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## DATASHEET

### 2,5-Dimethyl-celecoxib

#### Product overview

<b>Name</b>	2,5-Dimethyl-celecoxib
<b>Cat No</b>	HB3717
<b>Purity</b>	>99%
<b>Description</b>	Shows no COX-2 inhibitory function. Analog of celecoxib.

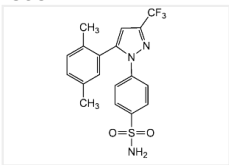
#### Biological Data

<b>Biological description</b>	Close structural analog of the selective cyclooxygenase-2 (COX-2) inhibitor celecoxib, that lacks COX-2 inhibitory function. Anti-proliferative, anti-tumorigenic and anti-angiogenic. Potent apoptosis inducer in many cancer cell lines. Down-regulates the expression of survivin, cyclins, MMPs and inhibits cyclin-dependent kinase activity. Modulates PDK-1, AKT, GSK3beta, p70 S6K, PKA and MAPKAP-K1alpha. Reduces phosphorylation of ERK1/2 but not AKT T-308 or AKT S-473. Increases intracellular free calcium levels and potently triggers the endoplasmic reticulum stress response (ESR), activating ER stress-associated proteins GRP78/BiP, CHOP/GADD153 and caspase-4.
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#### Solubility & Handling

<b>Storage instructions</b>	Room temperature
<b>Solubility overview</b>	Soluble in DMSO
<b>Important</b>	This product is for RESEARCH USE ONLY and is not intended for therapeutic or diagnostic use. Not for human or veterinary use.

#### Chemical Data

<b>Chemical name</b>	4-[5-(2,5-dimethylphenyl)-3-(trifluoromethyl)pyrazol-1-yl]benzenesulfonamide
<b>Molecular Weight</b>	395.4
<b>Chemical structure</b>	
<b>Molecular Formula</b>	C <sub>18</sub> H <sub>16</sub> F <sub>3</sub> N <sub>3</sub> O <sub>2</sub> S
<b>CAS Number</b>	457639-26-8
<b>PubChem identifier</b>	11545682
<b>SMILES</b>	CC1=CC(=C(C=C1)C)C2=CC(=NN2C3=CC=C(C=C3)S(=O)(=O)N)C(F)(F)F
<b>InChi</b>	InChI=1S/C18H16F3N3O2S/c1-11-3-4-12(2)15(9-11)16-10-17(18(19,20)21)23-24(16)13-5-7-14(8-6-13)27(22,25)26/h3-10H,1-2H3,(H2,22,25,26)
<b>InChiKey</b>	NTFOSUUWGCDXEF-UHFFFAOYSA-N
<b>MDL number</b>	MFCD19443858
<b>Appearance</b>	White solid