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Product Information

# Terazosin hydrochloride

≥98% (TLC), powder

#### T4680

Store at room temperature.

## **Product Description**

Molecular Formula: C<sub>19</sub>H<sub>25</sub>N<sub>5</sub>O<sub>4</sub>.HCl

Molecular Weight: 423.9 CAS Number: 63590-64-7 Melting Point: 278-279 °C

**Synonyms**: 1-(4-amino-6,7-dimethoxy-2-quinazolinyl)-4-[(tetrahydro-2-furanyl) carbonyl]

piperazine hydrochloride;

2-[(4-tetrahydro-2-furoyl)-1-piperazinyl]-4-amino-

6,7-dimethoxyquinazoline hydrochloride

Terazosin is an a1-receptor blocker that is structurally very similar to prazosin, differing in that terazosin contains a tetrahydrofuran unit at the amide linkage whereas prazosin contains a furan unit. The duration of action of terazosin is extended relative to that of prazosin. A review of the pharmacodynamic and pharmacokinetic properties of terazosin has been published.<sup>2</sup> Terazosin (2 μM) has been shown to abolish the norepinephrine response that leads to enhanced c-myc-encoded mRNA levels in cultured cardiac myocytes.<sup>3</sup> Terazosin has been used to probe apoptosis (15 µM) and the rate of DNA synthesis (1-100 µM) in cultured human prostate cancer cells.4 It has also been utilized to modulate the effects of brain epinephrine in the regulation of motor activity and movement in mice.<sup>5</sup> Assays for the detection of terazosin in plasma by HPLC and by HPLC/ESI-MS have been reported.<sup>6-8</sup>

#### Precautions and Disclaimer

For laboratory use only. Not for drug, household or other uses.

#### **Preparation Instructions**

This product is soluble in methanol (20 mg/mL), with heat as needed, yielding a clear, colorless solution.

#### References

- 1. Martindale The Extra Pharmacopoeia, 31st ed., Reynolds, J. E. F., ed., Royal Pharmaceutical Society (London, UK: 1996), p. 952.
- Wilde, M. I., et al., Terazosin. A review of its pharmacodynamic and pharmacokinetic properties, and therapeutic potential in benign prostatic hyperplasia. Drugs Aging, 3(3), 258-277 (1993).
- 3. Starksen, N. F., et al., Cardiac myocyte hypertrophy is associated with c-myc protooncogene expression. Proc. Natl. Acad. Sci. USA, 83(21), 8348-8350 (1986).
- Kyprianou, N., and Benning, C. M., Suppression of human prostate cancer cell growth by α1-adrenoceptor antagonists doxazosin and terazosin via induction of apoptosis. Cancer Res., 60(16), 4550-4555 (2000).
- 5. Stone, E. A., et al., Role of epinephrine stimulation of CNS a1-adrenoceptors in motor activity in mice. Synapse, 49(1), 67-76 (2003).
- 6. Cheah, P. Y., et al., Improved high-performance liquid chromatographic analysis of terazosin in human plasma. J. Chromatogr. B Biomed. Sci. Appl., 745(2), 439-443 (2000).
- 7. Sekhar, E. C., et al., Determination of terazosin in human plasma, using high-performance liquid chromatography with fluorescence detection. J. Chromatogr. B Biomed. Sci. Appl.,710(1-2), 137-142 (1998).
- 8. Zavitsanos, A. P., and Alebic-Kolbah, T., Enantioselective determination of terazosin in human plasma by normal phase highperformance liquid chromatography-electrospray mass spectrometry. J. Chromatogr. A., 794(1-2), 45-56 (1998).



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