



TSK-GEL® SP-STAT and CM-STAT Columns

INTRODUCTION

TSK-GEL SP-STAT and TSK-GEL CM-STAT cation exchange columns allow fast equilibration and analysis, as well as isolation, of complex biomolecules. Both TSK-GEL columns are packed with 7 or 10 μm mono-disperse, non-porous resin particles of which the surface consists of an open access network of multi-layered cation exchange groups (see Figure 1). The innovative bonding chemistry, combined with a relatively large particle size, result in a respectable loading capacity and a low operating pressure, attributes not found in traditional mono-disperse, non-porous resins.

PRODUCT HIGHLIGHTS

- Very efficient chromatography for high as well as low MW solutes made possible by novel bonding chemistry and the absence of micro-pores
- High speed and high resolution analysis of biomolecules
- Higher adsorption capacities and lower pressures compared with competitive non-porous columns
- 7 or 10 μm particles for SP and CM chemistries

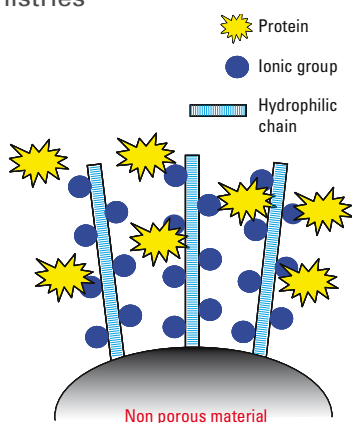


Figure 1

APPLICATIONS

Fast Separations

The fast separation of protein standards was investigated using short cation exchange columns (see Figure 2). A TSKgel SP-STAT column shows superior resolution, better peak shape, and a shorter analysis time (< 60 seconds) compared to a competitive monolithic SP-type column.

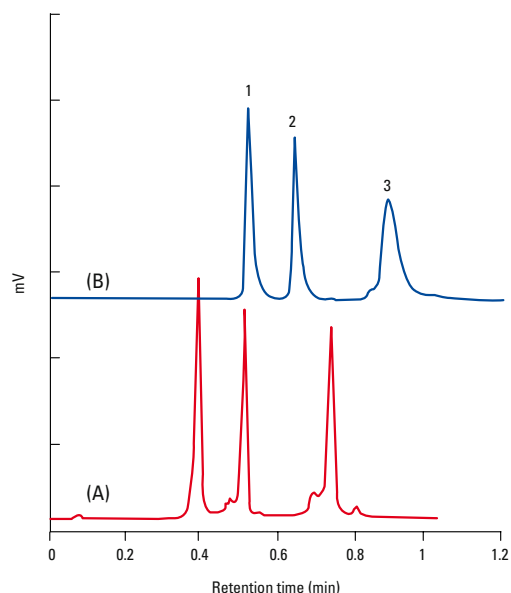


Figure 2

Column: A: TSKgel SP-STAT, 10 μm , 3.0 mm ID x 3.5 cm L
B: Competitor column 4.6 mm ID x 5.0 cm L
Eluent: A: 20 mmol/l sodium acetate (pH 5.0)
B: 1.0 mol/l NaCl in buffer A (pH 5.0) for column A
1.5 mol/l NaCl in buffer A (pH 5.0) for column B
Gradient: 0% B (0 min), 100% B (1 min)
Flow rate: A: 2.0 ml/min
B: 4.73 ml/min
Detection: UV @ 280 nm
Samples: 1. α -chymotrypsinogen A
2. cytochrome C
3. lysozyme

Reaction Monitoring

A sample of β -lactoglobulin (5 mg/mL) was reacted with polyethylene glycol (5 kDa) in a pH 6.5 phosphate buffer. The formation of PEGylated protein reaction products was monitored in 5 minute intervals on a 3.5 cm L TSKgel SP-STAT column. As demonstrated in Figure 3, peak areas of mono-, di-, and tri-PEGylated β -lactoglobulin increased with reaction time, while the area of unreacted β -lactoglobulin declined.

Antibody Analysis

The analysis profiles for five antibodies separated on a TSKgel CM-STAT column were compared with the profiles obtained on a competitive WCX column (Figure 4). Similar or higher resolution profiles were obtained on TSKgel CM-STAT in approximately half the time.

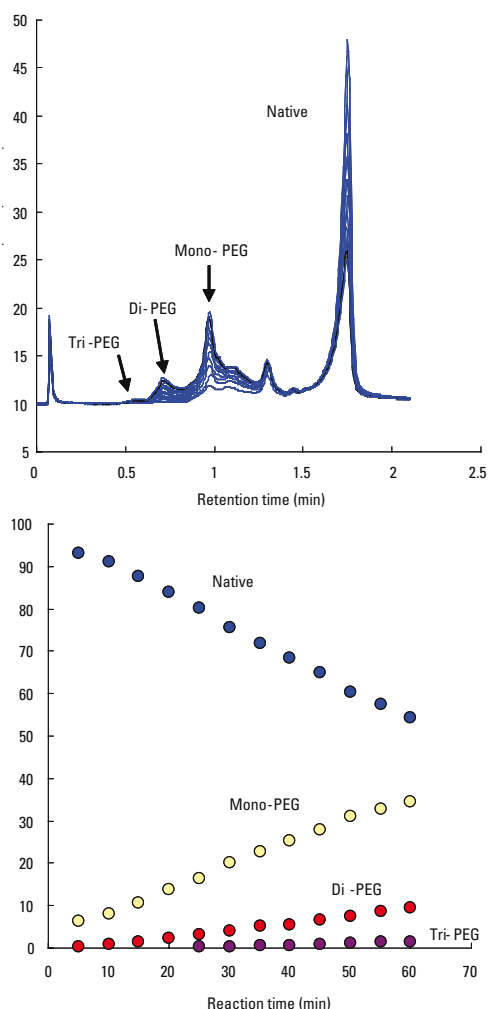


FIGURE 3

Column: TSKgel SP-STAT, 10 μ m, 3.0 mm ID x 3.5 cm L
 Eluent: A: 20 mmol/l sodium acetate (pH 5.0)
 B: 1.0 mol/l NaCl in buffer A (pH 5.0)
 Gradient: 0% B (0 min), 100% B (2 min)
 Flow rate: 2.0 ml/min
 Detection: UV @ 280 nm
 Samples: PEGylated β -lactoglobulin

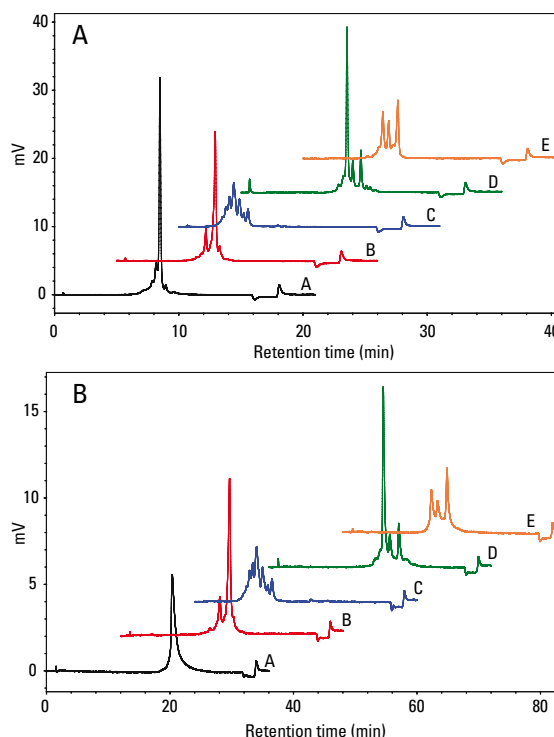


FIGURE 4

Column: A: TSKgel CM-STAT, 7 μ m, 4.6 mm ID x 10 cm L
 B: Competitor WCX, 10 μ m, 4.0 mm ID x 25 cm L
 Eluent: A: 20 mmol/L MES (pH 6.0)
 B: 20 mmol/L MES + 0.5 mol/L NaCl (pH 6.0)
 Gradient: A: 10% B (0 min), 30% B (15 min), 100% B (15 min),
 100% B (17 min), 10% B (17 min), 10% B (21 min)
 B: 10% B (0 min), 30% B (30 min), 100% B (30 min),
 100% B (32 min), 10% B (32 min), 10% B (36 min)
 Flow rate: A: 1.0 mL/min B: 2.0 mL/min
 Temp.: Ambient
 Detection: UV@280nm
 Inj. Vol.: 20 μ L
 Sample: monoclonal antibodies (mAb A through E)

For further details of choice and selection of the TSK-GEL® column that best suits your particular separation needs, please contact us:

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Ordering information

TSKgel STAT COLUMNS

Part-No	Description	Matrix	Housing	Dimensions
21963	TSKgel SP-STAT, 10 μ m	Polymer	Stainless steel	3.0 mm ID x 3.5 cm L
21964	TSKgel SP-STAT, 7 μ m	Polymer	Stainless steel	4.6 mm ID x 10 cm L
21965	TSKgel CM-STAT, 10 μ m	Polymer	Stainless steel	3.0 mm ID x 3.5 cm L
21966	TSKgel CM-STAT, 7 μ m	Polymer	Stainless steel	4.6 mm ID x 10 cm L