NVP-BSK805

Cat. No.: HY-14722
CAS No.: 1092499-93-8
Molecular Formula: \( C_{27}H_{28}F_2N_6O \)
Molecular Weight: 490.55
Target: JAK
Pathway: Epigenetics; JAK/STAT Signaling; Protein Tyrosine Kinase/RTK; Stem Cell/Wnt
Storage:
- Powder: -20°C, 3 years; 4°C, 2 years
- In solvent: -80°C, 6 months; -20°C, 1 month

SOLVENT & SOLUBILITY

In Vitro

DMSO : 100 mg/mL (203.85 mM; Need ultrasonic)

<table>
<thead>
<tr>
<th>Preparing Stock Solutions</th>
<th>1 mg</th>
<th>5 mg</th>
<th>10 mg</th>
</tr>
</thead>
<tbody>
<tr>
<td>Concentration 1 mM</td>
<td>2.0385 mL</td>
<td>10.1926 mL</td>
<td>20.3853 mL</td>
</tr>
<tr>
<td>Concentration 5 mM</td>
<td>0.4077 mL</td>
<td>2.0385 mL</td>
<td>4.0771 mL</td>
</tr>
<tr>
<td>Concentration 10 mM</td>
<td>0.2039 mL</td>
<td>1.0193 mL</td>
<td>2.0385 mL</td>
</tr>
</tbody>
</table>

Please refer to the solubility information to select the appropriate solvent.

BIOLOGICAL ACTIVITY

Description
NVP-BSK805 is an ATP-competitive JAK2 inhibitor, with \( IC_{50} \)s of 0.48 nM, 31.63 nM, 18.68 nM, and 10.76 nM for JAK2 JH1 (JAK homology 1), JAK1 JH1, JAK3 JH1, and TYK2 JH1, respectively\(^1\).

<table>
<thead>
<tr>
<th>IC(_{50}) &amp; Target</th>
<th>JAK2 JH1 0.48 nM (IC(_{50}))</th>
<th>FL JAK2 V617F 0.56 nM (IC(_{50}))</th>
<th>FL JAK2 wt 0.58 nM (IC(_{50}))</th>
<th>TYK2 JH1 10.76 nM (IC(_{50}))</th>
</tr>
</thead>
<tbody>
<tr>
<td>JAK3 JH1 18.68 nM (IC(_{50}))</td>
<td>JAK1 JH1 31.63 nM (IC(_{50}))</td>
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<td></td>
<td></td>
</tr>
</tbody>
</table>

In Vitro
NVP-BSK805 (BSK 805) is a JAK2 inhibitor, with IC\(_{50}\)s of 0.48 nM, 31.63 nM, 18.68 nM, and 10.76 nM for JAK2 JH1 (JAK homology 1), JAK1 JH1, JAK3 JH1, and TYK2 JH1, respectively. NVP-BSK805 inhibits the full-length wild-type JAK2 (FL JAK2 wt) and FL JAK2 V617F activity, with IC\(_{50}\)s of 0.58 ± 0.03 and 0.56 ± 0.04 nM. NVP-BSK805 is ATP-competitive, with calculated \( K_i \) of 0.43 ± 0.02 nM. NVP-BSK805 suppresses the growth of JAK2\(^{V617F}\)-bearing acute myeloid leukemia cell lines with GI\(_{50}\) of <100 nM. NVP-BSK805 blocks the STAT5 phosphorylation at ≥100 nM concentrations, and shows a bias for JAK2 over JAK1 and JAK3 inhibition in the JAK2\(^{V617F}\)-mutant cell lines\(^1\).
NVP-BSK805 (5 μM) improves P-gp inhibitory activity. NVP-BSK805 increases sensitization of drug-resistant KBV20C cancer cells to VIC treatment at 10 μM, and such an effect is more effective than a 5 μM dose[2].

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

In Vivo

NVP-BSK805 (50, 75, and 100 mg/kg, p.o.) also suppresses rhEpo-mediated polycythemia and splenomegaly in BALB/c mice [1].
MCE has not independently confirmed the accuracy of these methods. They are for reference only.

PROTOCOL

Cell Assay [1]
The antiproliferative activity of JAK2 inhibitors is determined by incubating cells for 72 hours (96 hours for MB-02 and MUTZ-8 cells) with an 8-point concentration range of NVP-BSK805 and cell proliferation relative to NVP-BSK805 is measured using the colorimetric WST-1 cell viability readout. Of each triplicate treatment, the mean is calculated and these data are plotted in XLfit 4 to determine the half-maximal growth inhibition (GI50) values [1].
MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Animal Administration [1]
Mice[1]
Concomitantly with NVP-BSK805 treatment, female BALB/c mice receive daily s.c. injections (in 100 μL saline buffer) of 10 units of rhEpo for 4 consecutive days. Controls are injected corresponding volumes of saline buffer. Mice are sacrificed 24 hours after the final dose and total blood, spleen, and bone marrow are taken for further analysis. Animals are 8 to 10 weeks of age at treatment start (20-25 g body weight) and are kept under optimal hygienic conditions with free access to food and water[1].
MCE has not independently confirmed the accuracy of these methods. They are for reference only.

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REFERENCES


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