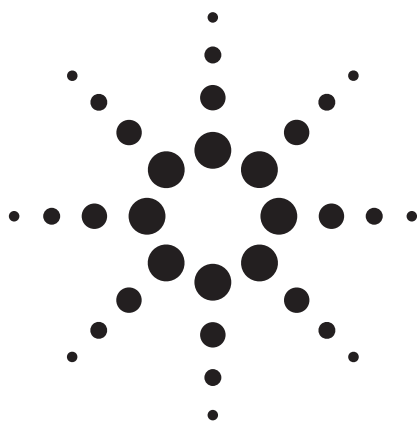




Agilent ZORBAX SB-C8 Rapid Resolution Threaded Column

Data Sheet

General Description

ZORBAX SB-C8 Rapid Resolution threaded columns are especially designed for higher pressure operation (up to 600 bar) and are packed with a high-performance microparticulate C8 packing for high-speed reversed-phase HPLC. The StableBond packing is made by chemically bonding a sterically protected C8 stationary phase to a specially prepared, high-purity ZORBAX porous silica microsphere. The special ZORBAX silica support is designed to reduce or eliminate strong adsorption of basic compounds. The densely covered, sterically protected, diisopropyl n-octylsilane stationary phase is chemically stable and gives longer column life. As a result, ZORBAX SB-C8 is a stable reversed-phase packing that can be used for basic, neutral, or acidic samples. It is particularly well suited for use with aggressive mobile phases (for example, pH < 2, high-ionic-strength, ion-pair additives, etc.) since the steric protection of the bonded phase resists degradation caused by such mobile phases. This material has also been shown to be stable when operated at temperatures up to 80 °C. These characteristics are particularly important for use in methods that need long-term stability and reproducibility. ZORBAX SB-C8 is especially suited in applications that use high-sensitivity detectors that require low backgrounds (for example, mass spectrometers).

The uniform, spherical ZORBAX SB-C8 particles have a controlled pore size of 80 Å and a surface area of 180 m²/g. Columns are loaded to a uniform bed density using a proprietary, high-pressure, slurry-loading technique to give optimum column efficiency.

Column Characteristics

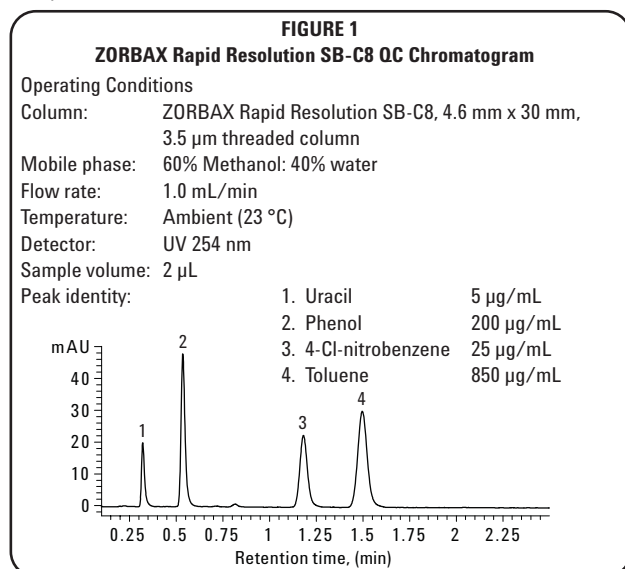
A typical quality control chromatogram for a 3.5-μm ZORBAX Rapid Resolution SB-C8 4.6 mm id × 30 mm threaded column is shown in Figure 1. The actual QC test and performance of your column is described in the Column Performance Report enclosed with your column.

Safety Considerations

- All points of connection in liquid chromatographic systems are potential sources of leaks. Users of liquid chromatographic equipment should be aware of the toxicity or flammability of their mobile phases.
- These Rapid-Resolution threaded columns are mechanically stable and have been tested to very high pressures to ensure safe lab operation on a variety of LC instruments. The 2.1- and 3.0-mm id columns will support 20,000 psi (1,300 bar) operation and 4.6-mm id columns will support 16,000 psi (1,000 bar) operation. Opening columns may compromise these pressure limits. Chromatographic performance has not been tested above 600 bar.
- Because of its small particle size, dry ZORBAX packings are respirable. Columns should only be opened in a well-ventilated area.

Operational Guidelines

- The direction of flow is marked on the column.
- While generally not harmful to the column, reversing flow should be avoided except to attempt removal of inlet plugging (see "Column Care" section).
- These columns are packed and assembled for high-pressure (up to 600 bar) use. Disassembling the column will degrade column performance.
- ZORBAX SB-C8 is compatible with water and all common organic solvents.



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- Avoid use of columns below pH 1.8 or above pH 8.0.
- Maximum operating pressure is 600 bar (9,000 psi).
- Maximum operating temperature is 80 °C.

NOTE: StableBond columns are designed for high stability at low pH (for example, pH < 5). However, all silica-based packings have some solubility in pH > 6 aqueous mobile phases. Therefore, when using silica-based columns under conditions of pH > 6, maximum column lifetime is obtained by operation at low temperatures (< 40 °C) using low buffer concentrations in the range of 0.01 to 0.02 M. Column stability at pH > 6 is also enhanced by avoiding phosphate and carbonate buffers (reference: H. A. Claessens, M. A. van Straten, and J. J. Kirkland, *J. Chromatogr. [A]*, 728 [1996] 259).

Mobile Phase Selection

The bonded stationary phase is nonpolar in nature and is best used with mobile phases such as methanol/water or acetonitrile/water mixtures. Increasing the amount of organic component usually reduces the retention time of the sample. Due to the relatively high viscosity of recommended mobile phases, increased efficiency can be achieved with the use of column temperatures in the range of 40 to 65 °C. Gradient elution techniques for this packing often use 5% methanol or acetonitrile in water as the initial solvent and 100% methanol or acetonitrile as the final solvent. Additional information on solvent selection may be found in Chapters Six and Seven, *Introduction to Modern Liquid Chromatography*, Second Edition, L. R. Snyder and J. J. Kirkland, (John Wiley & Sons, 1979), and Chapters Six, Seven, and Eight, *Practical HPLC Method Development*, Second Edition, L. R. Snyder, J. J. Kirkland, and J. L. Glajch, (John Wiley & Sons, 1997).

Applications

ZORBAX SB-C8 can be used with basic, neutral, or acidic analytes. For many basic compounds, it will normally not be necessary to use basic modifiers, such as triethylamine, to achieve efficient, symmetrical peaks. However, very basic compounds may require the addition of basic modifiers such as 10 to 20 mM dimethyl-octylamine or 20 to 30 mM triethylamine. Such samples are often best chromatographed with mobile phases of pH = 3. One highly recommended mobile phase for very basic compounds is 0.1% trifluoroacetic acid adjusted to pH = 3 with triethylamine and an appropriate concentration of methanol or acetonitrile.

ZORBAX SB-C8 can be used at 80 °C at low pH and is therefore a good choice for higher temperature separations. High temperature can reduce mobile phase viscosity, lower operating pressure, and change band spacing (selectivity).

Column Care

Samples that contain particulate matter may plug the column inlet frit and should be filtered before injection into the column. If solvent flow appears to be restricted

(unusually high column back-pressure), check first to see that solvent flow is unobstructed up to the column inlet. If the column has the restriction, there may be particulate matter on the inlet frit. An attempt should be made to remove any inlet debris by back-flushing 25 to 30 mL of mobile phase through the column. If this fails to return the column to near its original back-pressure, the column should be replaced.

To remove strongly retained materials from the column, flush the column with stronger (less polar) solvents. Solvents such as methanol, acetonitrile, or a 95/5 mixture of dichloromethane and methanol should remove most highly retained compounds. In extreme cases, dimethyl sulfoxide or dimethylformamide at low flow rates may also be used for this purpose. When switching between solvents with vastly different polarities, it may be necessary to first purge the column with a mutually miscible solvent such as isopropanol.

Since columns have 3/8-inch end nuts, a short 3/8-inch wrench should be used to attach the columns to the instrument to avoid any additional tightening of the end fittings. Overtightening the end fittings will cause damage and require column replacement.

Storage Recommendations

Long-term storage of silica-based, bonded-phase columns should be in a pure organic solvent, preferably an aprotic liquid such as 100% acetonitrile. If the column was previously used with a buffered mobile phase, the buffer should first be removed by purging the column with 20 to 30 column volumes of a 50/50 mixture of methanol or acetonitrile and water, followed by 20 to 30 column volumes of the pure solvent. Before storing the column, the end-fittings should be tightly capped with end plugs to prevent the packing from drying out. Columns may be safely stored for short periods in most mobile phases. However, to protect equipment, it is desirable to remove salts from the instrument and column by purging the column with the same mobile phase without the buffer (for example, using 60/40 ACN/H₂O to remove a 60/40 ACN/0.02 M phosphate buffered mobile phase). Re-equilibration is rapid with the original mobile phase when using this approach, and any danger of corrosion from the salts is eliminated.

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