

TSKgel® UP-SW2000 UHPLC COLUMNS

TSKgel UP-SW2000 columns packed with 2 µm silica based particles are the latest addition to the popular TSKgel SW series, the gold standard for QC analysis of proteins. The new TSKgel UP-SW2000 UHPLC columns with 12.5 nm pore size expand the existing UP-SW series with a smaller pore size version. They are based on the proven proprietary surface technology of the renowned TSKgel SW series and facilitate the transfer of existing HPLC methods from TSKgel G2000SW/SW_{XL} to UHPLC systems.

Aqueous size exclusion chromatography (SEC) is the method of choice for the analysis of peptides and proteins under non-denaturing conditions. Based on the flow of the sample through a porous stationary phase SEC separates molecules according to their size, or more precisely, their hydrodynamic volume. In aqueous elution systems SEC is also referred to as gel filtration chromatography (GFC). TSKgel SW_{XL} columns have been the industry's standard for quality control of protein biotherapeutics by SEC for decades.

HIGHLIGHTS

- Proven TSKgel SW SEC quality
- Virtual absence of nonspecific interaction
- Easy transfer of existing HPLC methods
- Optimized for peptides, small proteins, and oligonucleotides

TSKgel UP-SW series columns can be used with modern HPLC and UHPLC systems and are available with 15 or 30 cm length. The short version enables short analysis times; the long version provides higher resolution and maximum peak capacity. The lifetime of the columns can be improved when using the corresponding guard columns. A "direct connect" (DC) guard column allows minimizing extra column dead volume.

COMPARISON OF 12.5 nm TSKgel SW COLUMNS

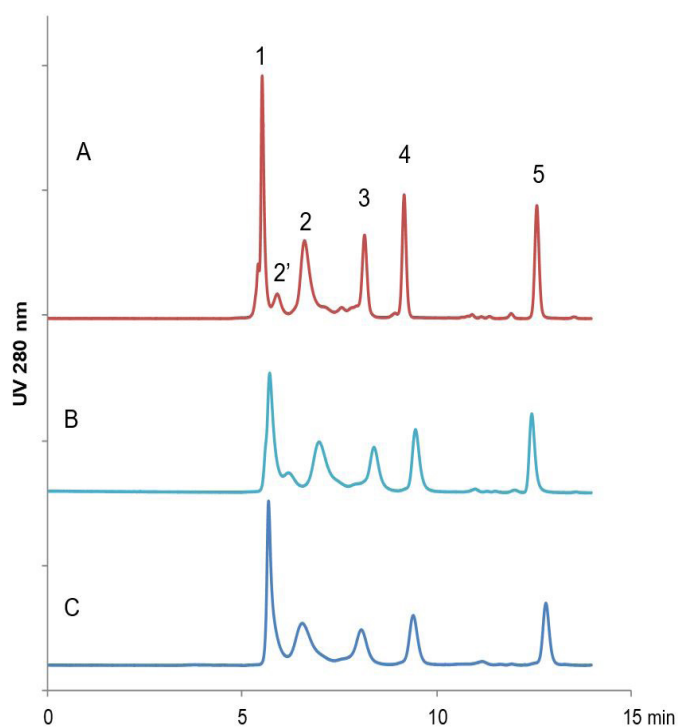


Figure 1

Columns: A) TSKgel UP-SW2000, 2 µm, 4.6 mm ID x 30 cm L
 B) TSKgel SuperSW2000, 4 µm, 4.6 mm ID x 30 cm L
 C) TSKgel G2000SW_{XL}, 5 µm, 7.8 mm ID x 30 cm L

Mobile phase: 100 mmol/L phosphate buffer (pH 6.7)
 + 100 mmol/L sodium sulfate + 0.05 % sodium azide

Flow rate: A), B) 0.35 mL/min, C) 1.0 mL/min
 Temperature: 25 °C
 Detection: UV @ 280 nm
 Injection vol.: 10 µL

Sample: 1. thyroglobulin (640,000 Da);
 2. γ-globulin (155,000 Da); (2' γ-globulin dimer)
 3. ovalbumin (47,000 Da)
 4. ribonuclease A (13,700 Da)
 5. p-aminobenzoic acid (137 Da)



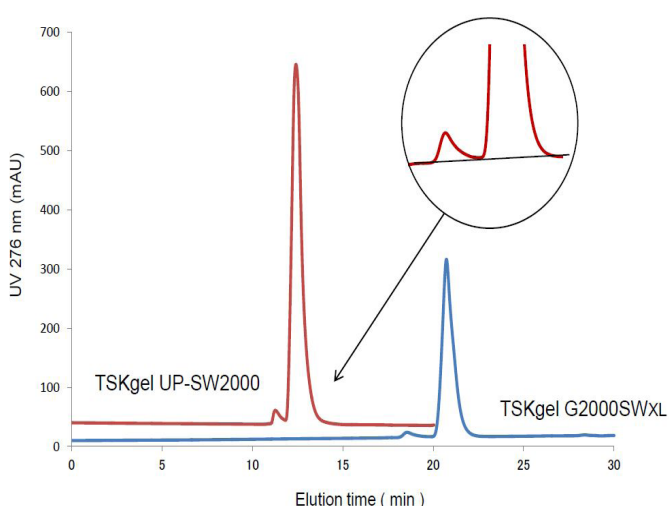
MASS RANGE

The molecular weight range (1-150 kDa) of the 2 µm TSKgel UP-SW2000 is identical to those of 5 micron TSKgel G2000SW_{XL} and 4 micron TSKgel SuperSW2000, which facilitates transfer of existing methods. While providing the same molecular mass separation range the new column has much higher column efficiency: **Figure 1** shows the similarity of the separation range between the 12.5 nm pore size TSKgel SW column portfolio and the increase in resolution achieved by reducing the particle size from 5 (respectively 4) micron to 2 micron.

APPLICATIONS

The separation range of TSKgel UP-SW2000 is ideally suited to analyze peptides and small molecular weight proteins. **Figure 2** shows the analysis of recombinant human insulin with a molecular weight of 5800 Da using the new two micron column in comparison with the analysis using a conventional five micron TSKgel G2000SW_{XL} column. Mobile phase conditions were chosen according to the Ph. Eur. monograph 838.

ANALYSIS OF RECOMBINANT HUMAN INSULIN



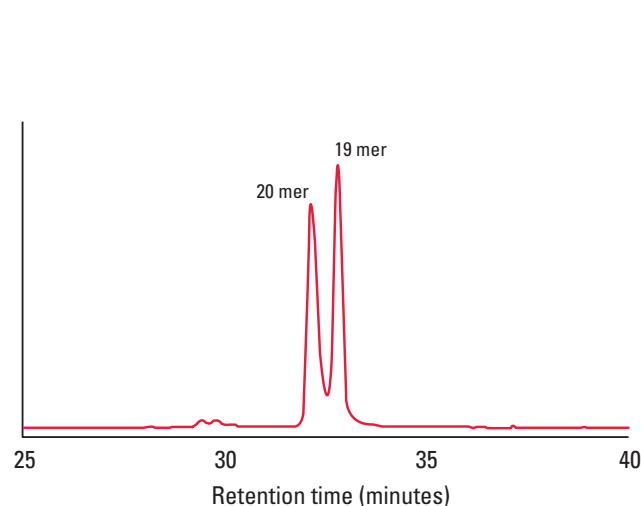
➔ **Figure 2**

Columns: TSKgel UP-SW2000, 2 µm, 4.6 mm ID x 30 cm L
 TSKgel G2000SW_{XL}, 5 µm, 7.8 mm ID x 30 cm L
 Mobile phase: 0.1 % L-arginine/acetonitrile/acetic acid =65/20/15
 Flow rate: 0.2 mL/min (TSKgel UP-SW2000)
 0.5 mL/min (TSKgel G2000SW_{XL})
 Detection: UV@276 nm
 Sample: rec. human insulin

A conventional HPLC system which has been optimized with regard to extra column dead volume was used for both analyzes. There is no need to use a high pressure UHPLC system for TSKgel UP-SW2000, although for any SEC analysis UHPLC systems typically provide ideal technical conditions with regard to system volume, injector and detector technology. Besides a shorter run time the analysis using TSKgel UP-SW2000 delivers sharper peaks.

The increasing importance of oligonucleotides used for therapeutic purposes rises the demand for suitable size exclusion columns. **Figure 3** shows the analysis of a mixture of N and N+1 synthetic oligonucleotides (19 mer and 20 mer) using two TSKgel UP-SW2000 column connected in line. This method achieved a very good separation of the two oligonucleotides.

ANALYSIS OF OLIGONUCLEOTIDES



➔ **Figure 3**

Column: TSKgel UP-SW2000, 2 µm, 4.6 mm ID x 30 cm L x 2
 Mobile phase: 0.05% NaN₃ and 0.3 mol/L NaCl in 0.05 mol/L phosphate, pH 6.7
 Flow rate: 0.2 mL/min
 Detection: UV @ 260 nm
 Temperature: 25 °C
 Sample: 20 mer: 5'-GAATTCATCGGTTTCAGAGAC-3'
 19 mer: 5'-AATTCATCGGTTTCAGAGAC-3'

Ordering information

Part-No	Description	Matrix	Housing	Dimensions
0023515	TSKgel UP-SW2000, 2 µm	Silica	Stainless steel	4.6 mm ID x 15.0 cm L
0023514	TSKgel UP-SW2000, 2 µm	Silica	Stainless steel	4.6 mm ID x 30.0 cm L
0023516	TSKgel Guardcolumn UP-SW2000	Silica	Stainless steel	4.6 mm ID x 2.0 cm L
0023517	TSKgel Guardcolumn UP-SW2000 DC	Silica	Stainless steel	4.6 mm ID x 2.0 cm L