
Product Data Sheet

Product Name: Doxapram

Cat. No.: GC16537

Chemical Properties

Cas. No. 309-29-5

Chemical Name 1-ethyl-4-(2-morpholin-4-ylethyl)-3,3-diphenylpyrrolidin-2-one

SMILES CCN1CC(C(C1=O))(C2=CC=CC=C2)C3=CC=CC=C3)CCN4CCOCC4Formula $C_{24}H_{30}N_2O_2$ M.Wt 378.51

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 **Background**

Doxapram is a strong, dose-dependent respiratory stimulant in mammals. The oral LD50 in rats is 211mg/kg [1].

After Doxapram treatment, it has been noted a pressor response action through stimulation of the central nervous system (CNS). In addition, Doxapram has shown not only the central sensitization but also the peripheral sensitization to hypercarbia in both animal and human studies. Furthermore, Doxapram has been reported to stimulate the carotid body in a dose-dependent fashion and have an additive but not synergistic effect on the carotid body response to hypercapnia. Apart from these, Doxapram has also shown the potent, direct and inhibitory effects on cloned TASK-1 and TASK-3 channels with the EC50 values of 410nM, 37nM and 9nM for TASK-1, TASK-3 and TASK-1/TASK-3, respectively. Doxapram has been revealed to augment neuromuscular transmission in a dose-related manner above a threshold concentration of 50µM [1].

References:

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

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[1] Yost CS. A new look at the respiratory stimulant doxapram. CNS Drug Rev. 2006 Fall-Winter;12(3-4):236-49.

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