Sielc

PromixTM

Enter a New Era in Biomolecule Analysis with Promix[™] Columns



"Unsurpassed Selectivity and Peak Capacity for Peptides and Proteins"

Applications:

Proteomics

•Peptide/Protein Analysis

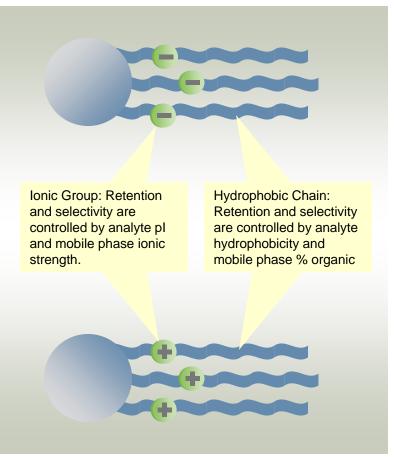
•Peptide/Protein Purification

Features & Benefits:

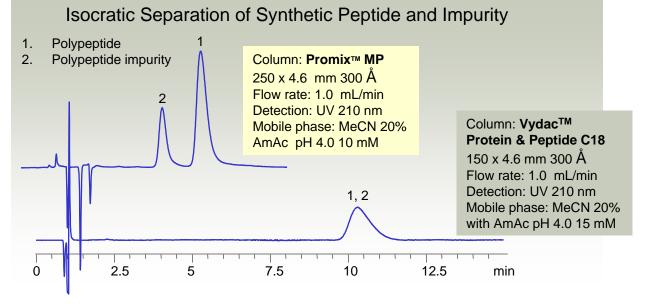
• 2-D HPLC with a Single Column

- Ionic Interactions
- •Reversed-Phase
- Separate peptides and proteins that differ only by a **single amino acid**.
- Enhanced selectivity for closely related peptides and proteins.
- **Increased peak capacity** compared to reversed-phase and ion exchange.
- Increased LC-MS sensitivity using formic acid and ammonium acetate.
- **Completely Scalable** from capillary to prep

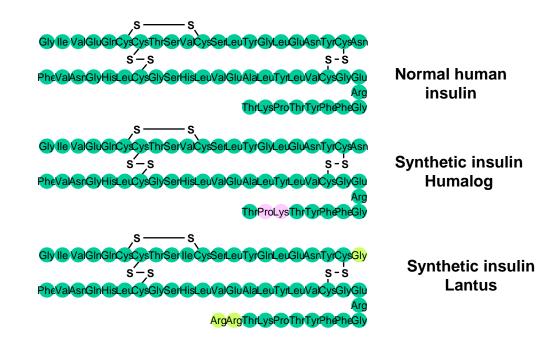
Promix[™] Technology

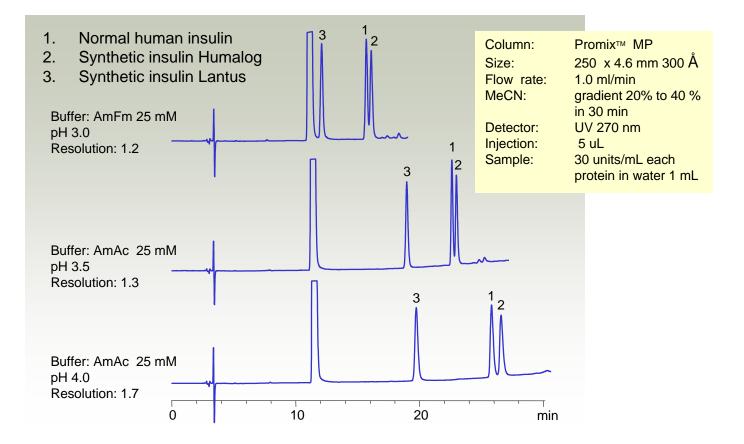


Independent control of acid/buffer concentration and organic modifier offers almost limitless control of retention and selectivity!



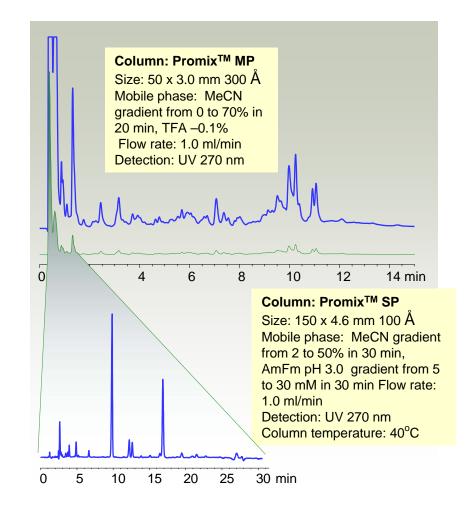
Separation of Insulin Analogs: Closely related peptides can be difficult to separate by reversed-phase. By combining ionic and hydrophobic interactions, Promix MP is able to separate these analogs, differing in sequence by a single amino acid pair, using the slight differences in pl as illustrated below.

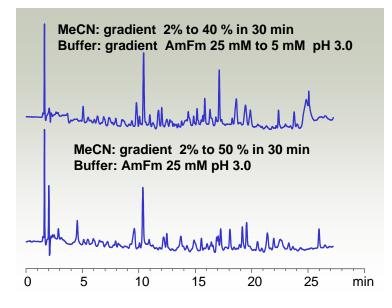




Separation of a Complex Mixture

containing very small peptides and very large peptides can be achieved using two different Promix columns. Very small peptides eluted in first 1.5 min on Promix[™] MP column were collected and resolved using Promix SP column.





Albumin Digest Selectivity Affected by Buffer Concentration:

Complex mixtures such as protein digests have different profiles at different ionic strengths. Multiple gradients can be applied to complex samples:

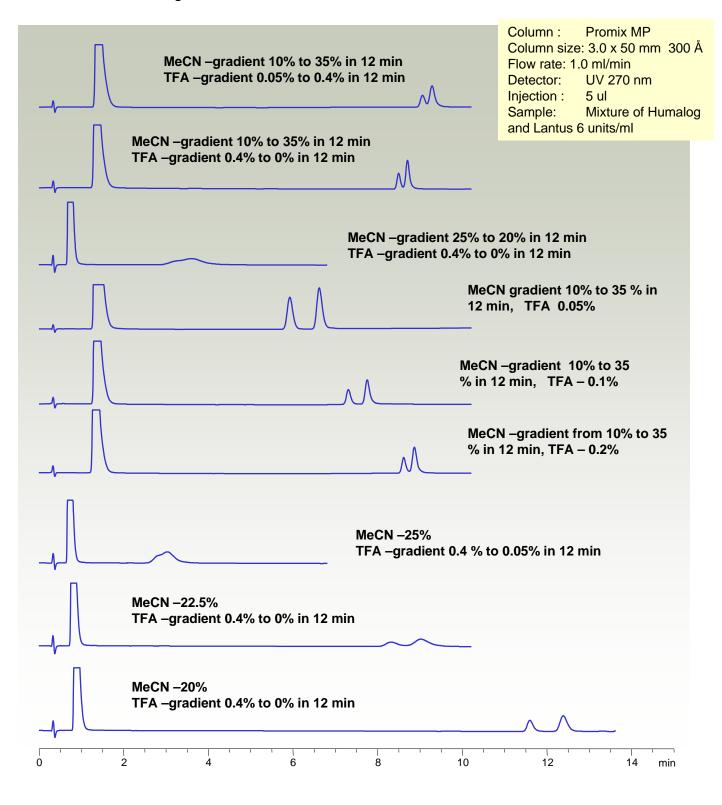
- •organic gradient
- •buffer gradient
- •or both

Each mode provides a different selectivity and elution profile.

Column: Promix[™] MP 150 x 4.6 mm 300 Å Flow rate: 1.0 ml/min Detector: UV 270 nm Injection: 50 uL Sample: 0.3 mg/ml

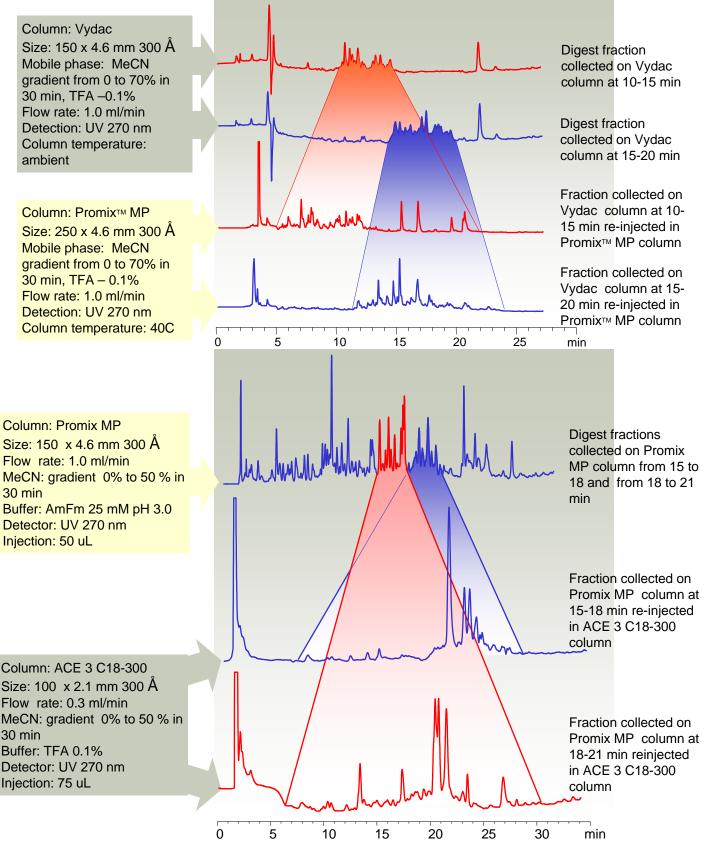
Separation Study of Lantus and Humalog

Similarly to a traditional ion separation, the buffer concentration plays an important role in the mixed-mode technology altering the degree of ionic interaction of the biomolecules with the stationary phase. The amount of the organic modifier is also important in changing the degree of hydrophobic interaction. Independent adjustment of the amount of buffer and organic modifier creates infinite number of separation conditions that are suitable for many types of biomolecules. Generally three types of gradient separation mode available on Promix columns. Gradient of MeCN with isocratic buffer concentration, gradient of buffer concentration with isocratic MeCN concentration, and dual gradient of both MeCN and buffer.



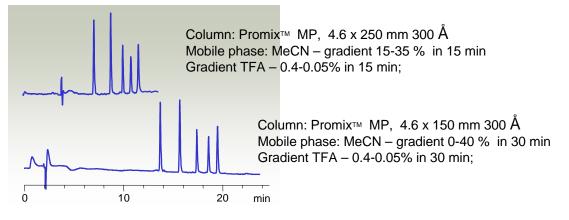
Alternative Selectivity of Promix[™] vs. RP 300Å Columns: An

indication of the alternative selectivity of Promix[™] columns can be observed when the digest fraction collected from a Reversed-Phase column is reinjected onto a Promix[™] column or visa versa. The elution time usually expands providing additional resolution and peak capacity.



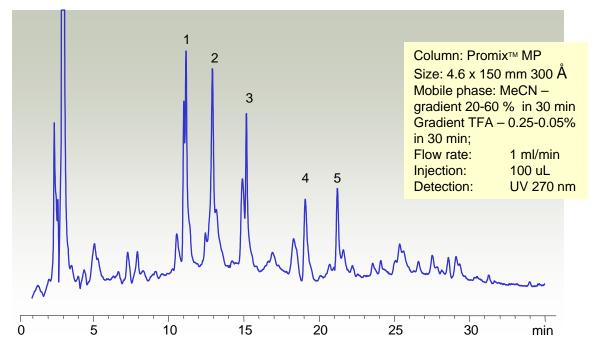
Small Peptides Test Mixture (5 Angiotensins)

- 1. ARG-VAL-TYR-VAL-HIS-PRO-ILE
- 2. ARG-VAL-TYR-VAL-HIS-PRO-PHE
- 3. ALA-PRO-GLY-ASP-ARG-ILE-TYR-VAL-HIS-PRO-PHE
- 4. ASP-ARG-VAL-TYR-VAL-HIS-PRO-PHE-HIS LEU
- 5. ASP-ARG-VAL-TYR-ILE-HIS-PRO-PHE-HIS-LEU



Large Peptides Mixture

- Ribonuclease A (mw 13,500)
- 2. Insulin (mw 5,700)
- 3. Lysozyme (mw 14,300)
- 4. B-Lactalbumin (mw 14,200)
- 5. Carbonic Anhydrase (mw 29,000)

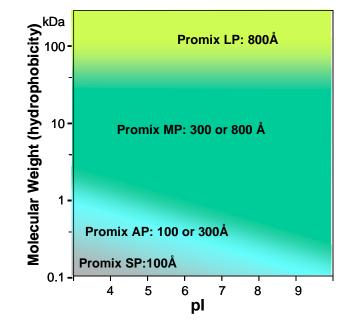


1.

Flow rate: 1.0 mL/min Injection: 40 uL Detection: UV 270 nm

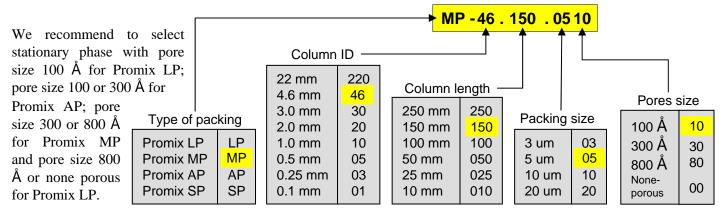
Column Selection and Use

- 1. Use the following chart to select a column based on the analyte's properties
- 2. Select the column length based on sample complexity
 - 150-250 mm for proteomics, protein digests, and complex samples
 - 50-150 mm for synthetic analysis and purification
- 3. Select the column i.d. based on sample loading and sensitivity needs
- 4. Mobile Phase should be pH 2-4 with:
 - TFA (0.05-0.3%)
 - Formic Acid (0.1-0.9%)
 - Ammonium Acetate or Ammonium Formate
- 5. Organic Modifier should be Acetonitrile
- 6. Ionic strength, pH, and organic modifier gradients can be used to optimize the separation



To order: By phone 847-229-2629 By fax 847-655-6079 By mail @sielc.com

Create your Part Number:



Promix[™] Column Selection Chart



formerly Allsep Technologies

For decades liquid chromatography stationary phase design has been dominated by the goal to eliminate multiple, or "unwanted", interactions and to obtain a simple and predictable retention mechanism. Unfortunately, the simplification of the retention process limits the ability to control elution order of the analytes and the scope of available applications this system can be used for. As a response to this limitation, hundreds of different reverse-phase columns were introduced in the last years to cover a variety of analytical situations.

In contrast, PrimesepTM stationary phases were intentionally designed with two major interactions offered on the same column. Both interactions are independently adjustable with mobile-phase composition producing unlimited states of the stationary phase. The hydrophobic interaction is controlled by the amount of organic modifier in the mobile phase (as in any reverse-phase column), while the ion-exchange interaction is controlled by the ion-strength and pH of the mobile phase (as in other ion-exchange columns). This unique property allows using one stationary phase for numerous applications, including analyses of polar and non-polar, ionizable and neutral, organic and inorganic compounds. The behavior of PrimesepTM columns is predictable and reproducible. The method development process is simple and versatile.

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