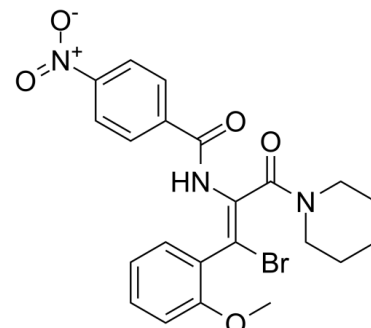


## Data Sheet

<b>Product Name:</b>	AT-130
<b>Cat. No.:</b>	CS-0017999
<b>CAS No.:</b>	211364-06-6
<b>Molecular Formula:</b>	C <sub>22</sub> H <sub>22</sub> BrN <sub>3</sub> O <sub>5</sub>
<b>Molecular Weight:</b>	488.33
<b>Target:</b>	DNA/RNA Synthesis; HBV
<b>Pathway:</b>	Anti-infection; Cell Cycle/DNA Damage
<b>Solubility:</b>	DMSO : 25 mg/mL (51.19 mM; Need ultrasonic)



### BIOLOGICAL ACTIVITY:

AT-130, a phenylpropenamide derivative, is a potent **hepatitis B virus (HBV)** replication non-nucleoside inhibitor. AT-130 inhibits the viral **DNA synthesis** with an **EC<sub>50</sub>** of 0.13 μM. AT-130 inhibits both wt and mutant HBVs. AT-130 has anti-HBV activity in hepatoma cells<sup>[1][2][3]</sup>. **In Vitro:** AT-130 inhibits Wt HBV (IC<sub>50</sub>=2.4 μM), rtL180M HBV (IC<sub>50</sub>=9.8 μM), rtM204I HBV (IC<sub>50</sub>=35.6 μM)<sup>[1]</sup>.

AT-130 (0.1, 1, 5, 10, 100 μM; for 7 days) causes dose-dependent inhibition of wt HBV replication in HepG2 cells transduced with HBV baculovirus. AT-130 at a concentration of 2.5 μM, reduces encapsidated HBV DNA by 50% (IC<sub>50</sub>) and at 18.5 μM by 90% (IC<sub>90</sub>)<sup>[1]</sup>.

AT-130 has no toxic to either HepG2 or Huh-7 cells at concentrations of up to 250 μM<sup>[1]</sup>.

AT-130 (0.005, 0.05, 0.5, 5, 50 μM) does not inhibit HBV DNA synthesis by blocking the HBV endogenous DNA polymerase reaction directly in Huh 7 or HepG2 cells. AT-130 inhibits HBV DNA replication in hepatoma cells but has no effect on viral DNA polymerase activity or core protein translation<sup>[3]</sup>.

AT-130 (2.5, 18.5 μM) has no effect on total HBV RNA production but does reduce encapsidated RNA. AT-130 does not affect core protein or nucleocapsid production and the activity of the protein expression vector<sup>[3]</sup>.

### References:

[1]. William E Delaney 4th, et al. Phenylpropenamide derivatives AT-61 and AT-130 inhibit replication of wild-type and lamivudine-resistant strains of hepatitis B virus in vitro. *Antimicrob Agents Chemother.* 2002 Sep;46(9):3057-60.

[2]. R B Perni , et al. Phenylpropenamide derivatives as inhibitors of hepatitis B virus replication. *Bioorg Med Chem Lett.* 2000 Dec 4;10(23):2687-90.

[3]. J J Feld, et al. The phenylpropenamide derivative AT-130 blocks HBV replication at the level of viral RNA packaging. *Antiviral Res.* 2007 Nov;76(2):168-77.

### CAIndexNames:

Benzamide, N-[(1E)-2-bromo-2-(2-methoxyphenyl)-1-(1-piperidinylcarbonyl)ethenyl]-4-nitro-

### SMILES:

O=C(N/C(C(N1CCCCC1)=O)=C(Br)C2=CC=CC=C2OC)C3=CC=C([N+][O-])C=C3

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

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