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Nota de aplicación

XBridge Prep Columns: Scalability and Loadability for Preparative Separations

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Abstract

This application note details about the efficiency of XBridge Prep Columns.

Benefits

BEH Technology, the second generation of patented organic-inorganic hybrid particle technology (HPT), is the new benchmark for HPLC columns. Waters XBridge Prep Columns reach a new level of maximum loadability and direct scalability.

Introduction

XBridge Columns were designed to be the most pH-stable phases commercially available, while still providing maximum efficiency, peak shape, and robustness. For method development consideration, we offer C_{18} , C_{8} , phenyl and RP_{18} chemistries, available 2.5, 3.5, and 5 μ m particle sizes and dimensions from analytical to prep. XBridge Prep Columns are manufactured with the patent pending Optimum Bed Density (OBD) design, which helps us to achieve direct scale-up from analytical to preparative columns, with the same efficiency and excellent column lifetimes.

Experimental

Scalability

Columns:	XBridge C ₁₈ 5 µm 4.6 x 100 mm; X	(Bridge Pren C
Columns.	Addinge C_{18} 3 µm 4.0 x 100 mm, A	Abridge Frep C

₁₈ 5 µm 19 x 100 mm

Mobile phase A: 10 mM ammonium bicarbonate buffer at pH 10

Mobile phase B: Acetonitrile/100 mM ammonium bicarbonate

buffer, pH 10 (90/10)

Flow rate: 1.06 mL/min (analytical); 18 mL/min

(preparative)

Gradient:	10-min linear from 5% to 95% B	
Injection volume:	30 μL (analytical); 510 μL (preparative)	
Sample:	Econazole and miconazole in DMSO (100 mg/mL each)	
Instrument:	Waters AutoPurification System	
Loadability		
Columns:	XBridge Prep C ₁₈ 5 μm 19 x 50 mm	
Mobile phase A:	0.1% diethylamine in water	
Mobile phase B:	0.1% diethylamine in acetonitrile	
Flow rate:	23.9 mL/min	
Gradient:	8-min linear from 5% to 95% B	
Injection volume:	660 μL	
Sample:	Labetolol (50 mg/mL), quinine (50 mg/mL), diltiazem (50 mg/mL), verapamil (100 mg/mL) and amitriptyline (50 mg/mL) in DMSO	
Instrument:	Waters AutoPurification System	

Results and Discussion

The retention and separation of two antifungal drugs on the analytical XBridge C_{18} Column is shown in

Figure 1A. Under the total load of 6 mg, we observe very symmetric peaks. The mass load was proportionally scaled-up and run on the preparative XBridge Prep C_{18} Column, as shown in Figure 1B. Note the direct scale up, excellent peak shapes, and total mass load of 102 mg.

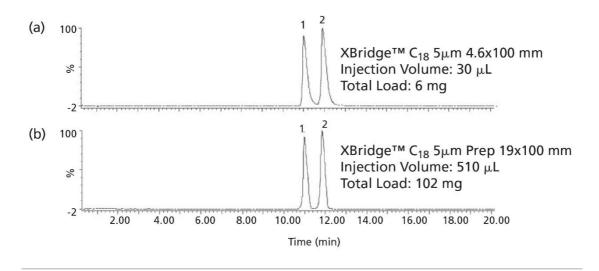


Figure 1. Scale-up of a critical pair of antifungal drugs from analytical to preparative XBridge columns. (A) XBridge C_{18} , 5 μ m 4.6 x 100 mm. (B) XBridge Prep C_{18} 5 μ m 19 x 100 mm. Analytes: (1) econazole, (2) miconazole.

The separation and loadability of five basic analytes on XBridge Prep C_{18} Column under high pH mobile phase conditions is shown in Figure 2. We successfully loaded 198 mg of bases on a 19 x 50 mm column without sacrificing peak shape.

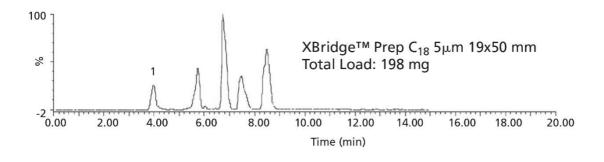


Figure 2. Separation of five basic drugs on XBridge Prep column in high-pH mobile phase.

Analytes in order of elution: labetolol, quinine, diltiazem, verapamil and amitriptyline.

Conclusion

XBridge Prep Columns provide highly efficient separations, direct scale-up, and maximum loadability, crucial for isolation of critical mixture components.

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AutoPurification System https://www.waters.com/10007147

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