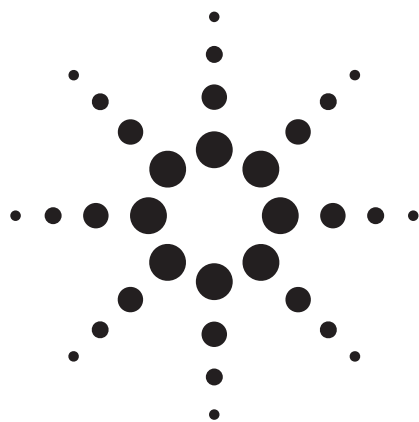




Agilent ZORBAX Eclipse Plus C18 Rapid Resolution Threaded Column

Data Sheet

General Description

ZORBAX Eclipse Plus C18 Rapid-Resolution threaded columns are specially designed for higher pressure operation (up to 600 bar) and are packed with a high-performance microparticulate C18 packing for high-speed reverse-phase HPLC. Eclipse Plus C18 columns are designed for superior peak shape with basic compounds and deliver high efficiency and excellent peak shape with all sample types. Eclipse Plus C18 packing is made by first chemically bonding a dense monolayer of dimethyl-n-octadecylsilane stationary phase to a specially prepared, improved ultra-high purity ($\geq 99.995\%$ SiO₂), ZORBAX Rx-SIL porous silica support. This special ZORBAX silica support (Type B) is designed to reduce or eliminate strong adsorption of basic and highly polar compounds. The bonded-phase packing is then doubly end-capped using proprietary reagents and procedures to obtain maximum deactivation of the silica surface. Eclipse Plus C18 columns can be used for acidic and neutral samples, but are especially suited for separations of basic compounds that produce poor peak shapes on other columns. These columns can be used for a wide range of applications and over a pH range of 2 to 9, accommodating most popular mobile phases.

The uniform, spherical Eclipse Plus C18 particles are based on an improved ZORBAX Rx-SIL support that has a nominal surface area of 160 m²/g and a controlled pore size of 95 Å. Columns are loaded to a stable, uniform bed density using a proprietary high-pressure slurry-loading technique to give maximum column efficiency.

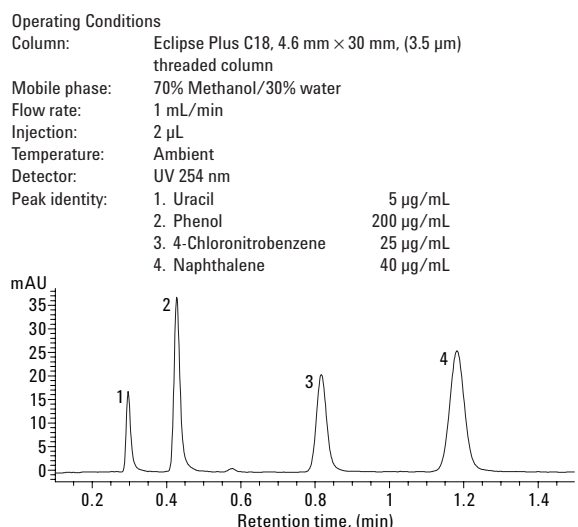
Column Characteristics

A typical quality control (QC) chromatogram for a ZORBAX Rapid Resolution Eclipse Plus C18 4.6 × 30 mm, 3.5-μm 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.

FIGURE 1
ZORBAX Rapid Resolution Eclipse Plus C18 QC Chromatogram



- These rapid resolution assembled columns are mechanically stable and have been tested to very high pressures to assure 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 the 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 it is not harmful to the column, reverse flow should be avoided except to attempt removal of inlet blockage (see "Column Care").
- These columns are packed and assembled for high pressure (up to 600 bar) use. Disassembling the column will degrade column performance.



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- Eclipse Plus C18 is compatible with water and all common organic solvents.
- Avoid use of this column below pH 2 or above pH 9
- Maximum operating pressure is 600 bar (9,000 psi).
- Maximum operating temperature is 60 °C.

NOTE: Eclipse Plus columns are designed for high stability over a wide pH range. 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 10 to 20 mM. 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).

- Columns should not be maintained at neutral or elevated pH or elevated temperature when not in use.

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 60 °C; however, best column lifetime is achieved with operation at ≤ 40 °C. Gradient elution techniques for this packing often use 5% methanol or acetonitrile 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. L. Glajch, and J. J. Kirkland, (John Wiley & Sons, 1997).

Applications

Eclipse Plus C18 can be used with basic, neutral, or acidic compounds. Ionizable compounds (basic, acidic) generally are best separated at pH ~ 3 with this column. However, Eclipse Plus C18 is especially suited for separating basic compounds when intermediate pH (4 to 8) must be used to maintain compound stability or to obtain desired band spacing (selectivity). For optimum results and long-term reproducibility, the use of 10- to 50-mM buffers is always recommended when separating ionizable compounds.

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. Over-tightening 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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