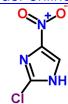


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IDNUMBER ST029401
CAS 57531-37-0
MDL NUMBER MFCD03419295
Molecular formula C3H2CIN3O2

Molecular weight 147.52 Purity 97%

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IUPAC2-chloro-4-nitroimidazoleSmilesc1(c[nH]c(n1)CI)[N+]([O-])=O

THERAPEUTIC CATEGORNY amoebic

ACCEPTORS 2
DONORS 1
ROTATION BONDS 0
N+O 5
Chiral Centers 0
LogP 0.49
LogS -2.48
LIPINSKI 4

Synonyms TIMTEC-BB SBB000100;1H-IMIDAZOLE, 2-CHLORO-4-NITRO-;2-CHLORO



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Xi Risk Statements 36/37/38 Safety Statements 26-36 HazardClass IRRITANT

2-Chloro-4-nitroimidazole

References

Watras, J., Widel M., Suwinski J., Salwinska E. 2-Chloro-4-nitroimidazole radiosensitizers of hypoxic tumor cells in vivo. Journal Neoplasma. 1987;34(3):253-9.

Abstract

The transplantable rhabdomyosarcoma in WAG/Rij rats was used to test the in vivo effectiveness of 1-methyl-2-chloro-4-nitroimidazole (P13) and its analog 1-(2-hydroxy-3-methoxy-propyl)-2-chloro-4-nitroimidazole (P40) as tumor-cell radiosensitizers after their i.p. administration at low doses. The results indicate that both compounds administered repeatedly at nontoxic concentrations (70-150 mg/kg body wt.) in combination with moderate fractional doses of irradiation (3.7 Gy) enhance the radiation effect on tumors. The local control of tumors on the 210th day in all experimental groups has been higher than in the control ones. In the case of P40 administered in the maximal dose, the increment in local control is statistically significant (p less than 0.05). The regrowth delay has also been longer after treatment with radiosensitizers. In the course of treatment we did not observe any symptoms of neurotoxicity of the compounds. The mean body weight of rats during the administration of compounds remained on the level of control rats whose tumors were irradiated only.