



SweetSep™ Columns

The New Benchmark for Carbohydrate Analysis

- New columns for reliable HPAEC-PAD and MS
- Superior and fast separations
- Works for all classes of carbohydrates

High-Performance Anion Exchange (HPAE) columns for the separation of mono-, oligo- and polysaccharides using PAD or MS detection.



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Why HPAEC?

High Performance Anion-Exchange Chromatography (HPAEC) is the most powerful analytical technique for carbohydrate analysis due to its ability to separate all classes of alditols (polyols), aminosugars, mono-, oligo- and polysaccharides including glycans, according to structural features such as size, composition, anomericity and linkage isomerism.

Monodisperse Particles for Highest Performance

Antec Scientific developed a novel pellicular anion-exchange stationary phase called SweetSep™ AEX. The phase is based on highly uniform monodisperse 5 µm resin particles of crosslinked poly(divinylbenzene-co-ethylvinylbenzene) copolymer. The particles are coated with latex nano beads, functionalized with a quarternary amine group or a bi-functional group containing a quaternary and tertiary amine.



SEM image of 5 µm SweetSep particles

The resin particles packed in inert PEEK columns result in exceptional column efficiencies with typical reduced plate height close to 2.0 with only moderate column back pressure. SweetSep AEX columns allow for rapid, high-resolution separations of carbohydrates that rival the performance of existing phases based on smaller particle size but operates with significantly lower system back pressures. The size and exchange capacity of the latex nanoparticles is optimized to enable the analysis of a wide variety of carbohydrates samples ranging from monosaccharides present in food, plants and glycoproteins up to oligosaccharides such as FOS (fructo-oligosaccharides) and N-linked glycans.

Available Chemistries

SweetSep columns are currently available in two different chemistries, AEX20 and AEX200. The main difference lies in the functionalization of the latex particle and the capacity. The AEX20 has a bifunctional exchange group consisting of a quarternary and tertiary amine. Meanwhile the AEX200 has a quaternary amine.

The exchange capacity for the analytical column of 4.0 mm ID x 20 cm is ca. 90 µeq for AEX 200 and ca. 160 µeq for the AEX20.

AEX20

Ideal for fast, high-resolution separation of monosaccharides from food samples, monosaccharides from glycoproteins, sialic acids, FDG, Heparin, etc.

AEX200

Universal column for separation of mono- to polysaccharides in food & beverages, plants and glycans.

Column Types and Dimensions

The following column types and dimensions are available:

- Analytical columns
- Pre-columns
- Trap columns

All wetted column parts are made of PEEK independent from the column inner diameter (ID). The 4.0 mm ID columns are made entirely from black PEEK. All 2.1 mm ID columns are PEEK-lined stainless steel columns.



Analytical columns are available as 4.0 and 2.1 mm ID x 20 cm in length.

Pre-columns used as guard column and/or to increase the length of the analytical column are available as 4.0 and 2.1 mm ID x 5 cm columns.

Trap columns are available in the same dimensions as the pre-columns, i.e., 4.0 and 2.1 mm ID x 5 cm.

Features 4.0 mm ID analytical columns

- Workhorse
- Maximum robustness
- Works at higher flow rates

The 4.0 mm ID x 20 cm HPAEC column is a true workhorse, designed for maximum robustness and reliability in demanding carbohydrate applications. Its optimized design allows it to operate efficiently at flow rates of typically 700 $\mu\text{L}/\text{min}$ without compromising performance. Ideal for high-throughput laboratories, this column delivers consistent, reproducible results even under challenging conditions.

Features 2.1 mm ID analytical columns

- Up to 3 times higher sensitivity
- Green chromatography: 4 times lower mobile phase consumption and generation of waste
- No notable loss in separation efficiency using same injection volume as 4.0 mm ID column

The 2.1 mm ID x 20 cm HPAEC column is the ideal choice for high-efficiency separations with enhanced performance. It provides up to 3 times higher sensitivity, ensuring precise and accurate detection of analytes. With typical flow rates of 180 $\mu\text{L}/\text{it}$ it offers 4 times lower mobile phase consumption and waste generation, making it an environmentally friendly solution “Green Chromatography”. Additionally, the column supports larger volume injections (up to 10 μL) with virtually zero loss in separation efficiency.

Instrumental Requirements

1. HPAEC-PAD

SweetSep columns can be used with any High Performance Anion Exchange Chromatography (HPAEC) system such as the IC systems of Thermo Fisher Scientific/Dionex, Metrohm, Shine, etc. Several bioinert HPLC systems are also suitable for use with HPAEC when equipped with a Pulsed Amperometric Detector (PAD) such as the DECADE™ Elite (Antec Scientific). For consistent results, ease of use, and highest reproducibility, the Antec Scientific ALEXYS™ Carbohydrate Analyzer is the best choice.

2. Borate Ion Trap

The use of a Borate Ion Trap (BIT) column installed between the pump and the autosampler is highly recommended. For more information about the BIT column, see page 10.

3. HPAEC-MS

Depending on the volatility of the buffer systems used, for the on-line coupling with MS, the installation of a desalter (ion suppressor) becomes necessary. Basically, any type of (ESI)-MS can be used for detection.

4. HPAEC-PAD/MS

Parallel detection using PAD and MS enables simultaneous quantification and identification of carbohydrates. This is achieved by splitting the flow rate after the SweetSep column into two streams. An in-line desalter (suppressor) is installed before MS introduction to remove Na⁺ ions from the mobile phase.

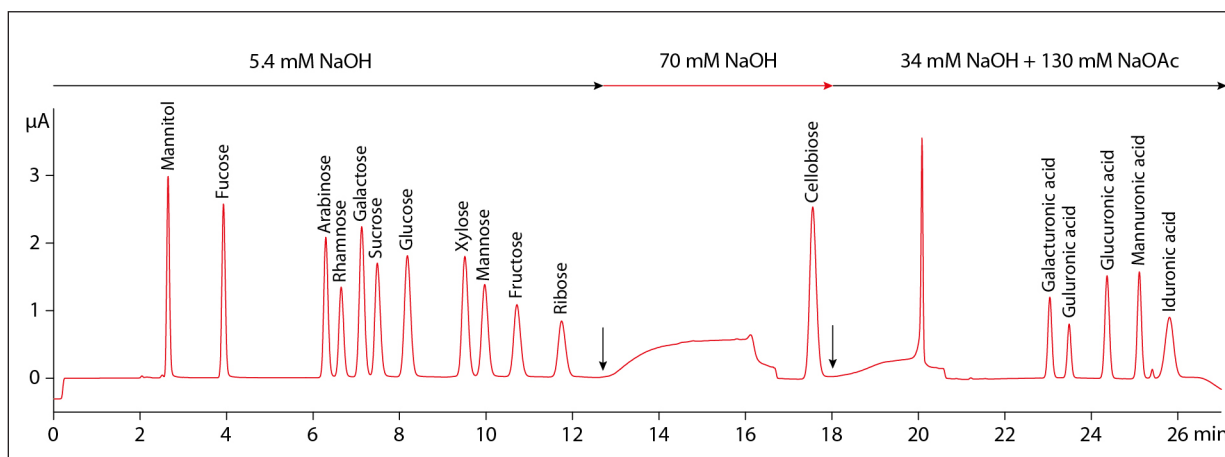
Analytical Columns (4.0 and 2.1 mm ID x 20 cm)

1. SweetSep™ AEX20

Fast, high-resolution separation of mono-saccharides from food samples and biomass, including monosaccharides and sialic acids from glycoproteins, FDG, etc. For specifications, see page 11.

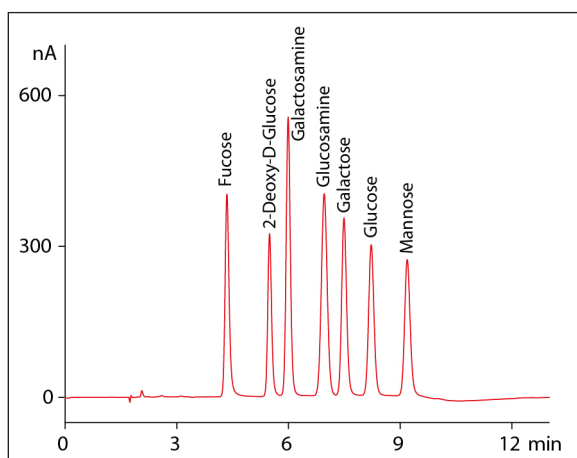


1.1 Monosaccharides and uronic acids from biomass

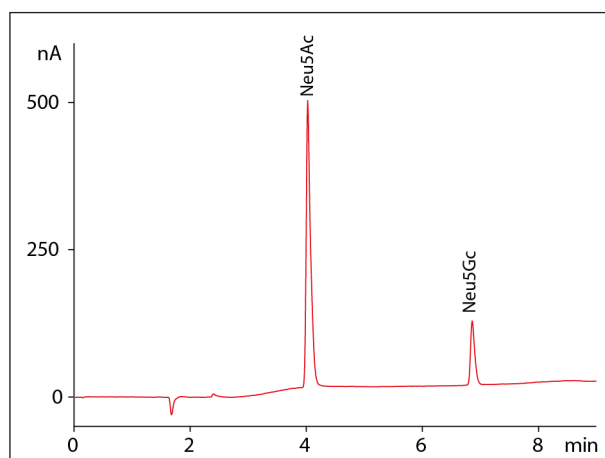


Separation of neutral sugars & uronic acids on a SweetSep™ **AEX20** column, 2.1 mm ID x 20 cm, 10 μ L inj. of a 100 μ M mixtures of neutral sugars & uronic acids std in water (HPAEC-PAD). Arrows indicate the start/end of the baseline disturbance due to step gradient.

1.2 Monosaccharides from glycoproteins including sialic acids



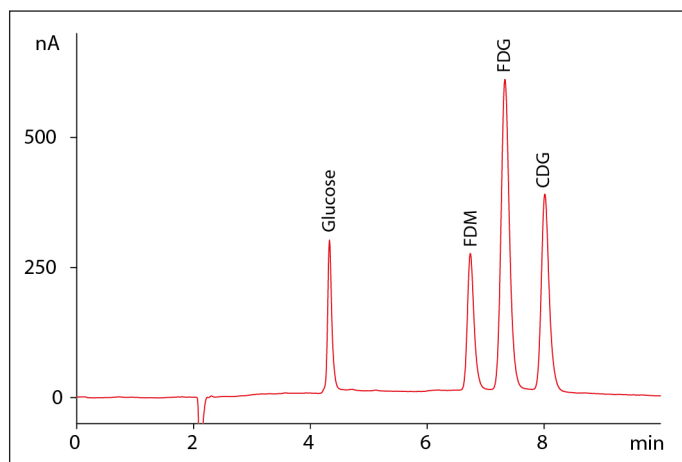
Isocratic separation of monosaccharides on a SweetSep™ **AEX20** column, 4.0 mm ID x 20 cm. 10 μ L inj. of a 10 μ M mixtures of monosaccharides std in water (HPAEC-PAD).



Gradient separation of sialic acids on a SweetSep™ **AEX20**, 4.0 mm ID x 20 cm. 10 μ L inj. of 10 μ M Neu5Ac and 1 μ M of Neu5Gc in water.

Analytical Columns (4.0 and 2.1 mm ID x 20 cm)

1.3 Fluorodeoxyglucose (^{18}F -FDG), optimized EP & USP methods



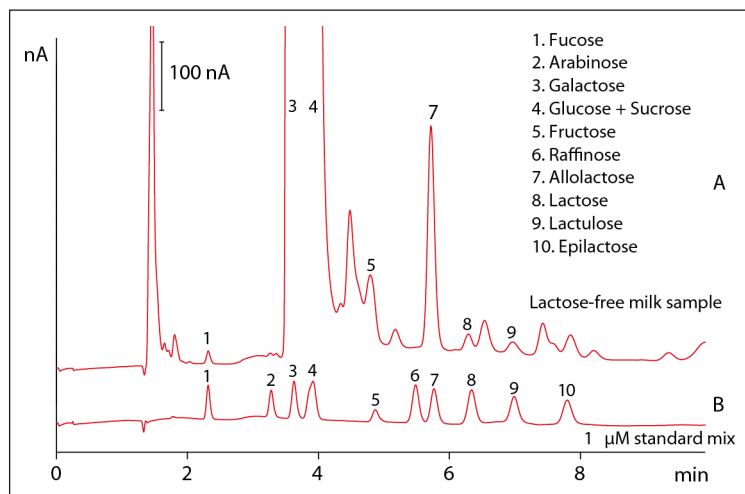
Separation of FDG and its by-products, FDM and CDG according to optimized USP, SweetSep™ **AEX200** column, 4.0 mm ID x 20 cm. 10 μL inj. of a standard mix of 10 $\mu\text{g/mL}$ FDG, FDM, CDG and 1 $\mu\text{g/mL}$ glucose. Resolution: FDM-FDG 2.5 and FDG-CDG 2.7.

2. SweetSep™ AEX200

Universal column for separation of mono- to polysaccharides in F&B, artificial sweeteners, plants, glycans, etc. For specifications, see page 11.



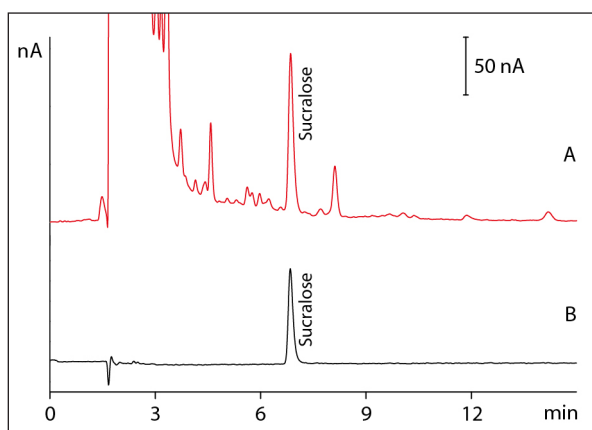
2.1 Mono- and disaccharides in F&B: Lactose Intolerance



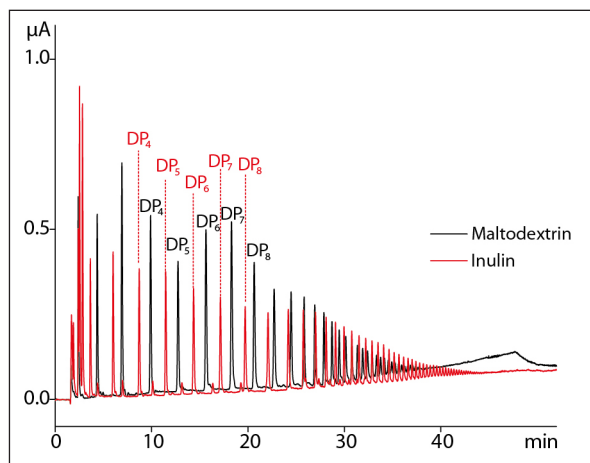
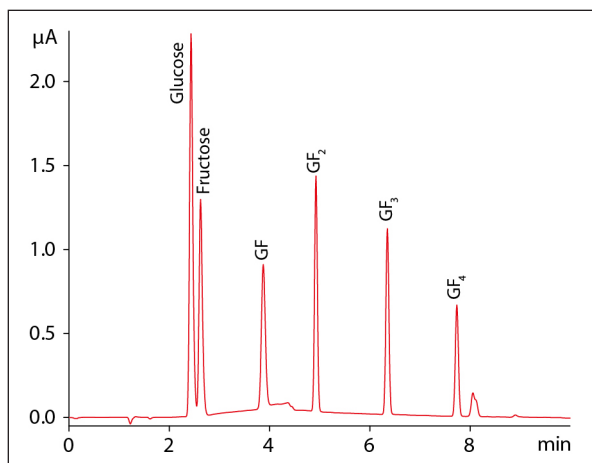
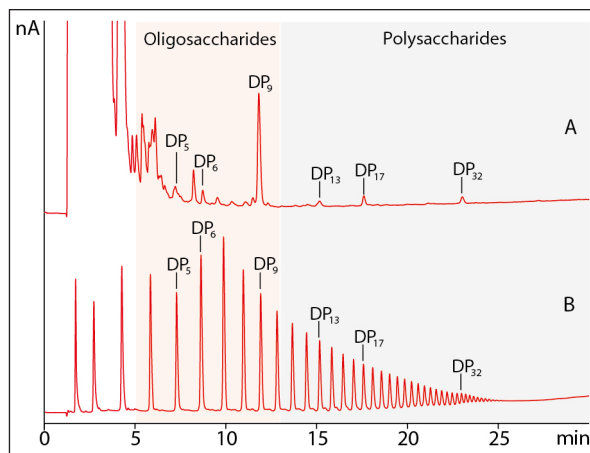
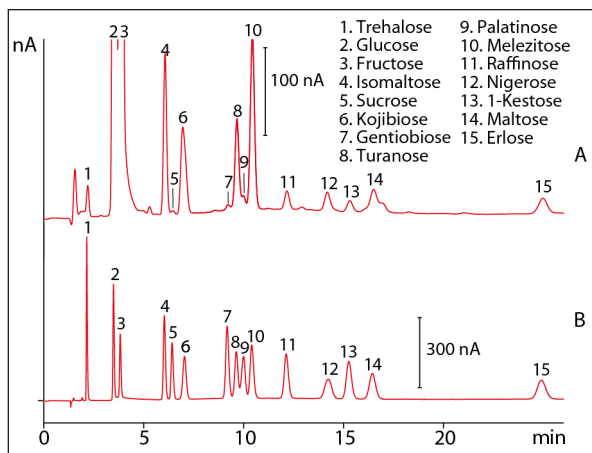
Analysis of milk on a SweetSep™ **AEX200** column, 4.0 mm ID x 20 cm. (A) 10 μL inj. of a 10 g/L lactose-free labelled milk. (B) 10 μL inj. of a 10 μM standard of 10 sugars commonly found in milk, incl. lactose (peak #8) and its isomers, allolactose (#7), lactulose (#9) and epilactose (#10).

Analytical Columns (4.0 and 2.1 mm ID x 20 cm)

2.2. Disaccharide: Sucralose artificial sweetener

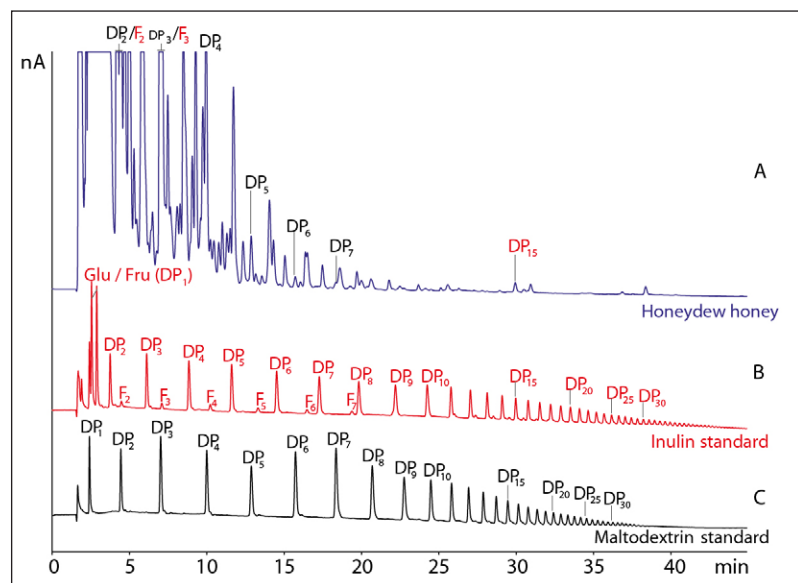


2.3. Mono-, oligo- and polysaccharides in F&B

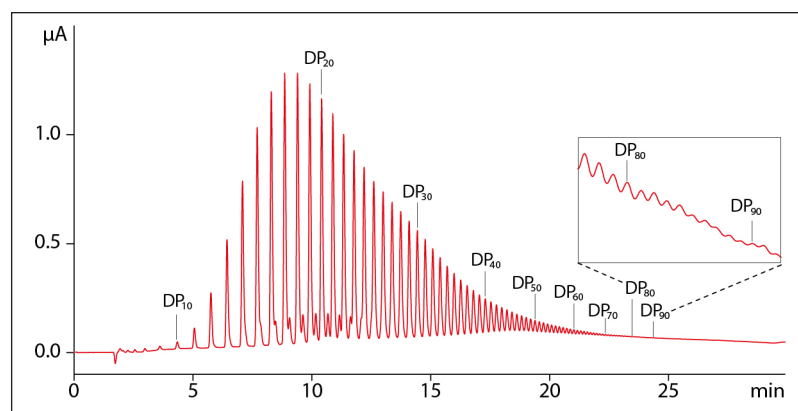


Analytical Columns (4.0 and 2.1 mm ID x 20 cm)

2.4. Oligo- and polysaccharides in F&B

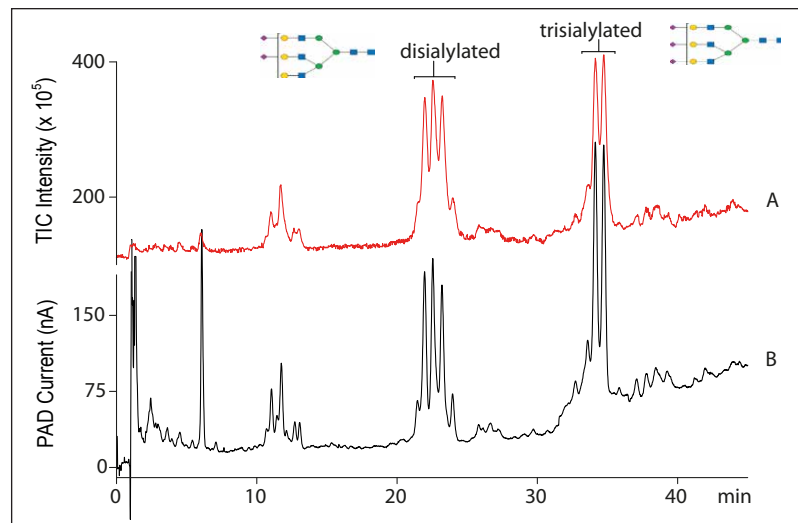


Authentic honeydew honey with natural oligosaccharides, (A) 10 µL injection of honeydew honey,, (B) 200 ppm inulin standard, and (C) 500 ppm maltodextrin standard (black line). Red peaks in (A) correspond to fructo-, black to malto-oligosaccharides. SweetSep **AEX200** column, 4.0 mm ID x 20 cm.



Separation of fructans up to DP 90. SweetSep™ **AEX200** column, 4.0 mm ID x 20 cm + Precolumn 4.0 mm ID x 5 cm

2.5. Glycans from Glycoproteins



Separation of N-glycans standard containing di-, tri-, and tetra sialylated oligosaccharides by HPAEC-PAD/MS on a SweetSep™ **AEX200** column, 4.0 mm ID x 20 cm. A) PAD for quantification, (B) MS for identification with desalter prior to MS

Trap Columns (4.0 and 2.1 mm x 5 cm)*

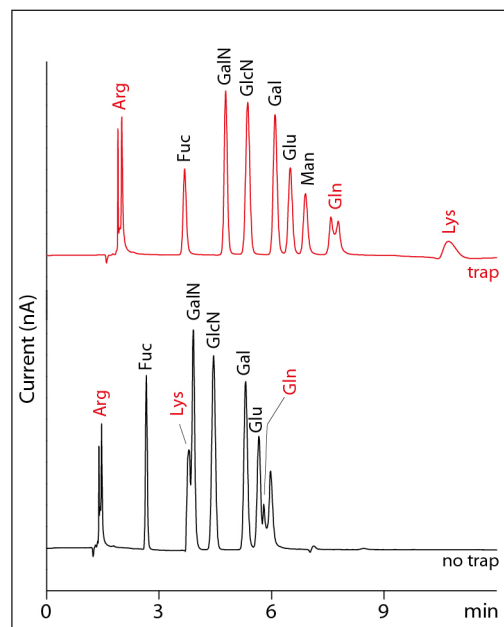
Amino Acid Trap (AAT)

- Efficient trapping of amino acids and small peptides
- Easy installation as precolumn
- Accurate quantification

In compositional analysis of monosaccharides from glycoproteins using HPAEC-PAD, amino acids and small peptides co-elute with the carbohydrates of interest making proper quantification impossible. These amino acids and small peptides are generated during the acid hydrolysis of glycoproteins. Moreover, amino acids can contaminate the Au electrode surface, which might lead to fouling and loss of response even under PAD conditions.

To eliminate the interference of amino acids and to assure accurate quantification of the monosaccharides the use of an Amino Acid Trap (AAT) column is highly recommended.

For specifications/ordering information, see page 11 and 12.



Analysis of monosaccharides with and without trap column. Interfering peaks of Glutamine (Gln) and Lysine (Lys) are efficiently trapped (upper trace) and elute later during the wash step. In both cases the amino acid Arginine (Arg) elutes in the void volume (t_0) of the column.

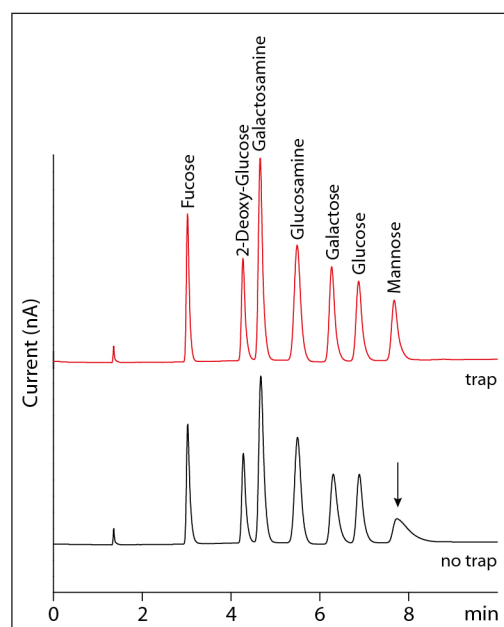
Borate Ion Trap (BIT)

- High borate trapping capacity
- Easy installation between pump and injector
- Reduce peak tailing

In carbohydrate analysis, the peak shape of certain sugars, such as mannose, fructose and sugar alcohols, are deteriorated when traces of borate are in the solvent. These borate contaminants can come from the laboratory deionized (DI) water system.

To eliminate the presence of borate ions and to assure optimal performance Antec Scientific introduced the Borate Ion Trap (BIT) column. The trap column is installed inline between pump and injector of the HPAEC-PAD system.

For specifications/ordering information, see page 11 and 12.



Analysis of carbohydrates with and without Borate Ion Trap (BIT) column. Without trap, the mannose peaks shows significant tailing, see arrow in lower trace.

* The 4 mm ID trap column should be used only with the 4 mm ID analytical column and the 2.1 mm trap with the 2.1 mm ID analytical, respectively.

Pre-Columns (4.0 and 2.1 mm x 5 cm)

- Used as guard column
- Used to extend column efficiency
- Available in all chemistries

The SweetSep™ Pre-columns are high performance columns, individually tested, and available in all chemistries (AEX 20 and 200). They are used as guard column to protect and increase longevity of analytical column and to increase the column length and efficiency by 25%.



The 4 mm ID Pre-column should be used only with the 4 mm ID analytical column and the 2.1 mm Pre-column with the 2.1 mm ID analytical, respectively.

Specifications

Specifications SweetSep™ Anion Exchange Columns

Parameter		AEX200	AEX20
Type		agglomerated pellicular resin	
Particle	Material	ethylvinylbenzene-divinylbenzene copolymer	
	Diameter (µm)	5	
	Functionality	surface sulfonated	
Latex	Material	vinylbenzylchloride-divinylbenzene	
	Functionality	quaternary amine	bifunctional quaternary and tertiary amine
Organic solvent limit		0-100% ACN or MeOH for cleaning	
T operating range (°C)		10-40	
pH range		0-14	
max (psi/bar)		4350/300	

Specifications Trap columns

Parameter		Borate ion trap	Amino acid trap
Type		Chemically derivatized polymeric resin	Polymer grafted film on porous polymeric resin
Particle	Material	Polyvinylbenzyl chloride	ethylvinylbenzene-divinylbenzene copolymer
	Diameter (µm)	10	5
	Pore size (Å)	n.d.	Macro-porous, 300
	Crosslinking (%)	12%	55%
	Functionality	polyol	hydroxyethyl quaternary ammonium
Organic solvent limit		0-90% ACN or MeOH for cleaning	0-80% ACN or MeOH for cleaning
T operating range (°C)		10-40	10-40
pH range		0-14	0-14
max (psi/bar)		4350/300	4000/280

Ordering Information

Ordering Information		
Part no.	Description	Additional info
Analytical columns (4.0 and 2.1 mm ID x 20 cm)		
260.0010	SweetSep™ AEX200, 4.0 mm ID x 20 cm, 5 µm	Universal column for separation of mono- to polysaccharides in F&B, plants and glycans.
260.0011	SweetSep™ AEX200, 2.1 mm ID x 20 cm, 5 µm	
260.0020	SweetSep™ AEX20, 4.0 mm ID x 20 cm, 5 µm	Fast, high-resolution separation of monosaccharides from food samples, incl. monosaccharides from glycoproteins, FDG, Heparin, etc.
260.0021	SweetSep™ AEX20, 2.1 mm ID x 20 cm, 5 µm	
Pre-columns (4.0 and 2.1 mm ID x 5 cm)		
260.0015	SweetSep™ AEX200, 4.0 mm ID x 5 cm, 5 µm	For use with the AEX200 analytical column.
260.0016	SweetSep™ AEX200, 2.1 mm ID x 5 cm, 5 µm	
260.0025	SweetSep™ AEX20, 4.0 mm ID x 5 cm, 5 µm	For use with the AEX20 analytical column.
260.0026	SweetSep™ AEX20, 2.1 mm ID x 5 cm, 5 µm	
Trap-columns (4.0 mm and 2.1 mm ID x 5 cm)		
260.0040	Amino acid trap, 4.0 mm ID x 5 cm, 5 µm	Traps amino acids present in the sample that interfere with the monosaccharide separation.
260.0041	Amino acid trap, 2.1 mm x 5 cm, 5 µm	
260.0030	Borate ion trap, 4.0 mm ID x 5 cm, 10 µm	Traps borate contaminants from mobile phase.
260.0031	Borate ion trap, 2.1 mm ID x 5 cm, 10 µm	
Accessories		
260.0100	Pre-column inlet filter PEEK, 0.5 µm	With replaceable PEEK frits 0.5 µm porosity, for direct connection into the analytical or pre-column.
260.0110	Replacement filters PEEK, 0.5 µm, 1 pcs	Replacement filters for Pre-column inlet filter.

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