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# **ProductInformation**

# N-(2-Chloroethyl)-N-ethyl-2-bromobenzylamine hydrochloride

Product number **C 8417**Store at Room Temperature

Synonyms: N-Ethyl-N-(2-chloroethyl)-2-bromobenzylamine hydrochloride, DSP-4

#### **Product description**

CAS RN: 40616-75-9

Molecular formula: C<sub>11</sub>H<sub>15</sub>BrCIN· HCI

Molecular weight: 313.06

N-(2-Chloroethyl)-N-ethyl-2-bromobenzylamine hydrochloride (DSP-4) is an adrenergic neurotoxin that induces acute and relatively selective degeneration of both central and peripheral noradrenergic nerve terminals. DSP-4 was found to intensify and modify the epileptic activity in the iron-induced chronic epilepsy in rat model. When the effect of DSP-4 on vigilance state was studied it provided evidence for differential involvement of the noradrenergic locus coeruleus system in sleep mechanisms.

DPS-4 was used in the study of the involvement of amygdala noradrenergic and serotonergic systems in memory storage processing, and the results suggested that the memory modulatory effect of peripheral E partially involved the amygdala noradrenergic system. <sup>4</sup> It was found that dopamine metabolism in the frontal cortex, as measured *ex vivo*, was increased in animals treated with low (10 mg/kg) but not with a high doses (50 mg/kg) of DPS-4. <sup>5</sup> In addition noradrenergic lesion with DSP-4 decreases dopamine release in the striatum and enhances catalepsy in experimental models of Parkinson's disease. <sup>6</sup>

#### Precautions and Disclaimer.

This product is for R&D use only, not for drug, household, or other uses. Please consult the Material Safety Data Sheet for information regarding hazards and safe handling practices.

### **Preparation instructions**

The product is soluble in ethanol and in water at 50 mg/ml yielding a clear colorless to faint yellow solution. Also soluble in 0.1 N NaOH at 7 mg/ml. Solutions should be prepared and used freshly.

## Storage and Stability

Store desiccated at room temperature. Under these conditions the product is stable for 3 years.

#### References

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- Sharma, V. and Singh, R., Electroencephalographic study of the effect of neurotoxin DSP-4 in iron model of chronic focal epilepsy., Indian J. Exp. Biol., 37, 468-75 (1999).
- 3. Gonzalez, M.M., et al., Noradrenaline neurotoxin DSP-4 effects on sleep and brain temperature in the rat., Neurosci. Lett., **248**, 93-6 (1998).
- 4. Liang, K.C., Pretraining infusion of DSP-4 into the amygdala impaired retention in the inhibitory avoidance task: involvement of norepinephrine but not serotonin in memory facilitation., Chin. J. Physiol., **41**, 223-33 (1998).
- Haidkind, R., et al., Denervation of the locus coeruleus projections by treatment with the selective neurotoxin DSP-4 [N (2-chloroethyl)-Nethyl-2-bromobenzylamine] reduces dopamine release potential in the nucleus accumbens shell in conscious rats., Neurosci. Lett., 332, 79-82 (2002).

6.	Srinivasan, J. and Schmidt, W.J., Functional recovery of locus coeruleus noradrenergic neurons after DSP-4 lesion: effects on dopamine levels and neuroleptic induced-parkinsonian symptoms in rats., J. Neural Transm,, <b>111</b> , 13-26 (2004).	NDH,PHC 01/05-