP4K2A	ADP-Glo	PI(5)P	10000	20	25	Mg	5	AG-183	
PIP4K2B	ADP-Glo	PI(5)P	10000	18	25	Mn	0.25	AG-183	
PIP5K1A	ADP-Glo	PI(4)P	10000	28	25	Mg	5	AG-183	
PIP5K1B	ADP-Glo	PI(4)P	10000	95	100	Mg	5	AG-183	
PIP5K1C	ADP-Glo	PI(4)P	10000	33	50	Mg	5	AG-183	
PIP5KL1	ADP-Glo	PI(4)P	10000	1.0	1	Mg	5	AG-183	
ΡΚΑCα	MSA	Kemptide	1000	2.6	5	Mg	5	Staurosporine	
РКАСβ	MSA	Kemptide	1000	4.7	5	Mg	5	Staurosporine	
PKACγ ¹⁾	MSA	Kemptide	1000	4.5	5	Mg	5	Staurosporine	
PKCα ⁵⁾	MSA	PKC peptide	1000	36	50	Mg+Ca	5+0.05	Staurosporine	
PKCβ1 ⁵⁾	MSA	PKC peptide	1000	79	75	Mg+Ca	5+0.05	Staurosporine	
PKCβ2 ⁵⁾	MSA	PKC peptide	1000	41	50	Mg+Ca	5+0.05	Staurosporine	
PKCγ ⁵⁾	MSA	PKC peptide	1000	74	75	Mg+Ca	5+0.05	Staurosporine	
$PKC\delta^{5)}$	MSA	PKC peptide	1000	26	25	Mg	5	Staurosporine	
PKCε ⁵⁾	MSA	PKC peptide	1000	16	25	Mg	5	Staurosporine	
ΡΚCζ	MSA	PKC peptide	1000	11	10	Mg	5	Staurosporine	
PKCη ⁵⁾	MSA	PKC peptide	1000	36	50	Mg	5	Staurosporine	
$PKC\theta^{5)}$	MSA	PKC peptide	1000	18	25	Mg	5	Staurosporine	
PKCι	MSA	PKC peptide	1000	27	25	Mg	5	Staurosporine	
PKD1	MSA	GS peptide	1000	25	25	Mg	5	Staurosporine	
PKD2	MSA	GS peptide	1000	26	25	Mg	5	Staurosporine	
PKD3	MSA	GS peptide	1000	34	50	Mg	5	Staurosporine	
PKN1	IMAP	S6K peptide	100	19	25	Mg	1	Staurosporine	
PKR	IMAP	SRPKtide	100	13	10	Mg	5	Staurosporine	
PLK1 ¹⁾	MSA	CDC25ctide	1000	5.6	5	Mg	5	GW843682X	
PLK2	IMAP	CHK2 peptide	50	30	30	Mg	10	GW843682X	
PLK3	MSA	CDC25ctide	1000	6.8	5	Mg	5	GW843682X	
PRKX ¹⁾	MSA	Kemptide	1000	20	25	Mg	5	Staurosporine	
PYK2	MSA	Blk/Lyntide	1000	56	50	Mg	5	Staurosporine	
QIK	MSA	AMARA peptide	1000	42	50	Mg	5	Staurosporine	
RET	MSA	CSKtide	1000	7.5	10	Mg	5	Staurosporine	
RET[G691S]	MSA	CSKtide	1000	13	10	Mg	5	Staurosporine	
RET[M918T]	MSA	CSKtide	1000	4.2	5	Mg	5	Staurosporine	
RET[S891A]	MSA	CSKtide	1000	11	10	Mg	5	Staurosporine	
RET[Y791F]	MSA	CSKtide	1000	29	25	Mg	5	Staurosporine	
ROCK1	MSA	LIMKtide	1000	3.1	5	Mg	5	Staurosporine	
ROCK2	MSA	LIMKtide	1000	7.4	5	Mg	5	Staurosporine	
RON	MSA	Srctide	1000	27	25	Mg	5	Staurosporine	
ROS	MSA	IRS1	1000	37	50	Mg	5	Staurosporine	
RSK1	MSA	S6K peptide (N-FL)	1000	21	25	Mg	5	Staurosporine	
RSK2	MSA	S6K peptide (N-FL)	1000	14	10	Mg	5	Staurosporine	
RSK3	MSA	S6K peptide (N-FL)	1000	9.9	10	Mg	5	Staurosporine	
RSK4	MSA	S6K peptide (N-FL)	1000	20	25	Mg	5	Staurosporine	
SGK	MSA	SGKtide	1000	52	50	Mg	5	Staurosporine	
SGK2	MSA	SGKtide	1000	58	50	Mg	5	Staurosporine	
SGK3	MSA	SGKtide	1000	17	25	Mg	5	Staurosporine	
SIK ¹⁾	MSA	AMARA peptide	1000	47	50	Mg	5	Staurosporine	
skMLCK ³⁾	MSA	MLCtide	1000	820	1000	Mg	5	Staurosporine	



17.	DI (C	Substrate		ATP	(μM)	Me	etal	D ''' (1
Kinase	Platform	Name	(nM)	Km	Assay	Name	(mM)	Positive control
SLK ¹⁾	MSA	Moesin-derived peptide	1000	36	50	Mg	5	Staurosporine
SPHK1	MSA	Sphingosine	1000	20	25	Mg	5	PF-543
SPHK2	MSA	Sphingosine	1000	620	600	Mg	5	PF-543
SRC	MSA	Srctide	1000	31	50	Mg	5	Staurosporine
SRM	MSA	Blk/Lyntide	1000	38	50	Mg	5	Staurosporine
SRPK1	IMAP	SRPKtide	100	200	100	Mg	10	Staurosporine
SRPK2 ¹⁾	MSA	DYRKtide-F	1000	14	10	Mg	5	Staurosporine
SYK	MSA	Blk/Lyntide	1000	59	50	Mg	5	Staurosporine
TAOK2 ¹⁾⁷⁾	MSA	TAOKtide	1000	39	50	Mg	5	Staurosporine
TBK1	MSA	CKtide	1000	21	25	Mg	5	Staurosporine
TEC	MSA	Srctide	1000	55	50	Mg	5	Staurosporine
TIE2	MSA	Blk/Lyntide	1000	94	100	Mg	5	Staurosporine
TNIK	MSA	Moesin-derived peptide	1000	16	25	Mg	5	Staurosporine
TNK1 ¹⁾	MSA	CSKtide	1000	71	75	Mg	5	Staurosporine
TRKA	MSA	CSKtide	1000	65	75	Mg	5	Staurosporine
TRKB	MSA	Srctide	1000	80	75	Mg	5	Staurosporine
TRKC	MSA	Srctide	1000	47	50	Mg	5	Staurosporine
TSSK1	MSA	GS peptide	1000	11	10	Mg	5	Staurosporine
TSSK2 ¹⁾	MSA	GS peptide	1000	8.8	10	Mg	5	Staurosporine
TSSK3 ¹⁾	MSA	GS peptide	1000	45	50	Mg	5	Staurosporine
TXK ¹⁾	MSA	Srctide	1000	110	100	Mg	5	Staurosporine
TYK2 ¹⁾	MSA	Srctide	1000	18	25	Mg	5	Staurosporine
TYRO3	MSA	CSKtide	1000	80	75	Mg	5	Staurosporine
WNK1 ¹⁾	MSA	SPAKtide	1000	140	150	Mg+Mn	5+3	K252a
WNK2 ¹⁾	MSA	SPAKtide	1000	48	50	Mg+Mn	5+3	K252a
WNK3 ¹⁾	MSA	SPAKtide	1000	48	50	Mg+Mn	5+3	K252a
YES	MSA	Srctide	1000	13	10	Mg	5	Staurosporine
YES[T348I]	MSA	Srctide	1000	8.5	10	Mg	5	Staurosporine
ZAP70	MSA	Blk/Lyntide	1000	3.3	5	Mg+Mn	5+1	Staurosporine

ATP 1mM

Kinase	Platform	Substrate		ATP	(µM)	Me	etal	Positive control
Kinase	Platform	Name	(nM)	Km	Assay	Name	(mM)	Positive control
ABL	MSA	ABLtide	1000	16	1000	Mg	5	Staurosporine
ABL[E255K]	MSA	ABLtide	1000	17	1000	Mg	5	Staurosporine
ABL[T315I]	MSA	ABLtide	1000	4.0	1000	Mg	5	Staurosporine
ACK ¹⁾	MSA	WASP peptide	1000	97	1000	Mg	5	Staurosporine
AKT1	MSA	Crosstide	1000	31	1000	Mg	5	Staurosporine
ALK	MSA	Srctide	1000	57	1000	Mg	5	Staurosporine
ALK[C1156Y]	MSA	Srctide	1000	64	1000	Mg	5	Staurosporine
ALK[F1174L]	MSA	Srctide	1000	49	1000	Mg	5	Staurosporine
ALK[G1202R]	MSA	Srctide	1000	31	1000	Mg	5	Staurosporine
ALK[G1269A]	MSA	Srctide	1000	27	1000	Mg	5	Staurosporine
ALK[L1196M]	MSA	Srctide	1000	57	1000	Mg	5	Staurosporine
ALK[R1275Q]	MSA	Srctide	1000	84	1000	Mg	5	Staurosporine
ALK[T1151_L1152insT]	MSA	Srctide	1000	110	1000	Mg	5	Staurosporine
EML4-ALK ¹⁾	MSA	Srctide	1000	43	1000	Mg	5	Staurosporine
NPM1-ALK	MSA	Srctide	1000	57	1000	Mg	5	Staurosporine



		Substrate		ATP	(µM)	Me	etal	Positive control
Kinase	Platform	Name	(nM)	Km	Assay	Name	(mM)	Positive control
ΑΜΡΚα1/β1/γ1	MSA	SAMS peptide	1000	130	1000	Mg	5	Staurosporine
ARG	MSA	ABLtide	1000	24	1000	Mg	5	Staurosporine
AurA	MSA	Kemptide	1000	27	1000	Mg	5	Staurosporine
AurB/INCENP	MSA	Kemptide	1000	16	1000	Mg	5	Staurosporine
AurC	MSA	Kemptide	1000	24	1000	Mg	5	Staurosporine
AXL	MSA	CSKtide	1000	32	1000	Mg	5	Staurosporine
BLK	MSA	Srctide	1000	62	1000	Mg	5	Staurosporine
BMX	MSA	Srctide	1000	75	1000	Mg	5	Staurosporine
BRK ¹⁾	MSA	Blk/Lyntide	1000	250	1000	Mg	5	Staurosporine
BRSK1	MSA	CHKtide	1000	30	1000	Mg	5	Staurosporine
BTK	MSA	Srctide	1000	22	1000	Mg	5	Staurosporine
BTK[C481S]	MSA	Srctide	1000	27	1000	Mg	5	Staurosporine
CaMK4 ³⁾	MSA	GS peptide	1000	20	1000	Mg	5	Staurosporine
CDC2/CycB1	MSA	Modified Histone H1	1000	34	1000	Mg	5	Staurosporine
CDC7/ASK ¹⁾	MSA	MCM2 peptide	1000	2.8	1000	Mg	10	Staurosporine
CDK2/CycA2	MSA	Modified Histone H1	1000	27	1000	Mg	5	Staurosporine
CDK2/CycE1	MSA	Modified Histone H1	1000	130	1000	Mg	5	Staurosporine
CDK4/CycD3 ¹⁾	MSA	DYRKtide-F	1000	200	1000	Mg	5	Staurosporine
CDK5/p25	MSA	Modified Histone H1	1000	10	1000	Mg	5	Staurosporine
CDK6/CycD3 ¹⁾	MSA	DYRKtide-F	1000	330	1000	Mg	5	Staurosporine
CDK7/CycH/MAT1 ¹⁾	MSA	CTD3 peptide	1000	32	1000	Mg	5	Staurosporine
CDK9/CycT1 ¹⁾	MSA	CDK9 substrate	1000	9.4	1000	Mg	5	Staurosporine
CHK1	MSA	CHKtide	1000	50	1000	Mg	5	Staurosporine
СНК2	MSA	CHKtide	1000	51	1000	Mg	5	Staurosporine
CK1α ¹⁾	MSA	CKtide	1000	4.1	1000	Mg	5	5-Iodotubercidin
CK1ε ¹⁾	MSA	CKtide	1000	16	1000	Mg	5	5-Iodotubercidin
CK2α1/β	MSA	CK2tide	1000	2.9	1000	Mg	5	TBB
CLK1	MSA	DYRKtide-F	1000	11	1000	Mg	5	Staurosporine
CLK2	MSA	DYRKtide-F	1000	140	1000	Mg	5	Staurosporine
CSK ¹⁾	MSA	Srctide	1000	4.8	1000	Mg+Mn	5+1	Staurosporine
DAPK1	MSA	DAPK1tide	1000	1.1	1000	Mg	5	Staurosporine
DDR1 ¹⁾	MSA	IRS1	1000	94	1000	Mg	5	Staurosporine
DDR2 ¹⁾	MSA	IRS1	1000	38	1000	Mg	5	Staurosporine
DYRK1A	MSA	DYRKtide-F	1000	16	1000	Mg	5	Staurosporine
DYRK1B	MSA	DYRKtide-F	1000	59	1000	Mg	5	Staurosporine
EGFR	MSA	Srctide	1000	2.7	1000	Mg+Mn	5+1	Staurosporine
EGFR[C797S/L858R]	MSA	Srctide	1000	4.1	1000	Mg+Mn	5+1	Staurosporine
EGFR[d746-750]	MSA	Srctide	1000	19	1000	Mg+Mn	5+1	Staurosporine
EGFR[d746-750/C797S]	MSA	Srctide	1000	8.2	1000	Mg+Mn	5+1	Staurosporine
EGFR[d746-750/T790M]	MSA	Srctide	1000	5.4	1000	Mg+Mn	5+1	Staurosporine
EGFR[d746-750/T790M/C797S]	MSA	Srctide	1000	1.8	1000	Mg+Mn	5+1	Staurosporine
EGFR[D770_N771insNPG]	MSA	Srctide	1000	2.3	1000	Mg+Mn	5+1	Staurosporine
EGFR[L858R]	MSA	Srctide	1000	9.8	1000	Mg+Mn	5+1	Staurosporine
EGFR[L861Q]	MSA	Srctide	1000	7.5	1000	Mg+Mn	5+1	Staurosporine
EGFR[T790M]	MSA	Srctide	1000	0.90	1000	Mg+Mn	5+1	Staurosporine
EGFR[T790M/C797S/L858R]	MSA	Srctide	1000	0.85	1000	Mg+Mn	5+1	Staurosporine
EGFR[T790M/L858R]	MSA	Srctide	1000	1.9	1000	Mg+Mn	5+1	Staurosporine
EPHA1	MSA	Blk/Lyntide	1000	22	1000	Mg	5	Staurosporine



	ni â	Substrate		ATP	(μΜ)	Me	etal	
Kinase	Platform	Name	(nM)	Km	Assay	Name	(mM)	Positive control
EPHA2	MSA	Blk/Lyntide	1000	67	1000	Mg	5	Staurosporine
ЕРНА3	MSA	Blk/Lyntide	1000	170	1000	Mg	5	Staurosporine
EPHA4	MSA	Blk/Lyntide	1000	52	1000	Mg	5	Staurosporine
EPHA5	MSA	Blk/Lyntide	1000	56	1000	Mg	5	Staurosporine
EPHA6	MSA	Blk/Lyntide	1000	27	1000	Mg	5	Staurosporine
EPHA7	MSA	Blk/Lyntide	1000	58	1000	Mg	5	Staurosporine
EPHA8	MSA	Blk/Lyntide	1000	69	1000	Mg	5	Staurosporine
EPHB1	MSA	Blk/Lyntide	1000	29	1000	Mg	5	Staurosporine
EPHB2	MSA	Blk/Lyntide	1000	86	1000	Mg	5	Staurosporine
ЕРНВ3	MSA	Blk/Lyntide	1000	49	1000	Mg	5	Staurosporine
EPHB4	MSA	Blk/Lyntide	1000	56	1000	Mg	5	Staurosporine
Erk1	MSA	Modified Erktide	1000	34	1000	Mg	5	K252a
Erk2	MSA	Modified Erktide	1000	33	1000	Mg	5	K252a
FAK ¹⁾	MSA	Blk/Lyntide	1000	25	1000	Mg	5	Staurosporine
FER	MSA	Srctide	1000	26	1000	Mg	5	Staurosporine
FES	MSA	Srctide	1000	43	1000	Mg	5	Staurosporine
FGFR1	MSA	CSKtide	1000	89	1000	Mg	5	Staurosporine
FGFR1[V561M]	MSA	CSKtide	1000	33	1000	Mg	5	Staurosporine
FGFR2	MSA	CSKtide	1000	66	1000	Mg	5	Staurosporine
FGFR2[V564I]	MSA	CSKtide	1000	21	1000	Mg	5	Staurosporine
FGFR3	MSA	CSKtide	1000	43	1000	Mg	5	Staurosporine
FGFR3[K650E]	MSA	CSKtide	1000	41	1000	Mg	5	Staurosporine
FGFR3[K650M]	MSA	CSKtide	1000	17	1000	Mg	5	Staurosporine
FGFR3[V555L]	MSA	CSKtide	1000	29	1000	Mg	5	Staurosporine
FGFR3[V555M]	MSA	CSKtide	1000	37	1000	Mg	5	Staurosporine
FGFR4	MSA	CSKtide	1000	230	1000	Mg	5	Staurosporine
FGFR4[N535K]	MSA	CSKtide	1000	30	1000	Mg	5	Staurosporine
FGFR4[V550E]	MSA	CSKtide	1000	210	1000	Mg	5	Staurosporine
FGFR4[V550L]	MSA	CSKtide	1000	160	1000	Mg	5	Staurosporine
FGR	MSA	Srctide	1000	34	1000	Mg	5	Staurosporine
FLT1	MSA	CSKtide	1000	140	1000	Mg	5	Staurosporine
FLT3	MSA	Srctide	1000	94	1000	Mg	5	Staurosporine
FLT4	MSA	CSKtide	1000	72	1000	Mg	5	Staurosporine
FMS	MSA	Srctide	1000	26	1000		5	Staurosporine
FRK	MSA	Srctide	1000	62	1000	Mg Mg	5	Staurosporine
FYN[isoform a]	MSA	Srctide	1000	36	1000		5	Staurosporine
FYN[isoform b]	MSA	Srctide	1000	20	1000	Mg Mg	5	Staurosporine
GSK3α	MSA	CREBtide-p	1000	12	1000		5	Staurosporine
GSK3β	MSA		1000	9.1		Mg	5	Staurosporine
•	-	CREBtide-p			1000	Mg		_
HCK	MSA	Srctide	1000	11	1000	Mg	5	Staurosporine
HER2	MSA	Srctide	1000	3.5	1000	Mn	5	Staurosporine
HER4	MSA	Srctide Moesin-derived	1000	27	1000	Mg	5	Staurosporine
HGK	MSA	peptide	1000	9.4	1000	Mg	5	Staurosporine
HIPK4	MSA	DYRKtide-F	1000	7.0	1000	Mg	5	Staurosporine
IGF1R	MSA	IRS1	1000	63	1000	Mg	5	Staurosporine
ІККВ	MSA	Modified IκBα-derived	1000	16	1000	Mg	5	Staurosporine
		pentide						
INSR	MSA	peptide IRS1	1000	58	1000	Mg	5	Staurosporine



Vinces	Dlotform	Substrate		ATP	(µM)	Me	etal	Positive control	
Kinase	Platform	Name	(nM)	Km	Assay	Name	(mM)	Positive control	
ITK	MSA	Srctide	1000	6.1	1000	Mg	5	Staurosporine	
JAK1 ¹⁾⁶⁾	MSA	JAK1 substrate peptide	1000	68	1000	Mg	5	Staurosporine	
JAK2	MSA	Srctide	1000	13	1000	Mg	5	Staurosporine	
JAK3	MSA	Srctide	1000	3.5	1000	Mg	5	Staurosporine	
JNK1	MSA	Modified Erktide	1000	29	1000	Mg	5	K252a	
JNK2	MSA	Modified Erktide	1000	21	1000	Mg	5	K252a	
JNK3	MSA	Modified Erktide	1000	6.0	1000	Mg	5	K252a	
KDR	MSA	CSKtide	1000	74	1000	Mg	5	Staurosporine	
KIT ⁶⁾	MSA	Srctide	1000	370	1000	Mg	5	Staurosporine	
KIT[D816E] ⁶⁾	MSA	Srctide	1000	40	1000	Mg	5	Staurosporine	
KIT[D816V] ⁶⁾	MSA	Srctide	1000	14	1000	Mg	5	Staurosporine	
KIT[D816Y] ⁶⁾	MSA	Srctide	1000	22	1000	Mg	5	Staurosporine	
KIT[T670I] ⁶⁾	MSA	Srctide	1000	100	1000	Mg	5	Staurosporine	
KIT[V560G] ⁶⁾	MSA	Srctide	1000	110	1000	Mg	5	Staurosporine	
KIT[V654A] ⁶⁾	MSA	Srctide	1000	220	1000	Mg	5	Staurosporine	
LCK	MSA	Srctide	1000	14	1000	Mg	5	Staurosporine	
LTK	MSA	Srctide	1000	49	1000	Mg	5	Staurosporine	
LYNa	MSA	Srctide	1000	14	1000	Mg	5	Staurosporine	
LYNb	MSA	Srctide	1000	18	1000	Mg	5	Staurosporine	
MAPKAPK2	MSA	GS peptide	1000	3.6	1000	Mg	5	Staurosporine	
MER	MSA	CSKtide	1000	36	1000	Mg	5	Staurosporine	
MET	MSA	Srctide	1000	27	1000	Mg	5	Staurosporine	
MET[D1228H]	MSA	Srctide	1000	25	1000	Mg	5	Staurosporine	
MET[M1250T]	MSA	Srctide	1000	17	1000	Mg	5	Staurosporine	
MET[Y1235D]	MSA	Srctide	1000	71	1000	Mg	5	Staurosporine	
MINK ¹⁾	MSA	Modified Erktide	1000	16	1000	Mg	5	K252a	
MST1 ¹⁾²⁾	MSA	IRS1	1000	50	1000	Mg	5	Staurosporine	
MUSK ¹⁾	MSA	CSKtide	1000	14	1000	Mg+Mn	5+1	Staurosporine	
NEK1 ¹⁾	MSA	CDK7 peptide	1000	64	1000	Mg	5	Staurosporine	
NEK2	MSA	CDK7 peptide	1000	65	1000	Mg	5	Staurosporine	
NEK6 ¹⁾	MSA	CDK7 peptide	1000	69	1000	Mg	5	PKR Inhibitor	
NEK7 ¹⁾	MSA	CDK7 peptide	1000	40	1000	Mg	5	PKR Inhibitor	
NEK9 ¹⁾	MSA	CDK7 peptide	1000	190	1000	Mg	5	Staurosporine	
p38α	MSA	Modified Erktide	1000	150	1000	Mg	5	SB202190	
р38β	MSA	Modified Erktide	1000	63	1000	Mg	5	SB202190	
р38ү	MSA	Modified Erktide	1000	13	1000	Mg	5	Staurosporine	
р38δ	MSA	Modified Erktide	1000	5.8	1000	Mg	5	Staurosporine	
p70S6K	MSA	S6K2 peptide	1000	14	1000	Mg	5	Staurosporine	
PAK2	MSA	DAPK1tide	1000	81	1000	Mg	5	Staurosporine	
PBK ¹⁾	MSA	Histone H3 peptide	1000	33	1000	Mg	5	Staurosporine	
PDGFRα	MSA	CSKtide	1000	28	1000	Mg	5	Staurosporine	
PDGFRα[D842V]	MSA	CSKtide	1000	21	1000	Mg	5	Staurosporine	
PDGFRα[T674I] ¹⁾	MSA	CSKtide	1000	11	1000	Mg	5	Staurosporine	
PDGFRα[V561D]	MSA	CSKtide	1000	35	1000	Mg	5	Staurosporine	
PDGFRβ	MSA	CSKtide	1000	23	1000	Mg	5	Staurosporine	
PDK1 ¹⁾⁸⁾	MSA	T308tide	1000	9.6	1000	Mg	5	Staurosporine	
PIM1	MSA	S6K2 peptide	1000	640	1000	Mg	5	Staurosporine	
PIM2 ¹⁾	MSA	S6K2 peptide	1000	4.0	1000	Mg	5	Staurosporine	



***	DI (C	Substrate		ATP	(μΜ)	Me	etal	Positive control
Kinase	Platform	Name	(nM)	Km	Assay	Name	(mM)	Positive control
ΡΚΑCα	MSA	Kemptide	1000	2.6	1000	Mg	5	Staurosporine
PKCα ⁵⁾	MSA	PKC peptide	1000	36	1000	Mg+Ca	5+0.05	Staurosporine
PKCε ⁵⁾	MSA	PKC peptide	1000	16	1000	Mg	5	Staurosporine
PKD2	MSA	GS peptide	1000	26	1000	Mg	5	Staurosporine
PLK1 ¹⁾	MSA	CDC25ctide	1000	5.6	1000	Mg	5	GW843682X
PLK3	MSA	CDC25ctide	1000	6.8	1000	Mg	5	GW843682X
PYK2	MSA	Blk/Lyntide	1000	56	1000	Mg	5	Staurosporine
QIK	MSA	AMARA peptide	1000	42	1000	Mg	5	Staurosporine
RET	MSA	CSKtide	1000	7.5	1000	Mg	5	Staurosporine
RET[G691S]	MSA	CSKtide	1000	13	1000	Mg	5	Staurosporine
RET[M918T]	MSA	CSKtide	1000	4.2	1000	Mg	5	Staurosporine
RET[S891A]	MSA	CSKtide	1000	11	1000	Mg	5	Staurosporine
RET[Y791F]	MSA	CSKtide	1000	29	1000	Mg	5	Staurosporine
ROCK1	MSA	LIMKtide	1000	3.1	1000	Mg	5	Staurosporine
RON	MSA	Srctide	1000	27	1000	Mg	5	Staurosporine
ROS	MSA	IRS1	1000	37	1000	Mg	5	Staurosporine
RSK1	MSA	S6K peptide (N-FL)	1000	21	1000	Mg	5	Staurosporine
RSK3	MSA	S6K peptide (N-FL)	1000	9.9	1000	Mg	5	Staurosporine
RSK4	MSA	S6K peptide (N-FL)	1000	20	1000	Mg	5	Staurosporine
SGK	MSA	SGKtide	1000	52	1000	Mg	5	Staurosporine
SIK ¹⁾	MSA	AMARA peptide	1000	47	1000	Mg	5	Staurosporine
SRC	MSA	Srctide	1000	31	1000	Mg	5	Staurosporine
SRM	MSA	Blk/Lyntide	1000	38	1000	Mg	5	Staurosporine
SYK	MSA	Blk/Lyntide	1000	59	1000	Mg	5	Staurosporine
TEC	MSA	Srctide	1000	55	1000	Mg	5	Staurosporine
TIE2	MSA	Blk/Lyntide	1000	94	1000	Mg	5	Staurosporine
TNIK	MSA	Moesin-derived peptide	1000	16	1000	Mg	5	Staurosporine
TNK1 ¹⁾	MSA	CSKtide	1000	71	1000	Mg	5	Staurosporine
TRKA	MSA	CSKtide	1000	65	1000	Mg	5	Staurosporine
TRKB	MSA	Srctide	1000	80	1000	Mg	5	Staurosporine
TRKC	MSA	Srctide	1000	47	1000	Mg	5	Staurosporine
TSSK1	MSA	GS peptide	1000	11	1000	Mg	5	Staurosporine
TXK ¹⁾	MSA	Srctide	1000	110	1000	Mg	5	Staurosporine
TYK2 ¹⁾	MSA	Srctide	1000	18	1000	Mg	5	Staurosporine
TYRO3	MSA	CSKtide	1000	80	1000	Mg	5	Staurosporine
YES	MSA	Srctide	1000	13	1000	Mg	5	Staurosporine
YES[T348I]	MSA	Srctide	1000	8.5	1000	Mg	5	Staurosporine
ZAP70	MSA	Blk/Lyntide	1000	3.3	1000	Mg+Mn	5+1	Staurosporine

- 1) Reaction time is 5 hours.
- 2) Cantharidin is added at the final concentration of 20 μM_{\cdot}
- 3) CaCl₂, Calmodulin are added at the final concentration of 1 mM and 10 µg/ml, respectively.
- 4) cGMP is added at the final concentration of $5 \mu M$.
- 5) Phosphatidylserine and Diacyl Glycerol are added at the final concentration of 50 μ g/mL and 5 μ g/mL, respectively.
- 6) Sodium orthovanadate is added at the final concentration of 25 μM .
- 7) Cantharidin is added at the final concentration of 10 μM .
- 8) PIFtide and Cantharidin are added at the final concentration of 2 μM and 20 μM , respectively.
- 9) TPX2 peptide is added at the final concentration of 200 nM.



Cascade assay

Kinase	Platform	Substrate		ATP (μM)		Metal		D ''' (1
	Platform	Name	(nM)	Km	Assay	Name	(mM)	Positive control
BRAF	MSA	MAP2K1	1	-	1000	Mg	5	ZM336372
BRAF[V600E]	MSA	MAP2K1	1	-	1000	Mg	5	ZM336372
COT	MSA	MAP2K1	1	-	1000	Mg	5	Staurosporine
DLK ¹⁾	MSA	MAP2K4/MAP2K7	0.5/0.5	-	1000	Mg	5	Staurosporine
MAP2K1	MSA	Erk2	2.5	-	1000	Mg	5	Staurosporine
MAP2K2	MSA	Erk2	2.5	-	1000	Mg	5	Staurosporine
MAP2K3	MSA	p38a	10	-	1000	Mg	5	Staurosporine
MAP2K4 ¹⁾	MSA	JNK2	50	-	1000	Mg	5	Staurosporine
MAP2K5 ¹⁾	MSA	Erk5	50	-	1000	Mg	5	Staurosporine
MAP2K6	MSA	p38a	10	-	1000	Mg	5	Staurosporine
MAP2K7 ¹⁾	MSA	JNK2	50	-	1000	Mg	5	Staurosporine
MAP3K1	MSA	MAP2K1	1	-	1000	Mg	5	Staurosporine
MAP3K2 ¹⁾	MSA	MAP2K4/MAP2K7	0.5/0.5	-	1000	Mg	5	Staurosporine
MAP3K3	MSA	MAP2K6	1	-	1000	Mg	5	Staurosporine
MAP3K4	MSA	MAP2K6	1	-	1000	Mg	5	Staurosporine
MAP3K5	MSA	MAP2K6	1	-	1000	Mg	5	Staurosporine
MLK1	MSA	MAP2K1	1	-	1000	Mg	5	Staurosporine
MLK2	MSA	MAP2K1	1	-	1000	Mg	5	Staurosporine
MLK3	MSA	MAP2K1	1	-	1000	Mg	5	Staurosporine
MOS	MSA	MAP2K1	1	-	1000	Mg	5	Staurosporine
RAF1	MSA	MAP2K1	1	ı	1000	Mg	5	ZM336372
TAK1-TAB1 ¹⁾	MSA	MAP2K4/MAP2K7	0.5/0.5	-	1000	Mg	5	Staurosporine

¹⁾ Reaction time is 5 hours.



Substrate information of cascade assay

Kinase	Substrate									
	MAP2K	(nM)	MAPK	(nM)	peptide	(nM)				
BRAF	MAP2K1	1	Erk2	2.5	Modified Erktide	1000				
BRAF[V600E]	MAP2K1	1	Erk2	2.5	Modified Erktide	1000				
COT	MAP2K1	1	Erk2	2.5	Modified Erktide	1000				
DLK	[MAP2K4/MAP2K7]	0.5/0.5	JNK2	50	Modified Erktide	1000				
MAP2K1	-	-	Erk2	2.5	Modified Erktide	1000				
MAP2K2	-	-	Erk2	2.5	Modified Erktide	1000				
MAP2K3	-	-	p38α	10	Modified Erktide	1000				
MAP2K4	-	-	JNK2	50	Modified Erktide	1000				
MAP2K5	-	-	Erk5	50	EGFR-derived peptide	1000				
MAP2K6	-	-	p38α	10	Modified Erktide	1000				
MAP2K7	-	-	JNK2	50	Modified Erktide	1000				
MAP3K1	MAP2K1	1	Erk2	2.5	Modified Erktide	1000				
MAP3K2	[MAP2K4/MAP2K7]	0.5/0.5	JNK2	50	Modified Erktide	1000				
MAP3K3	MAP2K6	1	p38α	10	Modified Erktide	1000				
MAP3K4	MAP2K6	1	p38α	10	Modified Erktide	1000				
MAP3K5	MAP2K6	1	p38α	10	Modified Erktide	1000				
MLK1	MAP2K1	1	Erk2	2.5	Modified Erktide	1000				
MLK2	MAP2K1	1	Erk2	2.5	Modified Erktide	1000				
MLK3	MAP2K1	1	Erk2	2.5	Modified Erktide	1000				
MOS	MAP2K1	1	Erk2	2.5	Modified Erktide	1000				
RAF1	MAP2K1	1	Erk2	2.5	Modified Erktide	1000				
TAK1-TAB1	[MAP2K4/MAP2K7]	0.5/0.5	JNK2	50	Modified Erktide	1000				

Data analysis

The readout value of reaction control (complete reaction mixture) is set as a 0% inhibition, and the readout value of background (Enzyme(-)) is set as a 100% inhibition, then the percent inhibition of each test solution is calculated.

 IC_{50} value is calculated from concentration vs. %Inhibition curves by fitting to a four parameter logistic curve.

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