

# Nedaplatin

**Catalog No: tcsc2772** 

Available Sizes

Size: 10mg

Size: 50mg

**Specifications** 

CAS No:

95734-82-0

## Formula:

 $\rm C_2H_8N_2O_3Pt$ 

Pathway: Cell Cycle/DNA Damage

## **Target:**

**DNA/RNA** Synthesis

Purity / Grade:

>98%

## Solubility: H2O : 13.6 mg/mL (44.86 mM; Need ultrasonic and warming)

## Alternative Names:

NSC 375101D

**Observed Molecular Weight:** 

303.18

## **Product Description**

Nedaplatin (NSC 375101D) is a derivative of cisplatin and DNA damage agent.

In Vitro:

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Nedaplatin (NSC 375101D, NDP) is a derivative of cisplatin which produced less nausea & vomiting and nephrotoxicity. the effect of NDP on the 7-ethyl-1-hydroxy-CPT (the active form of CPT-11)-induced inhibitory effect on DNA topoisomerase I was examined. The topoisomerase I-inhibitory effect of 7-ethyl-1-hydroxy-CPT was enhanced 10-fold in the presence of Nedaplatin (NSC 375101D, NDP) at microgram/milliliter concentrations<sup>[1]</sup>. Nedaplatin (NSC 375101D, NDP) was developed as a second generation platinum complex. Because it has greater antitumour activity and lower nephrotoxicity than cisplatin (CDDP). At the high-dose of Nedaplatin (NSC 375101D, NDP) in FN therapy, a reduction of tumour size and long-term tumour-free survival were frequently observed. The survival effect of the combinations of Nedaplatin (NSC 375101D, NDP) with 5-FU was superior to those of the combination of CDDP with 5-FU. In conclusion, the sequence-dependent antitumour efficacy and toxicity of the combination of NDP or CDDP with 5-FU was demonstrated in this study, and FN therapy appeared to be the most efficient regimen as a clinical therapy<sup>[2]</sup>.



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