
Product Data Sheet

Product Name: Vibunazole (BAY-N-7133)

Cat. No.: GC32321

Chemical Properties

Cas. No. 80456-55-9

SMILES OC(C(C)(C)C)(COC1=CC=C(Cl)C=C1)CN2N=CN=C2Formula C₁₅H₂₀ClN₃O₂ M.Wt 309.79

Solubility Soluble in DMSO Storage Store at -20°C

General tips For obtaining a higher solubility , please warm the tube at 37 °C and shake it in the ultrasonic bath for a while. Stock solution can be stored below -20°C for several months.

Shipping Condition Evaluation sample solution : ship with blue ice All other available size: ship with RT , or blue ice upon request.

Structure **Protocol****Animal experiment:**

Under pentobarbital anesthesia 26 to 30 g female Swiss-Webster albino mice are injected intratracheally with either 0.05 mL sterile 0.9 % NaCl solution or 150 arthroconidia of *Coccidioides imrnitis* (strain Silveira) suspended in 0.05 mL of 0.9 % NaCl solution. Treatment of cohorts of ten infected and ten uninfected mice is begun 72 h after inoculation. Daily i.v. injections (tail vein) of 0.1 mL of 5 % glucose solution delivering either 0, 2.5, 5 or 10 mg/kg/dose of Vibunazole are given for 30 days[1].

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

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References:

[1]. Hoeprich PD, et al. Activity of BAY n 7133 and BAY 1 9139 in vitro and in experimental murine coccidioidomycosis. Eur J Clin Microbiol. 1985 Aug;4(4):400-3.

Background

Vibunazole is a new antifungal azole.

Vibunazole is an antifungal azole. Low concentrations of all three drugs inhibit *Coccidioides immitis*, strain Silveira, in vitro with a descending order of activity ketoconazole>Vibunazole>BAY 1 9139[1].

The untreated, infected mice lost weight initially and progressively, whereas treated mice gain weight after an initial loss with Vibunazole (all doses), BAY 1 9139 and ketoconazole at 2.5 mg/kg/day. With both Vibunazole and BAY 1 9139, the 5 and 10 mg/kg doses yield serum concentrations exceeding the MICs for the *Coccidioides immitis* test strain (0.8 and 1.5 µg/mL respectively) for periods in excess of 30 min after injection[1].

[1]. Hoeprich PD, et al. Activity of BAY n 7133 and BAY 1 9139 in vitro and in experimental murine coccidioidomycosis. Eur J Clin Microbiol. 1985 Aug;4(4):400-3.

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