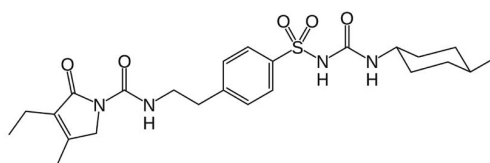
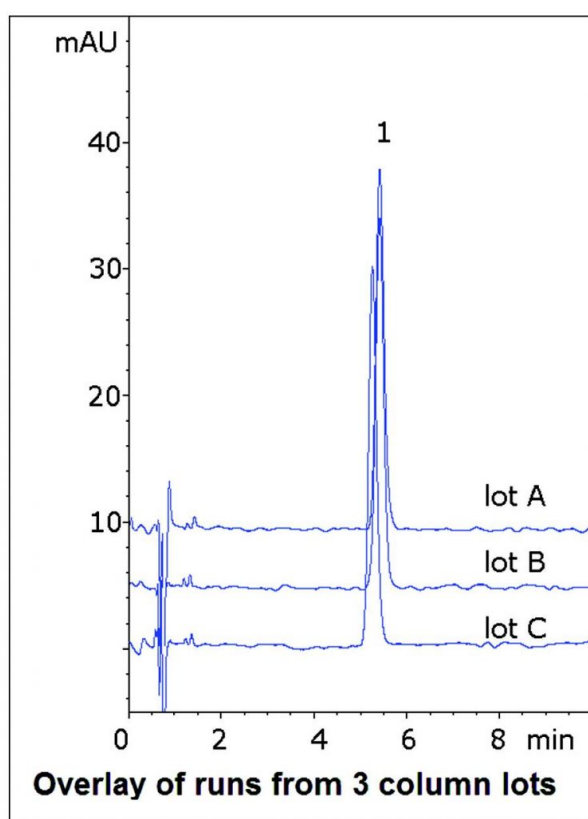


Glimepiride Tablet Analyzed with HPLC - AppNote

Isocratic Assay Method for a Diabetes Drug

In this study, an Isocratic, HPLC Method for Glimepiride Analysis in a tablet formulation is presented. The USP Assay Method for this drug calls for a Phosphate Buffer and is therefore not LCMS compatible. This Method uses Formic Acid instead and produces a Symmetrical, well-Retained Peak. Therefore the analysis could be applied to various LCMS studies as well if desired, such as with clinical laboratories dealing with biological extracts.

Data from three different Column lots is shown in the figure in order to demonstrate the Method Reproducibility and Robustness.



Glimepiride

Peak:

Method Conditions

Column: Cogent Bidentate C18™, 4µm, 100Å

Catalog No.: 40018-75P

Dimensions: 4.6 x 75mm

Mobile Phase: 50:50 DI Water / Acetonitrile with 0.1% Formic Acid (v/v)

Injection vol.: 5µL

Flow rate: 1.0mL / minute

Detection: UV @ 230nm

Sample Preparation: 1mg strength Glimepiride Tablet was ground and transferred to a 4mL volumetric flask. A portion of 50:50 Acetonitrile / DI Water was added and the flask was sonicated 10 minutes. It was then diluted to mark and mixed. A portion was filtered with a 0.45µm Nylon Syringe Filter (MicroSolv Tech Corp.) and diluted 1:10.

t₀: 0.9 minutes

Note: *Glimepiride is a sulfonylurea drug indicated to treat type 2 Diabetes Mellitus. It works by increasing insulin production in the pancreas. It is sold under trade names such as Amaryl®.*



Attachment

No 253 Glimepiride Tablet Analyzed by HPLC pdf 0.4 Mb [Download File](#)

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