

## SINCE 1992

## Nucleobases analyzed by HPLC – AppNote

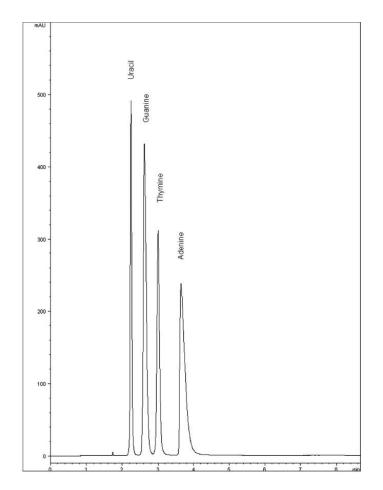
## Nucleotides Uracil, Guanine, Thymine, Adenine, Excellent Peak Shape and Resolution

This Method is easy to prepare, use and reproduce. Separation is accomplished under 100% Aqueous Conditions yet there is an alternate Selectivity. These bases may be difficult to retain on Columns with ordinary Silica that contain significant amounts of Silanols.



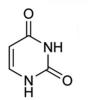
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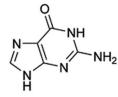
HPLC Columns and Chromatography Accessories





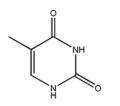
HPLC Columns and Chromatography Accessories



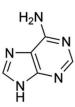


Uracil

Guanine



Thymine



Adenine



**Peaks:** 

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- 1. Uracil (U)
- 2. Guanine (G)
- 3. Thymine (T)
- 4. Adenine (A)

## **Method Conditions**



HPLC Columns and Chromatography Accessories



Column: Cogent Diamond Hydride<sup>™</sup>, 4µm, 100Å Catalog No.: 70000-75P Dimensions: 4.6 x 75mm Mobile Phase: DI Water / 0.1% Acetic Acid Temperature: 25°C Injection vol.: 2.5µL Flow rate: 1mL / minute Detection: UV @ 254nm

**Notes:** Nucleobases (or Nucleotide Bases) are the parts of DNA and RNA that may be involved in pairing. The main Bases are Cytosine, Guanine, Adenine (DNA and RNA), Thymine (DNA) and Uracil (RNA). They are usually simply called "Bases" in Genetics.



Attachment

Nucleobases Analyzed by HPLC pdf Download File