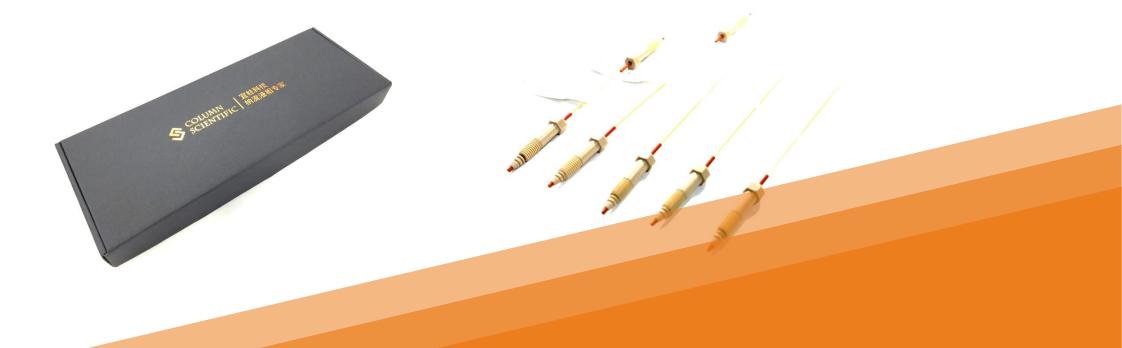


Smartube[®] Benchmark of Nano-LC

For your best ever separation on any Nano-LC system



The state of the art of Nano-LC

Smartube[®] is a high performance chromatographic column product line. It is made for high efficiency, high selectivity microscale separations.

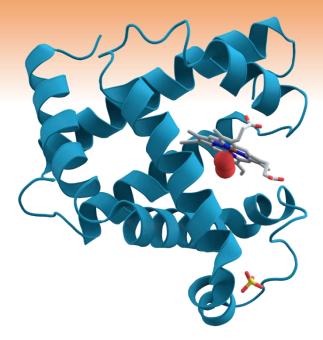
• Compared with other columns, Smartube® columns provide better efficiency, resolution, selectivity, throughput and lifetime.

• The unprecedented separation experience comes from meterlong separation path, excellent column-to-column reproducibility and wide-ranging choice of stationary phase chemistry.



Contents

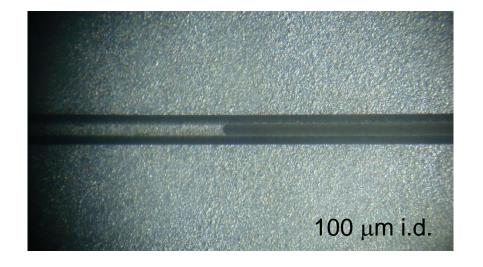
- Page 4 High Pressure Microscale Layer-by-Layer Assembly Technology (HiMLAT)
- Page 5 Nano-LC Columns
- Page 6High Resolution Proteomics
- Page 8 Meter-long Column
- Page 10 Stationary Phase Chemistry
- Page 11Customization & Total Solution



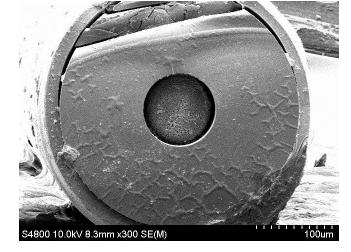
Secret of perfect column

High Pressure Microscale Layer-by-Layer Assembly Technology

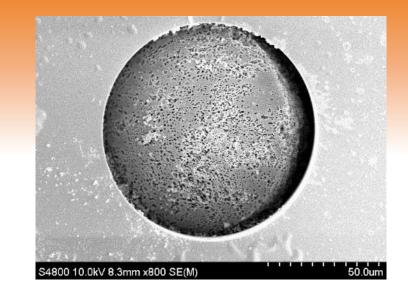
- Optimized slurry solvent to keep the dispersity of packing material
- Highly permeable end frits lead to 1-100 cm column bed
- Super short frit minimizes frit-induced resolution loss
- Layer-by-layer assembly builds up radial- and longitudinal- homogeneous packed bed



Homogeneous packed bed



Micro size highly permeable frit



Permeable frit



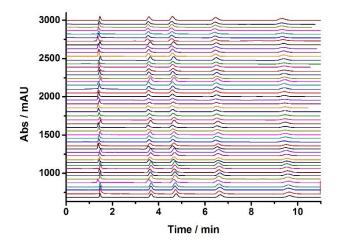
High Efficiency, High Stability, High Reproducibility, Long Lifetime ——Your best ever tool for Nano-LC method development and omics discovery

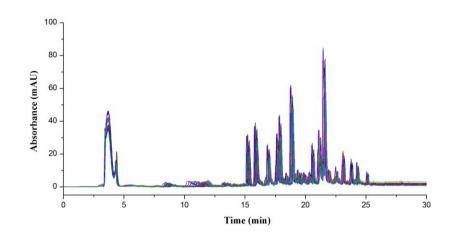
For general Nano-LC runs, **Smartube**® provides 0-50 cm columns for stable, reliable and reproducible separations

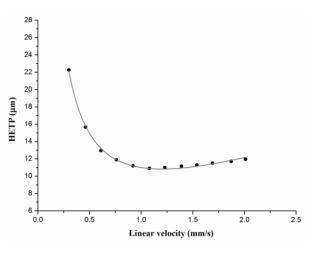
Inter-column reproducibility between 50 columns

Inter-run reproducibility for 35 injections in a week

Excellent efficiency and van Deemter curve







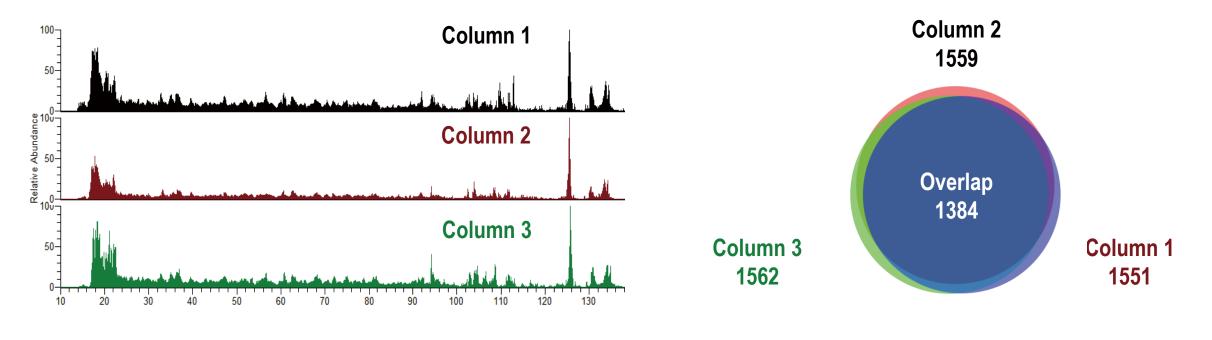
C18 5µm 300Å, 100 mm \times 100 µm i.d. UV214 nm. Analytes: thiourea, methyl-, ethyl-, propyl-, and butylbenzenes. RSD<1%

C18 5 μm 300Å, 150 mm \times 100 μm i.d. mobile phase. UV214 nm. Analytes: Tryptic digest of Cytochrome C

C18 5 μm 300 Å, 100 μm \times 200 mm Vopt = 350 nL/min (1.1 mm/s) $\,$ N = 92000/m H = 10.9 μm

High Efficiency, High Stability, High Reproducibility, Long Lifetime ——Your best ever tool for Nano-LC method development and omics discovery

Unlabeled identification of Hippocampal protein extracts using 3 different Smartube[®] columns

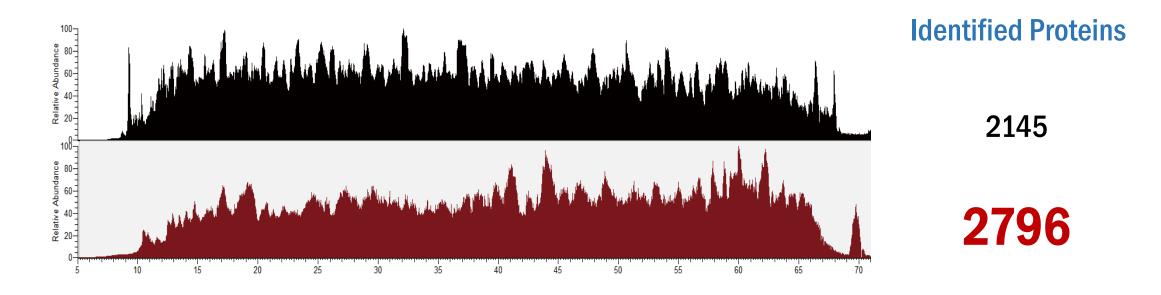


5 µm C18 100 µm * 50 cm gradient time: 140 min

Protein I.D. Coverage

Now you can see more

—Less Operation, More Details, Empower Your Discovery

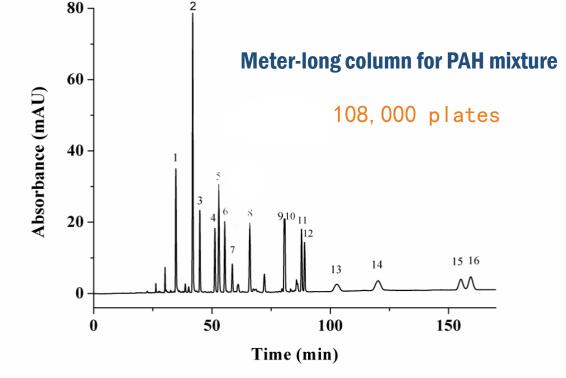


Sample : 293T Cells extracts Column parameters: 1.9 µm C18 15 cm Buffer A: 0.1% (v/v) formic acid (FA) in water ; Buffer B: 0.1% (v/v) FA in acetonitrile (ACN). 7%-22% buffer B for 50 min 22%-35% buffer B for 10 min. The full mass scan was acquired from m/z 350 to 1550



Separameter[®]: Meter-long Column your best tool for super-high resolution separation

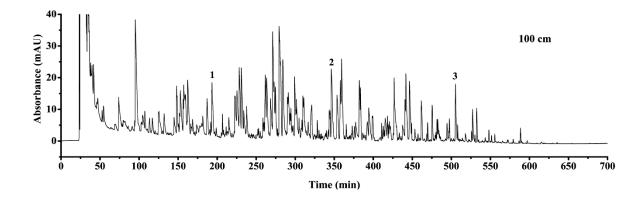
- Limited Sample? High Complexity? Demand Extreme Resolution?
- Meter-long column **Separameter**[®] is your best choice



One Meter is Enough

Meter-long column for protein digest

Peak Capacity 800

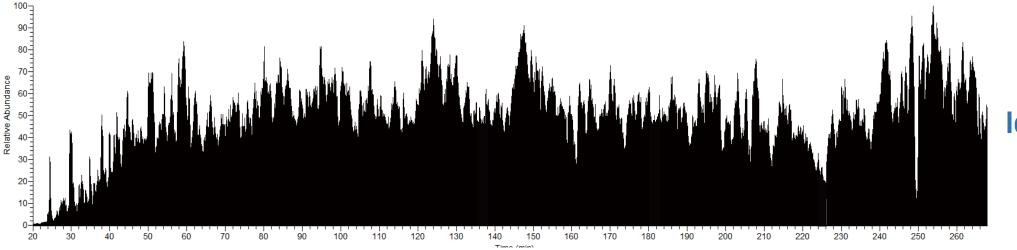


e EPA. (1) acene, (7) C18 5 μ m, 1000 mm \times 100 μ m i.d. UV214nm. Flow Rate: 250nL/min. Mobile phase:A: H₂O (0.05%TFA), B: ACN (0.05%TFA); 5%B-50%B. Sample: Tryptic digest of BSA

C18 5 μ m, 1000 mm \times 100 μ m i.d. UV214nm. Sample: 16 PAHs primary pollutants designated by the EPA. (1) Naphthalene, (2) acenaphthylene, (3) acenaphthene, (4) fluorene, (5) phenanthrene, (6) anthracene, (7) fluoranthene, (8) pyrene, (9) benz[a]anthracene, (10) chrysene, (11) benzo[b]fluoranthene, (12) benzo[k]fluoranthene, (13) benzo[a]pyrene, (14) dibenz[a,h]anthracene, (15) benzo[ghi]perylene, and (16) indeno[1,2,3-cd]pyrene.

Separameter[®]: Meter-long Column your best tool for super-high resolution separation

Sample: 293T Cells extracts Column: Separameter® Gradient time: 240 min

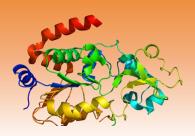


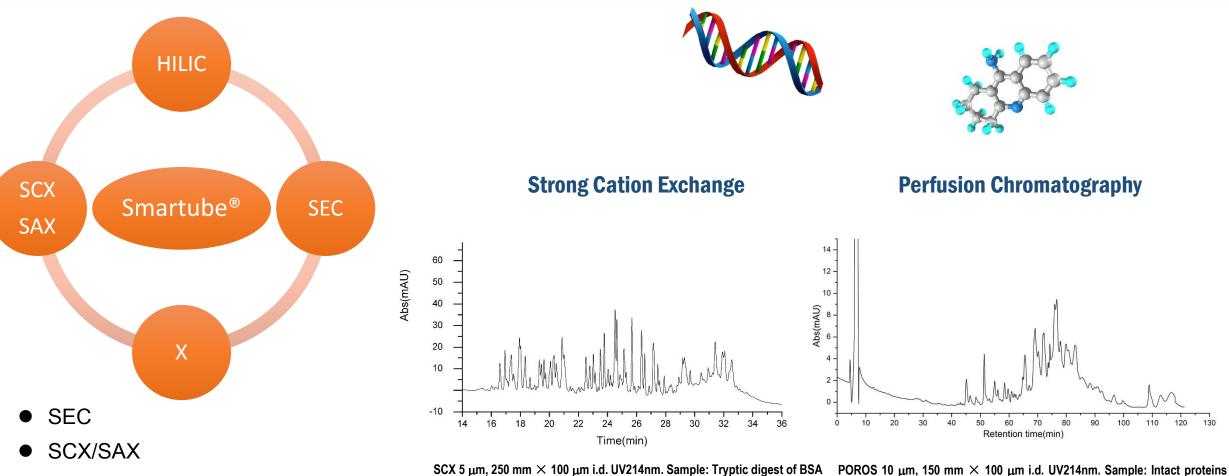
Identified Proteins: 4415

Database: Uniprot fasta Software: Proteome Discoverer (PD) The precursor tolerance: 10 ppm Fragment mass tolerance: 0.02 Da Cysteine carbamidomethylation was set as fixed modification, FDR of PSMs was validated by the Percolator algorithm at 1% based on q-values.

Wide range choice of column selectivity, comprehensively satisfy your needs

Support your microscale separations of biomolecules, pharmaceuticals, environmental pollutants and more





- HILIC
- X = Any phase you like)

from hepatoma cells

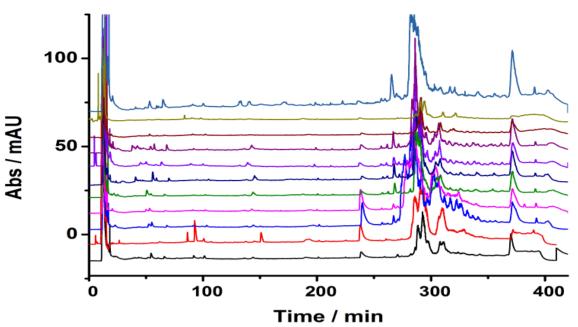
Custom make and total solution

• Specifically made for your needs:

Separation Tools, Purification Strategies, Total Solution

Case Study:

- ✓ Client: Sika deer plant of north China
- \checkmark Project: antler protein fingerprinting







Efficient, Green, Healthy

Column Scientific Inc.

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- Tel: 0592-2185880
- Email: info@columnscientific.com
- Web: www.columnscientific.com