Size-exclusion chromatography is a powerful technique for the size-dependent separation of biomolecules. While standard, non-denaturing mobile phases are often desirable, denaturing mobile phases containing guanidine hydrochloride (or SDS), may be used to insure un-aggregated, monomers for proper size determination. As shown below, denaturation shifts molecules toward larger apparent size and smaller retention volume.

**Comparison of Denaturing and Nondenaturing Mobile Phases in SEC: ZORBAX GF-250, +/- GuCl**

**Application**
Biochemical
Robert Ricker

**Highlights**
- Agilent ZORBAX GF-250 columns separate biomolecules in a size-dependent manner within a molecular-weight range of 4,000-400,000 daltons. Note, some molecules (depending on their characteristics) are retained by non-SEC mechanisms and are shifted from linearity.

- ZORBAX GF-250 columns are manufactured using extremely hard particles, allowing high flow rates and fast separations.

- When larger molecules are used in combination with denaturing mobile phases, ZORBAX GF-450 may be used to obtain more-linear separation in the upper molecular-weight range.

**Conditions:**
LC: Hewlett Packard 1050
Column: ZORBAX GF-250 (9.4 x 250), Agilent Part No. 884973-901
UV: 254 nm
Flow: 2.0 mL / min.; ambient
Inj. Vol.: 20 µL (1 µg / µL)

*Mean 2 expts.
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