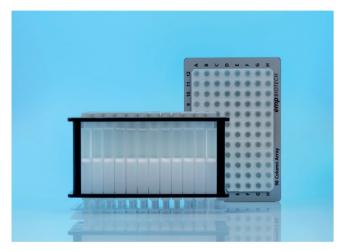
CentriPure 96

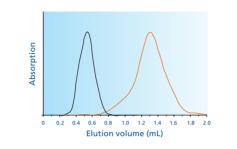
Gel Filtration Column Array 300-Z25M for rapid purification, desalting, and buffer exchange of:

- Nucleic acids
- Proteins and antibodies

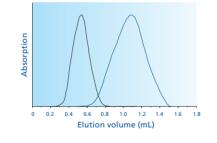


- Purifies samples of 150-300 µL into a final volume of 300-500 µL.
- Contains Zetadex-25 size exclusion resin.
- Simultaneously processes 96 samples.
- Designed specifically for automated systems using standard SLAS microplate footprint.

Typical application examples (proteins):



Elution profile overlay of ovalbumin (1 mg/mL) and free dye (TAMRA, 0,1 µmol) in a 200 µL sample volume.



Desalting of protein solution (1 mg albumin (OvA) in 1 mL 0.8 M NaCl), elution with water (200 µL sample volume)

CentriPure Gel Filtration Column Arrays

Prod. Code: CP-0581

CentriPure 96 Column Array 400-Z25M Gel filtration column array simultaneously processes 96 samples of 400 µL sample volume

Prod. Code: CP-0582

CentriPure 96 Column Array 500-Z25M Gel filtration column array simultaneously processes 96 samples of 500 µL sample volume

coming soon:

CentriPure 24 Column Array CentriPure 48 Column Array

Hazard and Precautionary Statements				
()	WARNING			
H317	Contains ProClin™ 150. May cause an allergic skin reaction.			
P262	Do not get in eyes, on skin, or on clothing.			
P280	Wear protective gloves, protective clothing, eye and face protection.			
P302/P352	If on skin: Gently wash with soap and water.			
P305/P338/P351	If in eyes: Rinse cautiously with water for several minutes. If possible, remove contact lenses and continue rinsing.			

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> emp BIOTECH GmbH is an ISO 9001:2015 certified company registration number 011001300789 (TÜV Rheinland)



Prod. Code: CP-0580

CentriPure 96

Gel Filtration Column Array 300-Z25M for rapid purification, desalting, and buffer exchange



For rapid desalting or buffer exchange

- of oligonucleotides longer than 10 bp/nt
- of proteins larger than 5 kDa
- of spheroidal nanoparticles greater than 2 nm Ø

Purifies samples of 150-300 μL into a final volume of 300-500 $\mu L.$

Instructions for use

1. Column Preparation

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Carefully remove the top mat and then remove the bottom mat of the **CentriPure 96** Column Array. The bottom mat may be replaced to store unused columns when less than 96 columns are processed.

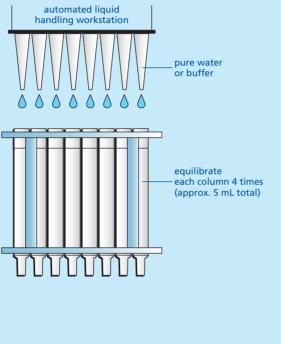
Allow excess column fluid to drain (via gravity) into a suitable waste reservoir.

If all 96 columns are processed simultaneously, a slightly reduced pressure may be used with a manifold to speed up this process.

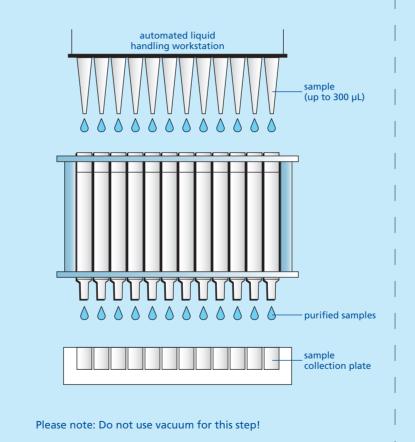
Please note: The Array is not intended to be centrifuged!



3. Sample Application and Elution



Choose a buffer for your specific application and use this same buffer for both equilibration and elution steps. To equilibrate the column, allow the buffer to enter the gel bed completely and to drain into a suitable waste reservoir.



Transfer your sample (up to $300 \ \mu$ L) to the **CentriPure 96 Column Array**. Allow the sample to enter the gel bed completely using a suitable waste reservoir. Do not use vacuum for sample application.

Using the chart below, determine the pre-run volume and elution volume for your sample size. Transfer the pre-run volume to each column and let it completely enter the gel bed without vacuum.

Place a collection plate for sample collection under the **CentriPure 96 Column Array**. Add the elution volume to the column and elute the purified sample.

Sample volume	Pre-run volume	Elution volume	Protein recovery*	Salt removed
150 µL	200 µL	300 µL	98 %	99.9 %
200 µL	150 µL	350 µL	98 %	99.6 %
250 µL	100 µL	400 µL	98 %	99.6 %
300 µL	0 µL	500 µL	98 %	98.9 %

* determined using 1 mg/mL ovalbumin in 0.8 M NaCl

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