LIFE SCIENCE



Receptor Tyrosine Kinase (RTK) Inhibitors

RTKs (Receptor tyrosine kinases) are transmembrane glycoproteins present on cell surface that activate various kinds of main cell signaling pathways like MAPK pathway and PI3K path-way. They're important molecules in biological phenomenon to control basic cell activities such as proliferation, differentiation, survival and death. Their excess expression and active mutation have been reported in cancer cells. Numerous specific low molecular-weight drugs and antibodies designed to inhibit their activities have been developed.

EGFR (ErbB) Family Inhibitors

EGFR is a transmembrane protein receptor for members of EGF family (Epidermal Growth Factor family) of extracellular protein ligands; EGFR is a member of ErbB family of receptors which includes EGFR (ErbB1), HER2 (ErbB2), HER3, HER4, Anti-tumor agents, including Dacomitinib (Product No. **D5450**). have been developed as a result of extensive studies of ligand binding to various ErbB receptors.

AEE 788

Butein PD 153035 Hydrochloride BMS-599626 **Curcumin** (Natural) Curcumin (Synthetic) Daphnetin Dacomitinib Genistein Gefitinib Methyl 2,5-Dihydroxycinnamate (= Erbstatin Analog) **Mubritinib** Pelitinib PD 158780 **Tyrphostin AG 1478** Tyrphostin AG 490 **Tyrphostin AG 494 Tyrphostin A23 Tyrphostin AG528** Tyrphostin AG 835 **Tyrphostin RG 13022** Tyrphostin RG 14620 **WHI-P154**

5mg / 25mg [A3271] 100mg / 1g [B3803] 25mg / 100mg [B4945] 10mg / 50mg [B5520] 1g / 25g [C0434] 5g / 25g [C2302] 1g / 5g [**D4001**] 50mg / 250mg [D5450] 100mg / 1g [G0272] 1g / 5g [G0546] 10mg / 100mg [M2520] 25mg / 100mg [M3058] 25mg / 100mg [P2529] 10mg / 50mg [P2567] 25mg [T2944] 20mg / 100mg / 1g [T2962] 20mg / 100mg [A2704] 25mg / 100mg [T3503] 25mg / 100mg [T3504] 25mg / 100mg [T3540] 25mg / 100mg [T3575] 25mg / 100mg [T3576] 10mg / 50mg [W0013]

FGFR Family Inhibitors

FGF (Fibroblast Growth Factor) is a protein that is involved in wide variety of processes related to normal development of various cells including fibroblasts, as well as angiogenesis, wound healing and so on. The mammalian FGFR (FGF receptor) family has four members : FGFR1, FGFR2, FGFR3 and FGFR4.

PD-161570	5mg / 25mg <mark>[P2531]</mark>
PD 173074	10mg / 50mg <mark>[P2474]</mark>
Regorafenib	25mg / 100mg [R0142]
*Regorafenib is a potent multikinase inhibitor, and it also inhibits other RTKs such as VE	EGFR, TIE2, KIT, RET and so on.

PDGFR Family Inhibitors

PDGF (Platelet-Derived Growth Factor) is mainly involved in the regulation of migration and proliferation of mesenchymal cells. PDGRF (PDGF Receptor) family belongs to type III class of RTK receptors, and includes PDGFR, CSF-1R, KIT, FLT3 among others. The over-expression of PDGF and PDGFR is believed to correlate with development of arteriosclerosis and fibrous growth disease.

AG-1296 TCS 359 GW 2580 Imatinib Imatinib Mesylate Regorafenib Tyrphostin A9 Tyrphostin RG 13022

5mg / 25mg [D4601] 25mg / 100mg [D5275] 50mg / 250mg [G0504] 100mg / 1g [I0906] 100mg / 1g [I0936] 25mg / 100mg [R0142] 25mg / 100mg [T3667] 25mg / 100mg [T3575]

VEGFR Family Inhibitors

VEGF (Vascular Endothelial Growth Factor) is signaling protein which plays an essential role in angiogenesis, enhancement of proliferation of endothelial cell and vasculogenesis. It deeply associated with cancer metastasis and inflammatory disease, therefore, VEGF and VEGFR (VEGF Receptor) are very important target molecules for Pharmaceutical R&D.

AEE 788 SKLB1002 NVP-BHG 712 *The inhibitory ability of NVP-BHG 712 against EphB4 is also reported. PD 173074 Regorafenib SU 1498 WHI-P154 5mg / 25mg [A3271] 25mg / 100mg [K0060] 10mg / 50mg [N1099]

10mg / 50mg [**P2474**] 25mg / 100mg [**R0142**] 5mg / 25mg [**U0103**] 10mg / 50mg [**W0013**]

RET Inhibitors

RET (Rearranged During Transfection) protein is activated as a receptor tyrosine kinase by interacting with GDNF (Glial Cell Line Derived Neuotrophic Factor), such interaction activates signaling pathways including PI3K-Akt signaling pathway and MAPK signaling pathway. It has been reported that a certain type of mutation in *RET*, known as 'rearrangement', fuse the *RET* gene with other nearby genes, resulting in a RET protein that contains parts of other proteins which makes it hyperactive oncogene and correlates with development of thyroid gland cancer and lung cancer.

RPI 1 Regorafenib

Other RTKs Inhibitors

NVP-BHG 712 PQ401 SB 431542 *SB 431542 selectively inhibits ALK (Activin Reveptor-Like Kinase) 5, a type1 receptor of TGFβ. SB-505124 10mg / 50mg [N1099]

10mg / 50mg [R0202]

25mg / 100mg [R0142]

10mg / 50mg [**P2479**] 25mg / 100mg [**B4003**]

5mg / 25mg [B6056]

For further information please refer to our website at www.TCIchemicals.com.

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