Neohesperidin dihydrochalcone

**Cat. No.:** HY-N0154  
**CAS No.:** 20702-77-6  
**Molecular Formula:** C_{28}H_{36}O_{15}  
**Molecular Weight:** 612.58  
**Target:** ROS  
**Pathway:** Protein Tyrosine Kinase/RTK  
**Storage:**  
- Powder: -20°C, 3 years; 4°C, 2 years; In solvent: -80°C, 6 months; -20°C, 1 month

**SOLVENT & SOLUBILITY**

<table>
<thead>
<tr>
<th>Solvent &amp; Concentration</th>
<th>1 mg</th>
<th>5 mg</th>
<th>10 mg</th>
</tr>
</thead>
<tbody>
<tr>
<td>1 mM</td>
<td>1.6324 mL</td>
<td>8.1622 mL</td>
<td>16.3244 mL</td>
</tr>
<tr>
<td>5 mM</td>
<td>0.3265 mL</td>
<td>1.6324 mL</td>
<td>3.2649 mL</td>
</tr>
<tr>
<td>10 mM</td>
<td>0.1632 mL</td>
<td>0.8162 mL</td>
<td>1.6324 mL</td>
</tr>
</tbody>
</table>

*“≥” means soluble, but saturation unknown.*

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

**BIOLOGICAL ACTIVITY**

**Description**  
Neohesperidin dihydrochalcone is a synthetic glycoside chalcone, is added to various foods and beverages as a low caloric artificial sweetener.

**In Vitro**  
Neohesperidin dihydrochalcone shows remarkable radical scavenging activity against stable radical and reactive oxygen species (ROS) in concentration dependent manner. Especially, neohesperidin dihydrochalcone is the most potent inhibitor of H_2O_2 and HOCl. Neohesperidin dihydrochalcone shows HOCl scavenging activity of 93.5% and H_2O_2 scavenging property of 73.5%. Neohesperidin dihydrochalcone shows extensive inhibitory effect especially on non-radical ROS H_2O_2 and HOCl with IC_{50} values of 205.1, 25.5 μM[1]. Neohesperidin dihydrochalcone is found to be an activator of porcine pancreatic alpha-amylase (PPA) with an IC_{50} of 389 μM[2].

**In Vivo**  
Neohesperidin dihydrochalcone administration results in significant reduction in activities of two useful markers of liver damage, AST and ALT. The relative levels of NF-κB, IL-6, IL-1β and TNF-α protein in the liver of PQ-treated mice are inhibited by neohesperidin dihydrochalcone[3]. The embryotoxicity/teratogenicity of neohesperidin...
Dihydrochalcone is examined in Wistar Crl:(WI)WU BR rats. No adverse effects are observed at neohesperidin dihydrochalcone levels of up to 5% of the diet, the highest dose level tested, at which the rats consumed about 3.3 g/kg body weight/day.[4]

**PROTOCOL**

**Cell Assay**[1]

WST-8 dye is used in the cell viability assay. HIT-T15 and HUVEC cells are grown and maintained in Dulbecco’s modified Eagle’s medium, supplemented with 10% fetal bovine calf serum. 1000 cells in each well are incubated with various concentrations of neohesperidin dihydrochalcone (50, 100, 500 μM, 1 mM) and other compounds. After treating HIT-T15 and HUVEC cells with 500 μM HOCl, WST-8 dye is added to each well, and the absorbance is detected at 420 nm with microplate reader[1].

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

**Animal Administration**[3][4]

Rats: The embryotoxicity/teratogenicity of neohesperidin dihydrochalcone is examined in Wistar Crl:(WI)WU BR rats. The study is comprised of four groups of 28 mated female rats each, i.e., a control group (0% neohesperidin dihydrochalcone) and three treatment groups (1.25, 2.5, and 5% neohesperidin dihydrochalcone). The general condition and behavior of the animals are observed twice daily. Body weight is determined on days 0, 7, 14, and 21 of gestation. Food consumption is determined during three consecutive periods (days 0-7, 7-14, and 14-21 of gestation)[4].

Mice: Neohesperidin dihydrochalcone is dissolved in a 0.5% CMC vehicle. Mice are randomized into four groups. The control group receives equal volume of vehicles throughout. The PQ group receives saline once daily for 6 consecutive days. One hour after final saline treatment, mice are injected with PQ (75 mg/kg body weight). The neohesperidin dihydrochalcone group receives a daily dose of 200 mg/kg body weight by oral gavage for 6 consecutive days. One hour after final neohesperidin dihydrochalcone treatment, mice are injected with PQ (75 mg/kg body weight).[3]

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

**REFERENCES**


