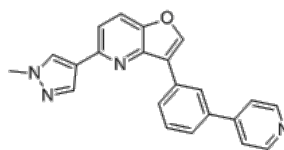


<b>Product Name</b>	: MU1210
<b>Synonyms</b>	: MU-1210; MU 1210
<b>Cat No.</b>	: M13620
<b>CAS Number</b>	: 2275601-87-9
<b>Molecular Formula</b>	: C <sub>22</sub> H <sub>16</sub> N <sub>4</sub> O
<b>Formula Weight</b>	: 352.40
<b>Chemical Name</b>	: 5-(1-methyl-1H-pyrazol-4-yl)-3-(3-(pyridin-4-yl)phenyl)furo[3,2-b]pyridine
<b>Description</b>	<p>MU1210 (MU-1210) is a potent and highly selective inhibitor of cdc-like kinases (CLKs) with IC<sub>50</sub> of 51 nM (cellular BRET for CLK1); profoundly affected the Mdm4 alternative splicing and showed high activity in cytotoxicity assays using MCF7 breast cancer cells; MU1210 has favorable pharmacokinetics (mouse, 10 mpk, IP: C<sub>max</sub> = 1.24 mm, T<sub>1/2</sub> = 58 min; no acute toxicity observed), suggesting its potential use in in vivo models; MU1210 is a quality probe for CLK1, CLK2, and CLK4.</p>
<b>Pathway</b>	: Cell Cycle/DNA Damage
<b>Target</b>	: CLK
<b>Receptor</b>	: CLK
<b>Solubility</b>	: —
<b>SMILES</b>	: —
<b>Storage</b>	: (-20°C)
<b>Stability</b>	: ≥ 2 years
<b>Reference</b>	:



1. Némec V, et al. Angew Chem Int Ed Engl. 2019 Jan 21;58(4):1062-1066.