

<b>Product Name</b>	: ON123300
<b>Synonyms</b>	: ON-123300; ON 123300
<b>Cat No.</b>	: M11467
<b>CAS Number</b>	: 1357470-29-1
<b>Molecular Formula</b>	: C <sub>24</sub> H <sub>27</sub> N <sub>7</sub> O
<b>Formula Weight</b>	: 429.52
<b>Chemical Name</b>	: Pyrido[2,3-d]pyrimidine-6-carbonitrile, 8-cyclopentyl-7,8-dihydro-2-[[4-(4-methyl-1-piperazinyl)phenyl]amino]-7-oxo-
<b>Description</b>	<p>ON123300 is a potent, multi-targeted kinase inhibitor with IC<sub>50</sub> of 5, 3.9, 26, 26, 9.2 and 11 nM for ARK5, CDK4, PDGFRβ, FGFR1, RET and FYN, respectively; inhibits U87 glioma cell proliferation with IC<sub>50</sub> 3.4 uM, and reduces phosphorylation of Akt, yet it also unexpectedly induces Erk activation and prevents phosphorylation of C-Raf S259; suppresses phosphorylation of Akt as well as activation of Erk in brain tumors in vivo; also exhibits potent activity against mantle cell lymphomas both in vitro and in vivo, triggers apoptosis and inhibition of the Rb and PI3K/AKT pathways.</p>
<b>Pathway</b>	: Angiogenesis
<b>Target</b>	: CDK
<b>Receptor</b>	: CDK
<b>Solubility</b>	: 10 mM in DMSO
<b>SMILES</b>	: <chem>N#CC1=CC2=CN=C(NC3=CC=C(N4CCN(C)CC4)C=C3)N=C2N(C5CCCC5)C1=O</chem>
<b>Storage</b>	: (-20°C)
<b>Stability</b>	: ≥ 2 years
<b>Reference</b>	:

1. Zhang X, et al. Mol Cancer Ther. 2014 May;13(5):1105-16. | 2. Divakar SK, et al. Leukemia. 2016 Jan;30(1):86-93. | 3. Perumal D, et al. Cancer Res. 2016 Mar 1;76(5):1225-36.

