

Introducing . . . Poroshell 300SB-C18

In the world of proteomics and protein chemistry you are creating a revolution in discovery. Now, here is a tool to revolutionize your separations — Poroshell 300SB-C18 (Figure 1). Constructed of a thin layer of porous silica on a solid core (Figure 2), Poroshell 300SB-C18 gives you improved speed, separating proteins in minutes or seconds, with high resolution (Figures 3, 5, 6). Bonded with the patented StableBond C18 phase (Figure 4A) that has given reliable separations, with unmatched column stability (Figure 4B), Poroshell 300SB-C18 offers the ability to separate even at high temperatures with minimal bleed, giving clean MS signals at ultra high speed. Built on a unique particle, Poroshell 300SB also exhibits low backpressure, so you can separate at very high flow rates. Now you have a tool to help conquer proteomics and protein

for Fast, High Resolution Protein Separations

Figure 1
Scanning Electron Micrograph
of Poroshell Particle

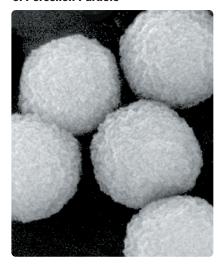
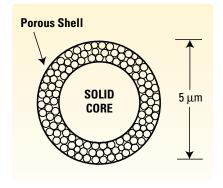


Figure 2 **High Efficiency at High Flow Rates**



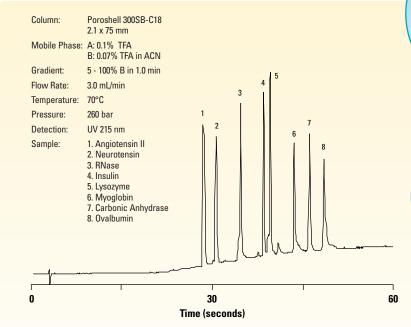
The Poroshell 300SB-C18 particle is a revolutionary chromatography media — superficially porous particles — that produce very fast, high resolution, RP-HPLC separation of proteins, DNA and other macromolecules. Poroshell works well for high resolution of macromolecules because of the rapid mass transfer in and out of the thin porous shell bonded phase. Mass transfer for macromolecules on comparable, completely porous particles, is slower due to the increased path the macromolecule must travel.



The Agilent 1100 LC/MSD

analysis - fast!

Figure 3
Fast, High Resolution Separation of Peptides and Proteins with Poroshell 300SB-C18... in Seconds

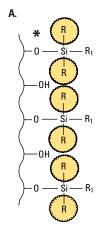


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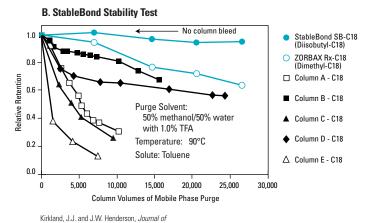
Separation between solutes indicates good peak capacity for rapidly separating complex samples.

Kirkland, J.J., Journal of Chromatographic Science, 38 (2000) 535-544.

Figure 4
Excellent for LC/MS, Poroshell 300SB-C18 Uses StableBond Chemistry to Minimize Column Bleed at Low pH



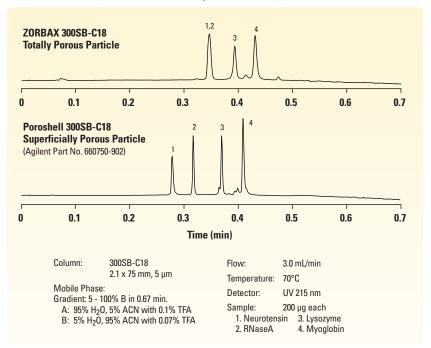
At low pH, reversed phase bonded phases silica break down by hydrolyzing the siloxane bond* (Figure 4A). This breakdown can be almost entirely eliminated by using bulky silanes shown above in StableBond RP-HPLC bonding chemistry.



Chromatographic Science, 32 (1994) 473-480.

Poroshell 300SB-C18 is bonded with StableBond surface chemistry employed in the widely used ZORBAX SB and 300SB columns. StableBond columns can be used at low pH with unsurpassed stability for difficult separations — even at high temperatures.

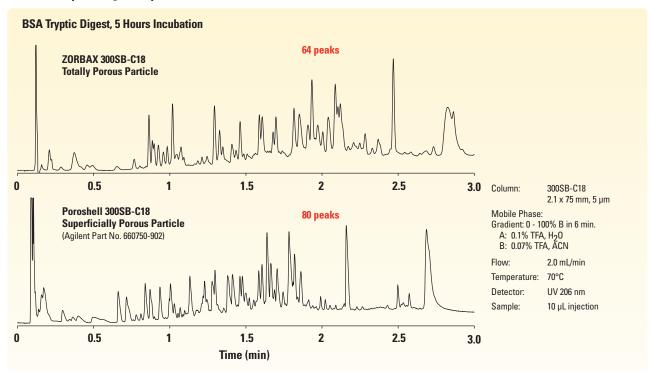
Figure 5
Poroshell Gives High Resolution Compared to Totally Porous Particles





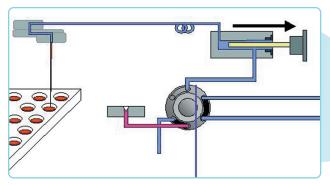
At high flow rates, protein peaks broaden on a totally porous particle, whereas on Poroshell, peaks remain sharp.

Figure 6
Poroshell 300SB Gives Additional Information . . . Fast!
3-Minute Peptide Digest Separation



With peptide digests, Poroshell can separate more peaks at high flow rate.

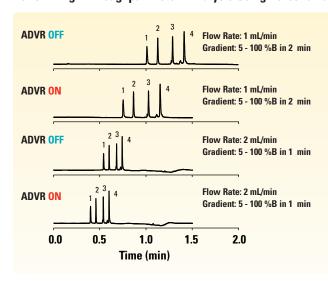
Figure 7 Poroshell Columns Can be Used with the Agilent 1100 Well-Plate Sampler (WPS) with **Overlapped Injection for Even Faster Results**



With Automatic Delay Volume Reduction (ADVR) on, sample draw can occur during a run (overlap mode), where the next sample is loaded during the preceding run.



Perform High-Throughput Protein Analysis Using Poroshell 300SB-C18 and Overlapped Injection



Using the Agilent 1100 well-plate sampler, with Automatic Delay Volume Reduction (ADVR), reduced runtime by approximately 20% in the examples above.

Agilent 1100 DAD Agilent 1100 WPS with ADVR

Column: Poroshell 300SB-C18 2.1 x 75 mm, 5 µm

Mobile Phase:

A: 95% H₂O, 5% ACN with 0.1% TFA B: 5% H₂O, 95% ACN with 0.07% TFA

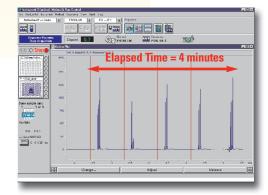
Temperature: 70°C Detector: UV 215 nm

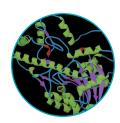
Sample:

1. Neurotensin 3. Lysozyme 2. RNaseA 4. Myoglobin

The default loop wash is 5X the injection volume. Pulse damper varies from 180 to 480 µL depending on pressure.

Instrument-Column Synergy in Optimizing Reversed-Phase HPLC Analysis of Protein and Peptides, ISPPP 2000, Ricker, R.D. and B.E. Boyes, Agilent Technologies, Inc.









Summary of Features, Advantages and Benefits of Poroshell 300SB-C18

Features	Advantages	Benefits	
Unique, 300Å porous shell on solid particle	Fast kinetics	Fast, high resolution protein and other macromolecule separations	
Uses 5 micron particle	Lower backpressure	Fast separations without associated higher pressures	
Uses StableBond chemistry	Extremely stable at low pH	Long column life at low pH	
		Excellent reproducibility for low pH protein separations, column-to-column, batch-to-batch	
	Extremely stable at high temperatures	Faster run times using elevated temperatures	
		Additional selectivity with different temperatures	
Uses high purity Zorbax Rx-Sil sols	Less acidic (fewer metals)	Good peak shape	
	5 μm particle size	High resolution	
	High strength, high-pressure packing	Better resolution	

Don't delay . . .

Place your order TODAY for the most revolutionary RP-HPLC column for proteomics — Poroshell 300SB-C18!

Poroshell 300SB-C18 Column Ordering Information

Column	Size	Particle	Part
Description	(mm)	Size (µm)	Number
Poroshell 300SB-C18 Analytical	2.1 x 75	5	660750-902

To place an order visit our online catalog at www.agilent.com on the World Wide Web, or contact your local Agilent Technologies authorized distributor.

Information, descriptions and specifications in this publication are subject to change without notice.

Call on Agilent Technologies, Inc. to provide your proteomics and protein separation needs. Other significant HPLC column products for protein analysis are porous ZORBAX 300SB (StableBond) C18, C8, C3 and CN, and porous ZORBAX 300Extend-C18.

Poroshell 300SB and all of our ZORBAX columns are the perfect fit for the Agilent 1100 LC and Agilent 1100 Capillary LC.



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