

CAPSULES

preliminary notes and applications from Bioanalytical Systems, Inc.

Clonidine Formulations

Purpose

Determination of clonidine in drug formulations.

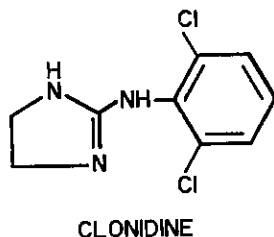


Figure 1. Structure of clonidine.

Clonidine (2-(2,6-dichloroanilino)-2-imidazoline, F1) is an antihypertensive agent used in the treatment of high blood pressure. The drug works by stimulating α -adrenergic receptors in the central nervous system. Side effects include analgesia, hypothermia, sedation, and a withdrawal syndrome which includes hypertension, tachycardia and anxiety. Effective therapeutic concentrations are low (0.2- 2.2 ng/mL

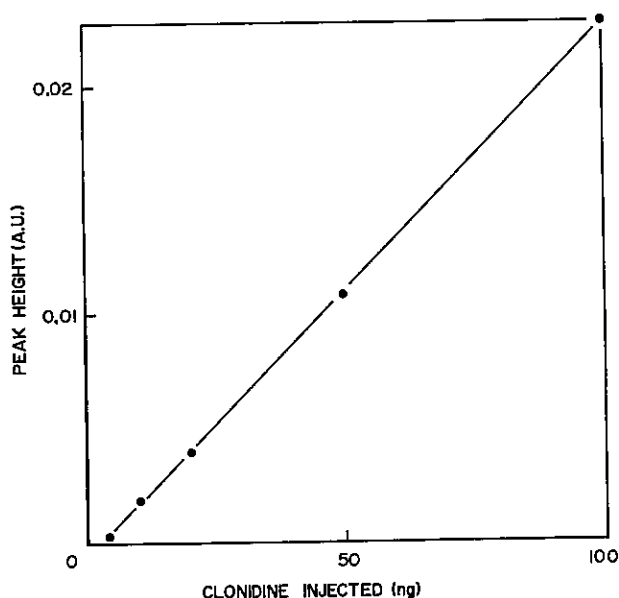


Figure 2. Calibration curve for clonidine standards. Each point represents the mean of 2 determinations.



Figure 3. Sample chromatogram of 2 ng clonidine.

plasma). At the time of this writing we were not aware of any published LC procedures for clonidine in body fluids.

Existing Methods

GC, GC-MS, GC-ECD [1], radioimmunoassay and radiolabelling have been used for therapeutic monitoring or pharmacokinetic studies. The USP method for tablets requires UV spectrophotometric measurement, which is non-specific [2].

Conditions

System: BAS 400 Liquid Chromatograph
 Detector: BAS UV-108 variable wavelength (220 nm)
 Column: BAS 3 μ m Phase II Octyl reverse-phase (100 x 3.2 mm) (PN MF-6214)
 Mobile Phase: 82.7% (v:v) 0.02 M KH_2PO_4 , pH 6.0; 17.5% acetonitrile. Flow rate was 0.9 mL/min.
 Detection Limit: 200 pg injected standard (S/N = 3)
 Linear Range: 0.5-100 ng injected standards

Sample Preparation

Appropriate amounts of clonidine hydrochloride (corrected for the amount of free base) were dissolved in mobile phase and injected in 20 μ L aliquots.

Notes

The determination of clonidine also can be performed on the BAS 200 Problem Solver.

References

1. Nazarali, A.J., *J. Chromatogr.* 380 (1986): 393-400.
2. Walters, S.M. and D.B. Stonys, *J. Chromatogr. Sci.* 21 (1983): 43-45.

