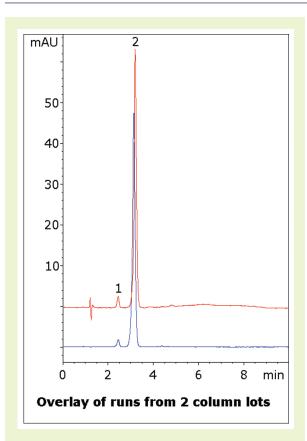


## **Famotidine Tablet**

## Separation from matrix component



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

**Note:** Famotidine is an acid reducer that is used to treat ulcers, gastroesophageal reflux disease, heartburn, and other related conditions. It is sold under trade names such as Pepcid® and Calmicid®.

## **Method Conditions**

Column: Cogent Diamond Hydride™, 4µm, 100Å

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

Solvents: A: DI H<sub>2</sub>O / 0.1% trifluoroacetic acid (v/v)

B: Acetonitrile / 0.1% trifluoroacetic acid (v/v)

 Gradient:
 time (min.)
 %B

 0
 95

 2
 95

 6
 50

 7
 95

Post Time: 3 min
Injection vol.: 1µL
Flow rate: 1.0 mL/min

Detection: UV 265 nm

Sample: 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 min. It was then diluted to mark and filtered with a  $0.45\mu m$  nylon syringe filter (MicroSolv Tech Corp.).

**Peaks:** 1. Matrix component 2. Famotidine

to: 0.9 min

## **Discussion**

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 Cogent Diamond Hydride column lots.