Ascentis Express HILIC Care & Use Sheet

Ascentis Express Description
Ascentis Express HILIC is a high-speed, high-performance liquid chromatography column based on a new Fused-Core® particle design. The Fused-Core particle provides a thin porous shell of high-purity silica surrounding a solid silica core. This particle design exhibits very high column efficiency due to the shallow diffusion paths in the 0.5 µm thick porous shell and the small overall particle size of 2.7 µm. The bare silica stationary phase of Ascentis Express HILIC provides a column that can be used for traditional normal phase separations (not discussed in this document) or for aqueous normal phase chromatography with the typical reversed-phase mobile phases for hydrophilic interactive liquid chromatography (HILIC) of basic, acidic, or neutral compounds.

Column Characteristics
The Fused-Core particle has a surface area of ~ 150 m²/g and an average pore size of 90Å. The Fused-Core particles are 30% to 50% heavier than commercially available totally porous particles due to the density of the solid cores. Therefore, the effective surface area per column is similar to columns packed with totally porous particles having surface areas in the 225-300 m²/g range.

Operation Guidelines
- The direction of flow is marked on the column label.
- Reversed flow may be used to attempt removal of inlet pluggage or contamination.
- A new column is shipped in 100% acetonitrile.
- Water and all common organic solvents are compatible with Ascentis Express HILIC columns.
- Ascentis Express HILIC columns are best used at temperatures below 60 °C for maximum column life.
- Mobile phase pH for Ascentis Express HILIC columns is best maintained in the range of pH = 2 to 8 for maximum column stability.
- Ascentis Express HILIC columns are stable to operating pressures up to 600 bar (9000 psi).

Column Care
To maximize column life, ensure that samples and mobile phases are particle-free. The use of guard columns or an in-line filter with 0.5 µm porosity between the sample injector and the column is highly recommended. The 2 µm porosity frits on Ascentis Express HILIC columns are less subject to pluggage than are the 0.5 µm frits typically used with other small-particle columns. Should the operating pressure of the column suddenly increase beyond normal levels, reversing the flow direction of the column may be attempted to remove debris on the inlet frit.

Column Storage
Long-term storage of silica-based columns is best in 100% acetonitrile. Columns may be safely stored for short periods (up to 3 or 4 days) in most common mobile phases. However, when using buffers, it is best to remove the salts to protect both the column and the HPLC equipment by first flushing mobile phases. However, when using buffers, it is best to remove the salts to protect both the column and the HPLC equipment by first flushing mobile phases.

Applications
HILIC is a useful and complimentary method to reversed-phase chromatography (RPC) and is especially attractive in situations where compound retention is poor in RPC and very high levels of water are required in the mobile phase for adequate retention. Retention in HILIC appears to be a combination of hydrophilic interaction, ion-exchange and some reversed-phase retention. The aqueous layer which forms on the surface of HILIC particles promotes interaction with polar solutes. Retention in HILIC as a function of the mobile phase is just opposite from that in RPC. The strongest mobile phase has a high concentration of water and the weakest has a high concentration of organic solvent. Therefore, for gradient separations, the initial mobile phase has a high concentration of organic solvent and the gradient is formed by increasing the aqueous concentration.

Guidelines for Low-Volume Columns
High performance columns with small internal volumes (shorter lengths, internal diameters < 3 mm) are being increasingly used for Fast HPLC. These low-volume columns generate peaks having less volume than those eluting from columns of larger dimensions. The efficiency of separations performed in low-volume columns is highly dependent on the HPLC system having components designed to minimize band spreading. All low-volume columns perform best when used with proper attention to the following factors:
- Detector – Cells should be < 2 µL.
- Detector – Detector response time should be set to ~ 0.1 second and integration set at least 20 points per second.
- Injector – Should be of a low-volume design (e.g., Rheodyne® Model 8125). Auto-samplers may be used for convenience with the expectation of some loss in column efficiency.
- Connecting Tubing – The shortest possible lengths of connecting tubing with narrow internal diameters (at most 0.005-inch, 0.12 mm I.D.) should be used to connect the column to the injector and the detector cell.
- Peak Retention – As retention is increased, the volume of a peak increases, decreasing the effects on band spreading caused by components of the instrument.
- Sample Solvent – For isocratic separations, the sample should be dissolved in the mobile phase or in a solvent that is weaker than the mobile phase. For gradient separations, dissolve sample in the initial mobile phase or in a weaker solvent.
- Injection Volume – For isocratic separations, the volume of sample injected should be 2 µL or less. Sample volumes are less critical for gradient separations, especially if the sample is dissolved in a weak solvent.

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