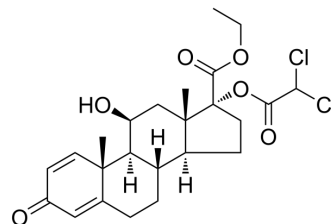


## Etiprednol dicloacetate

<b>Cat. No.:</b>	HY-106215
<b>CAS No.:</b>	199331-40-3
<b>Molecular Formula:</b>	C <sub>24</sub> H <sub>30</sub> Cl <sub>2</sub> O <sub>6</sub>
<b>Molecular Weight:</b>	485.4
<b>Target:</b>	Others
<b>Pathway:</b>	Others
<b>Storage:</b>	Please store the product under the recommended conditions in the Certificate of Analysis.



### BIOLOGICAL ACTIVITY

<b>Description</b>	Etiprednol dicloacetate (BNP 166) is an anti-inflammatory agent. Etiprednol dicloacetate inhibits eosinophil accumulation. Etiprednol dicloacetate can be used in the research of inflammatory airway diseases, such as asthma <sup>[1][2]</sup> .								
<b>In Vitro</b>	Etiprednol dicloacetate (1 nM-1 μM) inhibits LPS-induced TNF-α production by human blood <sup>[1]</sup> . <b>Caution: Product has not been fully validated for medical applications. For research use only.</b> MCE has not independently confirmed the accuracy of these methods. They are for reference only. Tel: 609-228-6898 Fax: 609-228-5909 E-mail: tech@MedChemExpress.com								
<b>In Vivo</b>	<p>Etiprednol dicloacetate (0-100 μg/kg, intranasal inhalation) attenuates peribronchial eosinophilia in the lung of the allergen sensitized and challenged rats<sup>[1]</sup>.</p> <p>Etiprednol dicloacetate (0.1 μg/kg, intranasal inhalation) inhibits eosinophil accumulation in rats (antigen-induced airway eosinophil infiltration)<sup>[3]</sup>.</p> <p>Etiprednol dicloacetate (0-20 mg/kg, p.o., 28 days) is well tolerated in rats and dogs<sup>[2]</sup>.</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <table border="1"> <tr> <td>Animal Model:</td> <td>Allergen sensitized and challenged Brown Norway rats<sup>[1]</sup></td> </tr> <tr> <td>Dosage:</td> <td>0.1, 1, 10, and 100 μg/kg</td> </tr> <tr> <td>Administration:</td> <td>Intranasal inhalation</td> </tr> <tr> <td>Result:</td> <td>Decreased the number of eosinophils at the perivascular area of the lung tissue. Decreased the number of mucus secreting cells and antigen-induced formation of perivascular edema.</td> </tr> </table>	Animal Model:	Allergen sensitized and challenged Brown Norway rats <sup>[1]</sup>	Dosage:	0.1, 1, 10, and 100 μg/kg	Administration:	Intranasal inhalation	Result:	Decreased the number of eosinophils at the perivascular area of the lung tissue. Decreased the number of mucus secreting cells and antigen-induced formation of perivascular edema.
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### REFERENCES

- [1]. István Kurucz, et al. Potency and specificity of the pharmacological action of a new, antiasthmatic, topically administered soft steroid, etiprednol dicloacetate (BNP-166). J Pharmacol Exp Ther. 2003 Oct;307(1):83-92.
- [2]. A Miklós, et al. 28-day oral toxicity study with soft corticosteroid BNP-166 in rats and dogs, followed by a 14-day recovery period. Pharmazie. 2002 Feb;57(2):142-6.
- [3]. I Kurucz, et al. Anti-inflammatory effect and soft properties of etiprednol dicloacetate (BNP-166), a new, anti-asthmatic steroid. Pharmazie. 2004 May;59(5):412-6.