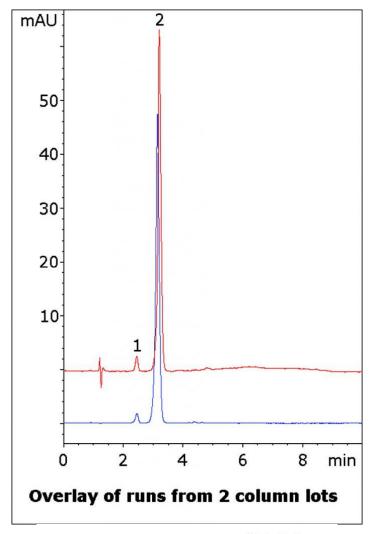


# Famotidine Tablet Analyzed with HPLC - AppNote

## **API Separation from Matrix Component**

This Method for Analysis of Famotidine Tablets is easy to perform and produces a Symmetrical Peak Shape for the API. This compound has numerous amines which can be problematic in terms of Peak Shape with conventional Columns. Separation from a component from the tablet extract matrix is obtained as well, illustrating specificity of the Method.

Reproducibility is shown by the overlay of runs from two different Column lots.



$$\begin{array}{c|c} & & & & & & & & \\ H_2N & & & & & & \\ NH_2 & S & & & & & \\ NH_2 & S & & & & \\ N & S & O & & \\ \end{array}$$

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#### Peaks:

1. Matrix Component

2. Famotidine

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### **Method Conditions**

**Column**: Cogent Diamond Hydride<sup>™</sup>, 4μm, 100Å

**Catalog No.**: 70000-7.5P **Dimensions**: 4.6 x 75mm

**Mobile Phase:** 

A: DI Water with 0.1% Trifluoroacetic Acid (*TFA*) v/v B: Acetonitrile with 0.1% Trifluoroacetic Acid (*TFA*) v/v

#### **Gradient**:

Time (minutes)	%B
0	95
2	95
6	50
7	95

Post Time: 3 minutes
Injection vol.: 1µL

Flow rate: 1.0mL / minute Detection: UV @ 265nm

**Sample Preparation**: 10mg strength Famotidine tablet was ground and added to a 25mL volumetric flask. A portion of 50:50 Solvent A / Solvent B diluent was added and the flask was sonicated 10 minutes. It was then diluted to mark and filtered with a 0.45µm Nylon Syringe Filter (MicroSolv Tech Corp.).

**to**: 0.9 minutes



#### Attachment

No 221 Famotidine Tablet Analyzed with HPLC pdf 0.4 Mb Download File

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