
Underneath The Stars Book

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Determination Of Phenylpropanolamine In Pharmaceutical

A convenient HPLC method to determine phenylpropanolamine (PPA) in addition to phenylephrine (PE) and chlorpheniramine (CPA) in commercially available over-the-counter (OTC) preparations has been developed. Sample solutions were prepared by dilution with water or methanol followed by filtration and direct injection into the HPLC system.

HPLC determination of phenylpropanolamine in ...

Phenylpropanolamine (PPA) hydrochloride used as nasal decongestant agent was determined by second derivative spectrophotometry after derivatization with 2-hydroxynaphthaldehyde. The absorbance was measured between minimum-maximum of 386 nm-392 nm. The linear calibration range was obtained within 0.5~2.0 µg/mL.

Determination of Phenylpropanolamine in Pharmaceutical ...

Determination of phenylpropanolamine hydrochloride and acetaminophen in pharmaceutical preparations by Raman spectroscopy. Thomas H. King. The Department of Chemistry, Florida State University, Tallahassee, FL 32306

Determination of phenylpropanolamine hydrochloride and ...

The amount of PPA in the pharmaceutical preparations was evaluated from an external calibration curve prepared from standards of pure PPA. 3. Results and discussion. Phenylpropanolamine (PPA) absorbs only weakly in the UV region at 210 and 256 nm with a molar absorptivity of 3778 and 156 l mol⁻¹ cm⁻¹. In order to increase the spectrophotometric sensitivity for the determination of PPA,

a derivatization process with DAB was carried out.

HPLC determination of phenylpropanolamine in ...

HPLC determination of phenylpropanolamine in pharmaceutical preparations using 4-dimethylaminobenzaldehyde as a derivatizing reagent

HPLC determination of phenylpropanolamine in ...

A convenient HPLC method to determine phenylpropanolamine (PPA) in addition to phenylephrine (PE) and chlorpheniramine (CPA) in commercially available over-the-counter (OTC) preparations has been developed. Sample solutions were prepared by dilution with water or methanol followed by filtration and direct injection into the HPLC system.

HPLC determination of phenylpropanolamine in ...

A rapid, accurate and precise method for the determination of phenylpropanolamine hydrochloride in a cough-mixture is described. The method is based on the determination by normal-phase HPLC of benzaldehyde formed after periodate oxidation. The method is applicable to other phenalkanolamine drugs that yield benzald

Determination of phenylpropanolamine hydrochloride as ...

HPLC determination of phenylpropanolamine in pharmaceutical preparations using 4-dimethylaminobenzaldehyde as a derivatizing reagent Rind, Rind; Khuhawar, Khuhawar; Rajper, Rajper Fluorimetric method of analysis for D -norpseudoephedrine hydrochloride, glycine and L -glutamic acid by reversed-phase high performance liquid chromatography

HPLC determination of phenylpropanolamine in ...

A selective and simple spectrophotometric method has been developed for the determination of phenylpropanolamine HCl (PPA) in its dosage forms. The method was based on the formation of a colored N-vinyl chlorobenzoquinone derivative of PPA through its reaction with 2,3,5,6-tetrachloro-1,4-benzoquinone in presence of acetaldehyde.

A New Spectrophotometric Method for Determination of ...

determination of chlorpheniramine maleate,3-11 phenylpropanolamine hydrochloride12-19 and combination of chlorpheniramine maleate & phenylpropanolamine hydrochloride20-22 have been mentioned. These methods are time consuming; therefore an alternative method of multi wavelength spectroscopy by UV spectrophotometry is rendered.

SPECTROPHOTOMETRIC DETERMINATION OF CHLORPHENIRAMINE ...

High-performance liquid chromatography (HPLC) has been used for the determination of phenylpropanolamine alone and in pharmaceutical formulations (6-11), but little information has appeared on the HPLC analysis of phenylpropanolamine in biological fluids (12, 13).

Determination of Phenylpropanolamine in Serum and Urine by ...

A colorimetric method for the determination of phenylpropanolamine hydrochloride, based on the reaction with ninhydrin, was applied to various forms of pharmaceutical products. The method is applicable to pharmaceutical products that do not contain other primary or secondary amines. The determination is dependent upon the combined hydroxyl and primary amino moieties. It is postulated that ...

Selective determination of phenylpropanolamine ...

Analytical procedure has been developed for the gas chromatographic determination of phenylpropanolamine (PPA) using trifluoroacetylacetone (FAA) as derivatizing reagent. Elution is carried out from the column HP-5 (30 m × 0.32 mm i.d.) with film thickness 0.25 µm at initial column temperature 70 °C for 5 min, followed by heating rate 10 °C/min up to 120 °C.

Capillary gas chromatographic determination of ...

(1995). Simultaneous Determination of Buzepide, Phenylpropanolamine, and Clocinizine in Pharmaceutical Preparations by Ion-Pair Reversed-Phase HPLC. Journal of Liquid Chromatography: Vol. 18, No. 2, pp. 227-234.

Simultaneous Determination of Buzepide ...

Journal of Pharmaceutical and Biomedical Analysis 41 (2006) 998-1001 Short communication
Capillary gas chromatographic determination of phenylpropanolamine

Capillary gas chromatographic determination of ...

Phenylpropanolamine hydrochloride (PP), 2-amino-1-phenyl-1-ol (14838-15-4) is a largely indirect acting sym-pathomimetic agent and less active as a central nervous stimulant [15]. Several methods have been reported for the quantitative determination of phenylpropanolamine hydrochloride in pure and in dosage forms, including

Spectrophotometric microdetermination of nefopam ...

Objective: A particular, easy and precise reverse phase liquid chromatographic technique was developed for the synchronous determination of Bromohexine and...

Analytical Method Development and Validation for ...

The linear calibration curve was obtained with 30-150 microg/ml PPA with detection limit of 6.0 microg/ml. The method was used for the determination of PPA from Sinutab and Tavegyl-D tablets. The relative standard deviation (R.S.D.) for the analysis of pharmaceutical preparation was obtained within 0.4-0.9%.