



## Product Information

### Xylazine

Product Number **X 1126**

Storage Temperature -0 °C

#### Product Description

Molecular Formula: C<sub>12</sub>H<sub>16</sub>N<sub>2</sub>S

Molecular Weight: 220.3

CAS Number: 7361-61-7

Melting Point: 136-139 °C; 140-142 °C<sup>1</sup>

Xylazine is an  $\alpha_2$ -adrenergic receptor agonist, sedative, and muscle relaxant. A ketamine hydrochloride and xylazine hydrochloride solution (Product No. K 113) has been used to produce rapid and reversible anesthesia in experimental animals.<sup>2</sup> Xylazine elicited a diuretic response in ketamine-anesthetized animals, probably through activation of complex peripheral and CNS  $\alpha_2$ -adrenergic receptor systems.<sup>3</sup> Xylazine did not discriminate among the four known  $\alpha_2$  B-adrenergic receptor subtypes (A<B<C and D) in membranes of particular tissues.<sup>4</sup>

#### Precautions and Disclaimer

For Laboratory Use Only. Not for drug, household or other uses.

#### Preparation Instructions

Xylazine is soluble in methanol (50 mg/ml), yielding a clear, colorless solution. It is also soluble in dilute acid (HCl) and in chloroform. Xylazine is practically insoluble in water and in alkali solutions.

#### References

1. The Merck Index, 13th ed., Entry# 10135.
2. Soltesz, I., and Deschenes, M., Low- And High-Frequency Membrane Potential Oscillations During Theta Activity In CA1 And CA3 Pyramidal Neurons Of the Rat Hippocampus Under Ketamine-Xylazine Anesthesia. *J. Neurophysiol.* **70(1)**, 97-116 (1993).
3. Cabral, A. D., et al., Central Alpha2-receptor Mechanisms Contribute To Enhanced Renal Responses During Ketamine-Xylazine Anesthesia. *Am. J. Physiol.*, **275(6 pt 2)**, R1867-R1874 (1998).
4. Schwartz, D. D., and Clark, T. P., Affinity Of Detomidine, Medetomidine And Xylazine For Alpha-2 Adrenergic Receptor Subtypes. *J. Vet. Pharmacol. Ther.*, **21(2)**, 107-111 (1998).
5. Ware, T. D., and Paul, D., Cross-tolerance Between Analgesia Produced By Xylazine And Selective Opioid Receptor Subtype Treatments. *Eur. J. Pharmacol.*, **389(2-3)**, 181-185 (2000).

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