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## NLG919 (RG6078)

产品编号: D50917

CAS: 1402836-58-1

分子式: C<sub>18</sub>H<sub>22</sub>N<sub>2</sub>O

纯度: ≥98%

InChi: InChi=1S/C<sub>18</sub>H<sub>22</sub>N<sub>2</sub>O/c21-18(13-6-2-1-3-7-13)10-16-14-8-4-5-9-15(14)17-11-19-12-20(16)17/h4-5,8-9,11-13,16,18,21H,1-3,6-7,10H2

InChi Key: YTRRAUACYORZLX-UHFFFAOYSA-N

Smiles: OC(CC1C2=CC=CC=C2C2=CN=CN21)C1CCCCC1

外观: 固体粉末

作用通路: Indoleamine 2, 3-Dioxygenase (IDO)

溶解性: DMSO up to 50 mM

保存条件: Store in dry, dark place for one year.

产品介绍: NLG919 is a potent, selective and orally bioavailable IDO (indoleamine-(2, 3)-dioxygenase) pathway inhibitor with  $K_i \sim 7$  nM ( $EC_{50} \sim 75$  nM) in vitro and in cell based assays. Oral administration of NLG919 reduces the [Kyn] in plasma and tissue by  $\sim 50\%$ . Using human IDO+ pDCs in allogeneic MLR reactions, NLG919 potently blocked IDO- induced T cell suppression and restored robust T cell responses with an  $EC_{50} \sim 90$  nM. NLG919 abrogated IDO-induced suppression of antigen-specific T cells (OT-I or pmel- 1) in vitro, ( $ED_{50} \sim 130$  nM ) using mouse IDO+ pDCs from tumor-draining lymph nodes. In mice bearing large established B16F10 tumors, administration of NLG919 markedly enhanced the antitumor responses of naive, resting pmel-1 cells to vaccination with cognate hgp100 peptide plus CpG-1826 in IFA. Currently NLG919 is in phase I trial for the treatment of immunosuppression associated with cancer.