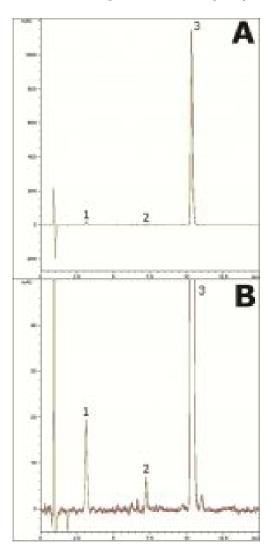


Hormone Replacement Capsule Analysis by HPLC - AppNote

Separation of Estriol, Estradiol, & Progesterone

This gradient method features a separation of the three components of a hormone replacement formulation. *Figure A* shows a five run overlay of the formulation extract injections. *Figure B* shows a zoomed-in view so that the Estriol and Estradiol Peaks, which are present in much lower concentration than Progesterone, can be seen clearly. *Figure B* also shows separation of an impurity from the Progesterone Peak.



Peak:



2. Estradiol

3. Progesterone

Method Conditions

Column: Cogent UDC Cholesterol[™], 4μm, 100Å

Catalog No.: 69069-7.5P **Dimensions:** 4.6 x 75mm

Mobile Phase:

A: DI Water / 0.1% Formic Acid (v/v)
B: Acetonitrile / 0.1% Formic Acid (v/v)

Gradient:

Time (minutes)	%B
0	33
2	33
11	65
12	33

Post Time: 3 minutes

Flow rate: 1.0 mL / minute Detection: UV @ 210nm

Injection vol.: 1µL

Sample Preparation: The contents of a capsule containing 0.124 mg Estradiol, 1.001 mg Estriol, and 50 mg Progesterone were added to a 25 mL volumetric flask. The flask was diluted to mark with solvent B and sonicated 10 minutes. Then A portion was filtered with a $0.45\mu m$ Nylon Syringe Filter (MicroSolv Tech. Corp.). Peak identities were confirmed by individual standards of 0.1 mg / mL in a Solvent B diluent.



Attachment

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