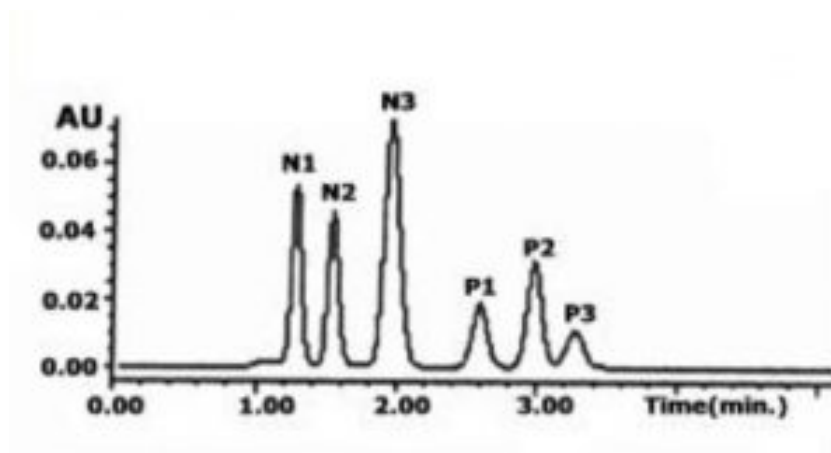


Nucleoside Analogs Analyzed with HPLC – AppNote

Polar and Non-Polar Anticancer Prodrugs Analyzed in the Same Run

In this Method, we display Retention of both polar and nonpolar compounds in a simple Isocratic run. Adjustment of the aqueous to organic ratio can be used to make either the polar or more hydrophobic compounds elute first. In the example presented, the more hydrophobic compounds are retained longer but retention and separation of the polar constituents is still achieved. If greater retention of the polar compounds is desired then a higher percentage of Acetonitrile is required.



Method Conditions

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

Catalog No.: 69069-7.5P

Dimensions: 4.6 x 75mm

Mobile Phase: 60% Acetonitrile / 40% DI Water / 0.5% Formic Acid

Injection vol.: 5µL

Flow rate: 1 mL / minute

Detection: UV @ 270nm

Sample Preparation: Proprietary Drugs. N1-N3 Nucleoside Analogues. P1-P3 have greater hydrophobicity with polar functional groups. 1µg / mL of each dissolved in the Mobile Phase

Notes: Nucleoside analogue anticancer “pro” drugs are difficult to retain on conventional HPLC Columns because of their highly polar natures. As such, Ion Pair Reagents are used in order to achieve sufficient Retention. These additives interfere with MS detection and are not preferred. With this Method, you can achieve good Retention paired with a MS-Compatible Mobile Phase.

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Attachment

No 05 Nucleoside Analogs Analyzed with HPLC pdf 0.1 Mb [Download File](#)

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