

Evaluation of Sub-2µm Zirconia-PBD Particles for Multi-Modal UHPLC

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Introduction

- Lately, efficiency has received most of the attention in HPLC. As we study and debate optimum particle geometry and instrument design, higher efficiency columns are being adopted by analysts to improve *resolution, peak capacity, speed, sensitivity and solvent economy*.
- Most of the progress with small particles has been made with silica RP columns so it is important to investigate whether the high efficiency observed with ultra-small silica RP particles can be translated to other substrates and phases, which may retain and separate by other selective modes.
- Zirconia phases often separate by a multi-modal mechanism so they are good candidates to see if the performance advantages of sub-2µm particles can be observed (at ambient or elevated temperature) for other packings.



Brief Review of Multi-Modal Zirconia Behavior



- Zirconia substrate exhibits polar and ionic solute interaction: mainly cation-exchange.
- With stable organic coatings, reversed-phase interaction creates Multi-Modal behavior.
- Extreme resistance to temperature, pH and mechanical stress are potential advantages.



Addition of RP Behavior with Coated Zirconia Phases



- Ionic solute retention (and selectivity) is modulated by pH, buffer/salt type and concentrations, and temperature.
- RP solute retention is modulated by organic solvent.
- *Five* important mobile phase variables must be controlled.



Zr-PBD and Si-C18 are Orthogonal for Basic Drugs²

C18 (RP) columns separate mainly by hydrophobic forces and Zr-PBD columns separate by a combination of ionic and hydrophobic forces





LC Conditions: Machine-mixed 80/20 ACN/10 mM ammonium acetate pH=6.7 without pH adjustment; Flow rate, 1.0 mL/min.; Injection volume 0.1 µL; Temperature, 35 °C; Detection at 254 nm; Columns, Zr-PBD, 50 x 4.6 mm i.d. (3 µm particles); Silica-C18 150 x 4.6 mm i.d., (3.5 µm particles).



Difficult Compounds for Silica Often Separate on Zirconia

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Quaternary amines paraquat and diquat are retained and resolved on Zr-PS (also Zr-PBD or bare ZrO₂) due to the cation exchange mechanism; 50% ACN is useful to suppress or regulate retention by RP mode.



column: Discovery C18, 15 cm x 4.6 mm l.D., 3μm mobile phase: 5% acetonitrile in 25 mM phosphate, pH 7 flow rate: 1 mL/min. temp.: 35 °C det.: UV 290 nm

Zirconia-PS: primarily ion-exchange



column: Discovery Zr-PS, 7.5 cm x 4.6 mm, 3µm mobile phase: 50% acetonitrile in 25 mM phosphate, pH 7 flow rate: 3 mL/min. temp.: 65 °C det.: UV 290 nm

Data provided by Sigma-Supelco



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Evaluation of Smaller Diameter Porous Zirconia Particles



1µm 25000X

Specifications:

- Particles produced by a sol-gel process with 1000Å sol
- •Pore diameter 250-300Å
- •Density: 2.6 g/cc (2.5X silica)
- •Surface area: 25 m²/g
- •Particle diameters: 3µm and sub-2µm
- •Totally porous (porosity: 0.45)





Comparison Between Sub-2µm and 3µm Zr-PBD



Efficiency for the larger particle is about right, but the smaller particle should be better (lower H). Could be the phase coating or the column, but instrument dispersion cannot be ruled out with this data.

Sub-2µm Pressure Drop at Different Temperatures*



* 3µm particles (not shown) have about half the pressure drop

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cm/sec); Temp.: 30 °C; UV@254nm; Agilent 1100.



Flow Studies on 3µm Zr-PBD: Alkylbenzenes



Plate height based on van Deemter Equation vs linear velocity at various temperatures for retained solutes: Alkylbenzenes, Temperature: 30 °C, Mobile phase: 50/50 ACN/water, Column: ZirChrom PBD, 50 x 4.6mm, Agilent 1100/UV micro cell (0.007" i.d. tubing).



Plate height based on van Deemter equation vs linear velocity for retained solutes: Alkylbenzenes, Temperature 30 °C, Mobile phase: 50/50 ACN/water (keep k in the same range as 3µm particles), Column: 50 x 4.6mm, Agilent 1100/UV micro cell (0.007" i.d. tubing).

Horvath 2009 Conclusion: systematic investigation of instrument dispersion needed. 15



Flow Studies on Sub-2µm Zr-PBD: Factory Instrument at Ambient



Plate height vs linear velocity for retained solutes: Alkylbenzenes, Temperature 30 °C, Mobile phase: 50/50 ACN/water (keep k' in the same range as 3µm particles), Column: 50 x 4.6mm, Agilent 1100/UV with Standard Cell and 0.007" i.d. tubing.



Plate height vs linear velocity for retained solutes: Alkylbenzenes, Temperature 30 °C, Mobile phase: 50/50 ACN/water (keep k' in the same range as 3µm particles), Column: 50 x 4.6mm, Agilent 1100/UV with Micro Cell and 0.007" i.d. tubing.



Plate height vs linear velocity for retained solutes: Alkylbenzenes, Temperature 30 °C, Mobile phase: 50/50 ACN/water (keep k' in the same range as 3µm particles), Column: 50 x 4.6mm, Agilent 1100/UV with Micro Cell and optimized 0.005" i.d. tubing.



Plate height vs linear velocity for retained solutes: Alkylbenzenes, Temperature 30 °C, Mobile phase: 50/50 ACN/water (keep k' in the same range as 3µm particles), Column: 50 x 4.6mm, Agilent 1100/UV with Micro Cell, high pressure fitting and heat exchanger.

Optimization and Configuration for Elevated Temperature Operation with High-Pressure Column Fitting



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ZirChrom-PBD sub-2µm; Reduced Instrument Volume

Alkylbenzenes











Calibrating Background Pressure Drop for Optimized Agilent 1100

100% H ₂ O at 30 °C		100% H ₂ O at 75 °C	
Flow (mL/min)	BP (bar)	Flow (mL/min)	BP (bar)
1	26	1	21
2	50	2	39
3	77	3	60

* Reference point: Waters Acquity (0.005" ID inlet / 0.0025" ID outlet), 60/40 ACN/water, 0.5 mL/min, background pressure = 1700 psi (113 bar).

Conclusions and Plans for Further HPLC Development with Zirconia

- Performance results with a sub-2µm Zr-PBD column in an Agilent 1100 are encouraging.
- The study of ultra-high speed applications using sub-2µm Zr-PBD, especially at higher pH and temperature ("extreme conditions for silica") will be continued; generic conditions for LC-MS will be investigated.
- Additional advantages of optimizing the IBW of an Agilent Model 1100 HPLC instrument using a high performance (Model 1200) heat exchanger will be studied.
- Other sub-2µm Zr phases (such as CARB) will be prepared and compared to Zr-PBD under ambient and extreme conditions.



References and Acknowledgements

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