Roquinimex

Cat. No.: HY-13743
CAS No.: 84088-42-6
Molecular Formula: C₁₈H₁₆N₂O₃
Molecular Weight: 308.33
Target: TNF Receptor
Pathway: Apoptosis
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: ≥ 83.3 mg/mL (270.17 mM)
* “≥” means soluble, but saturation unknown.

Preparing Stock Solutions

<table>
<thead>
<tr>
<th>Concentration</th>
<th>1 mg</th>
<th>5 mg</th>
<th>10 mg</th>
</tr>
</thead>
<tbody>
<tr>
<td>1 mM</td>
<td>3.243 mL</td>
<td>16.2164 mL</td>
<td>32.4328 mL</td>
</tr>
<tr>
<td>5 mM</td>
<td>0.6487 mL</td>
<td>3.2433 mL</td>
<td>6.4866 mL</td>
</tr>
<tr>
<td>10 mM</td>
<td>0.3243 mL</td>
<td>1.6216 mL</td>
<td>3.2433 mL</td>
</tr>
</tbody>
</table>

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

BIOLOGICAL ACTIVITY

Description
Roquinimex (Linomide; PNU212616; ABR212616) is a quinoline derivative immunostimulant which increases NK cell activity and macrophage cytotoxicity; inhibits angiogenesis and reduces the secretion of TNF alpha. IC50 value:

Target: TNF alpha

Prophylactic administration of DSS-treated mice with roquinimex significantly reduced clinical signs of colitis, MDS and the CH-reduction. Moreover, in roquinimex treated animals, the MPO activity was significantly reduced by more than 50% compared to DSS control mice. Notably, therapeutic administration of roquinimex in DSS-treated mice also significantly inhibited the MDS, CH-reduction and MPO activity [2]. Linomide, a synthetic immunomodulator, at concentrations effective in vivo reduces the number of MBP-reactive TNF-alpha and increases MBP-reactive IL-10 and TGF-beta mRNA expressing MNC from MS patients’ blood when analysed in vitro. Compared to dexamethasone, Linomide up-regulated levels of blood MNC expressing mRNA of TGF-beta after culture in presence of MBP [3].

REFERENCES


Caution: Product has not been fully validated for medical applications. For research use only.
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