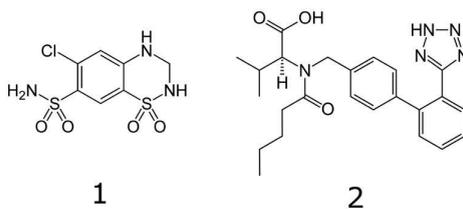
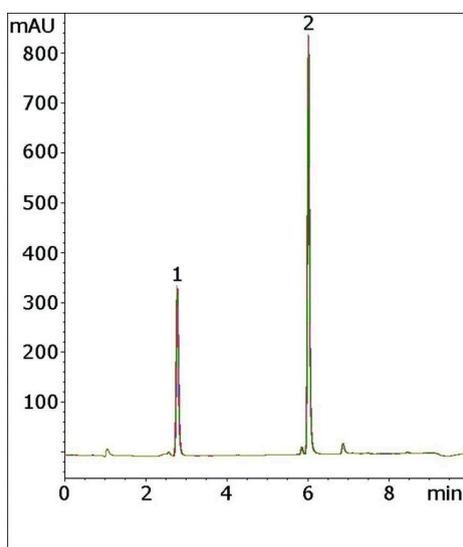


Valsartan and Hydrochlorothiazide Analyzed with HPLC USP Comparison - AppNote

Improved Gradient Method with Faster Equilibration

The USP Assay Method for Valsartan in combination with Hydrochlorothiazide features a 27 minute gradient with a 13 minute re-equilibration for a total run time of 40 minutes. In this method, the run time was a quarter of the USP method, and the Column equilibrates much faster when gradients are used. This demonstrates a substantial time and solvent savings for the analytical laboratory.

Five Chromatograms are overlaid below which shows the Robustness and Precision of this Method.



Peaks: 1. Hydrochlorothiazide (HCT), 2. Valsartan

Method Conditions

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

Catalog No.: 40018-75P

Dimensions: 4.6 x 75 mm

Mobile Phase:

- A: DI Water / 0.1% Trifluoroacetic Acid (TFA)
- B: Acetonitrile / 0.1% Trifluoroacetic Acid (TFA)

Gradient:

Time (minutes)	%B
0	10
8	90
9	10

Post Time: 1 minute**Injection vol.:** 10 µL**Flow rate:** 1.0 ml / minute**Detection:** UV @ 265 nm**Sample Preparation:**

- **Stock Solution:** A Diovan® HCT brand tablet containing 160 mg Valsartan and 25 mg Hydrochlorothiazide was ground and added to a 50 mL volumetric flask. The flask was diluted to mark with 50:50 Solvent A / Solvent B mixture and sonicated. A portion was then filtered with a 0.45 µm Nylon Syringe Filter AQ™ Brand (MICROSOLV Tech Corp.).
- **Working Solution:** 100 µL of the stock solution was diluted with 900 µL of a 50:50 Solvent A / Solvent B mixture

t₀: 1 minute

Comparison of This Method and the USP Method		
	Bidentate C18	Ordinary C18
Total Run Time	10 Minutes	40 Minutes
Column Volume Equilibration	1 Column	10 Columns
Solvent Usage per Run	10ml	40ml

**Attachment No 150 Valsartan and Hydrochlorothiazide Analyzed with HPLC pdf 0.5 Mb**[Download File](#)

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MicroSolv Technology Corporation

9158 Industrial Blvd. NE, Leland, NC 28451

Tel: (732) 380-8900

Fax: (910) 769-9435

Email: customers@mtc-usa.comWebsite: www.mtc-usa.com