

## Selection Guide

Peptides	Polar	Mid & non-polar	Hydrophobic	Natural, Fatty Acids	
< 40AA MW: up to 5KDa					
pH: 1.5 to 8.0	puriFlash® BIO 100 C18N	puriFlash® BIO 100 C18T	Screening Of puriFlash® BIO 100 (C18N /C18T)		
max. pH: 10		puriFlash® BIO 100 C18XS	puriFlash® BIO 100 C18XS		
< 80AA MW: up to 10KDa					
pH: 1.5 to 8.0	puriFlash® BIO 200 C18N	puriFlash® BIO 200 C18T	Screening Of puriFlash® BIO 200 (C18N /C18T)		
max. pH: 10		puriFlash® BIO 200 C18XS	puriFlash® BIO 200 C18XS		
< 160AA MW: up to 20KDa					
pH: 1.5 to 8.0	puriFlash® BIO 200 C18N	puriFlash® BIO 200 C8N	puriFlash® BIO 200 C8N		
< 80AA MW: up to 100KDa					
pH: 1.5 to 8.0				puriFlash® BIO 300 C4AQ	
In-Process QA/QC of	Peptides Synthesis		In-Process QA/QC of Peptides Synthesis puriFlash® BIO CS 2.6C18N => puriFlash® BIO 100 2.5C18N		

### Notes

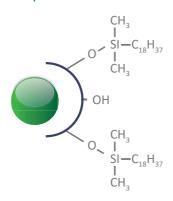
Polar Peptides => Hilic mode using higher % of ACN 95 -to- 85%

Hydrophobic Peptides => it is useful to work with Water/ACN using a few % Formic Acid or 0.05% TFA ~ pH 2. In case your peptides have Lysine, Arginine etc. it is better to have an alki environment in the solvent. You need real buffer and according to buffer solubility it is to suggest to switch to MeOH instead of ACN. Usually step-Gradients (Ramp Gradients) or Pseudo-Isocratic or very flat gradients lead to highest capacity.

# **Stationary Phases & Columns**

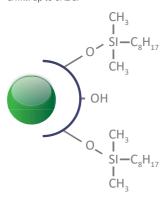
## **BIO-Stationary Phases - Peptides Purification**

## **Peptides**



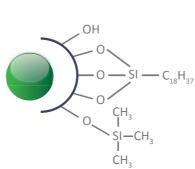
## puriFlash® BIO C18-N

100Å - 320m²/g 2.5, 3.5, 5, 10, 15 & 30µm C18 - octadecyl Mono-functional %C: 15.0 End-capping: None pH stability: 1.5 to 8.0 Use mode: Reverse In-Process QA/QC of Peptides Synthesis. Analysis & Purification of polar Peptides with les than 40AA & mw. up to 5KDa under pseudo hilic mode with 85% -to- 95% ACN. Analysis & Purification of hydrophobic Peptides with less than 40AA & mw. up to 5KDa.



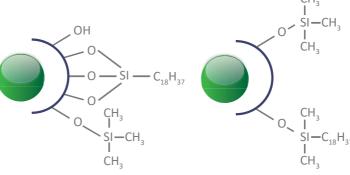
### puriFlash® BIO C8-N

200Å - 200m<sup>2</sup>/g 2.5, 3.5, 5, 10, 15 & 30µm C18 - octadecyl Mono-functional %C: 7.0 End-capping: None pH stability: 1.5 to 8.0 Use mode: Reverse Analysis & Purification of polar Peptides less than 160AA & mw. up to 20KDa under pseudo hilic mode with 85% -to- 95% ACN. Analysis & Purification of hydrophobic Peptides with less than 80AA & mw. up to 10KDa.

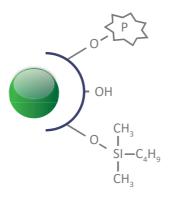


## puriFlash® BIO C18-T

100Å - 320m<sup>2</sup>/g 2.5, 3.5, 5, 10, 15 & 30µm C18 - octadecyl Tri-functional %C: 17.0 End-capping: One-step pH stability: 1.5 to 8.0 Use mode: Reverse Analysis & Purification of mid & non-polar Peptides, hydrophobic Peptides with less than 40AA & mw. up to 5KDa.



puriFlash® BIO C18-XS 100Å - 320m<sup>2</sup>/g 2.5, 3.5, 5, 10, 15 & 30µm C18 - octadecyl Mono-functional %C: 17.0 End-capping: Multi-step pH stability: 1.0 to 10.0 Use mode: Reverse Analysis & Purification of mid & non-polar Peptides, hydrophobic Peptides with less than 40AA & mw. up to 5KDa under basic conditions up to pH: 10.0



### puriFlash® BIO C4-AQ

300Å - 100m²/g 3.5, 5, 10, 15 & 30µm C4 - butyl Mono-functional %C: 3.0 End-capping: Mixte pH stability: 1.5 to 8.0 Use mode: Reverse Analysis & Purification of natural Peptides, fatty acids with larger than 80AA & mw. up to 100KDa.

