



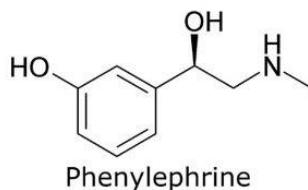
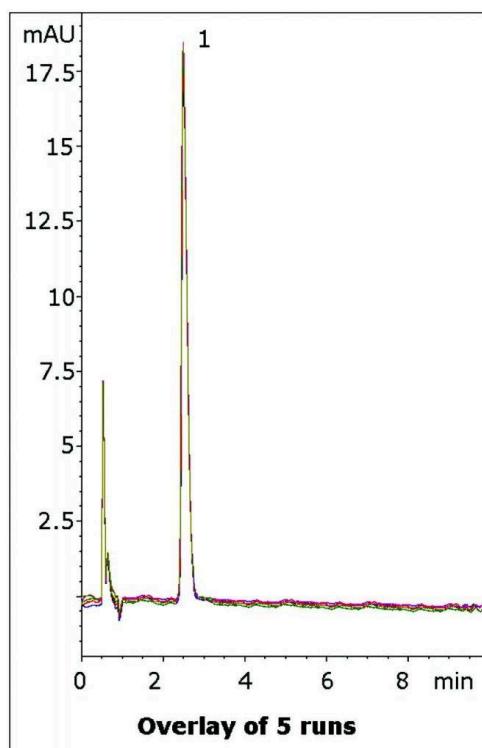
Phenylephrine HCL Tablet Analyzed with HPLC - AppNote

Analysis of Phenylephrine, a Highly Polar Compound

Phenylephrine was first studied under various Reversed Phase conditions using a C18 Stationary Phase, but even using a 95% Aqueous Mobile Phase a Retention factor of no more than 0.6 could be obtained. With such low retention, it may be difficult to Separate Phenylephrine from excipients or other APIs using this type of Column.

This Method produces a Retention factor of 3.7 with an excellent Peak Shape and could be suitable for routine assay of this API when in a Tablet Formulation.

Five Chromatograms are shown in the Figure to illustrate the repeatability of the data.



Peak:
Phenylephrine

Method Conditions

Column: Cogent Diamond Hydride 2.0TM, 2.2 μ m, 120 \AA

Catalog No.: [70200-05P-2](#)

Dimensions: 2.1 x 50mm

Mobile Phase: 5:95 A: DI Water with 0.1% Trifluoroacetic Acid (*TFA*) v/v, / B: Acetonitrile with 0.1% Trifluoroacetic Acid (*TFA*) v/v

Injection vol.: 0.2 μ L

Flow rate: 0.4mL / minute

Detection: UV @ 225nm

Sample Preparation: 10mg strength Phenylephrine HCL Tablet was ground and added to a 25mL volumetric flask. A portion of the Mobile Phase was added and the flask was sonicated 10 minutes. Then it was diluted to mark and mixed. A portion was filtered through a 0.45 μ m Nylon Syringe Filter (MICROSOLV Tech Corp.).

t₀: 0.5 minutes

Note: Phenylephrine is a common decongestant used in many over-the-counter cold medications. Combination with other drugs is also prevalent in many formulations, so there is a need for a method for phenylephrine from other APIs.



Attachment No 272 Phenylephrine HCL Tablet Analyzed with HPLC pdf 0.4 Mb [Download File](#)

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