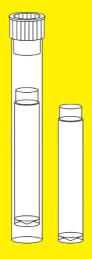
Instructions for Use

Vivaspin® Filtrate

Centrifugal Ultrafiltration Devices for General Laboratory Use



3104669-001-00





Contents

1	Abo	ut thes	e Instructions	
	1.1	Scope	e	
	1.2		t Groups	
	1.3		ools Used	
		1.3.1	Warnings in Operation Descriptions	
		1.3.2	Other Symbols	
2	Safe	ty Instr	ructions	
	2.1	Gene	ral Functions	
	2.2	Perso	nnel Qualification	
	2.3	Signif	icance of these Instructions	
	2.4	_	tionality of the Product	
3	Proc	luct De	escription	9
	3.1		uct Overview	
	3.2	Produ	uct Symbols	10
4	Proc	ess Pre	eparation	11
	4.1	Scope	e of Delivery	
	4.2	Unpa	cking	
5	Оре	ration.		12
	5.1	Pre-R	Rinsing the Product	
	5.2	Saniti	zing the Product	
	5.3	Perfo	rming Filtration	
		5.3.1	Applying the Sample	
		5.3.2	Performing Filtration	
		5.3.3	Removing the Sample	
6	Stor	age		16
	61	Storin	na the Product	16

Contents

7	Disp	osal	17
	7.1	Decontaminating the Product	17
	7.2	Disposing of the Product	17
8	Tech	inical Specifications	18
	8.1	Dimensions	18
	8.2	Materials	18
	8.3	Ambient Conditions	19
	8.4	Equipment Required	19
		8.4.1 Centrifuges	19
		8.4.2 Pipettes	19
	8.5	Operating Conditions	20
	8.6	Sanitizing Methods	20
	8.7	Typical Performance Characteristics	21

1 About these Instructions

1.1 Scope

These instructions are part of the product. These instructions apply to the following versions of the product:

Vivaspin® Filtrate	Quantity	Prod. no.
5 kDa CTA	12	13229E
10 kDa CTA	12	13239E
20 kDa CTA	12	13249E
100 kDa PES	12	13269GE
300 kDa PES	12	13279E

1.2 Target Groups

The instructions are addressed to the following target groups. The target groups must possess the knowledge specified below.

Target Group	Knowledge and Qualifications
Operator	The operator is familiar with the device and the associated work processes. The operator understands the hazards which may arise when working with the device, and knows how to prevent them.

1.3 Symbols Used

1.3.1 Warnings in Operation Descriptions

NOTICE

Denotes a hazard that may result in property damage if it is **not** avoided.

1.3.2 Other Symbols

- Required action: Describes actions that must be carried out.
 The actions in the sequence must be carried out in succession.
- Result: Describes the result of the actions carried out.

2 Safety Instructions

2.1 General Functions

The product is intended for the ultrafiltration of biological and aqueous solutions with small sample volumes, e.g. for protein removal.

The filtration process must be carried out in a centrifuge. For this purpose, the sample solution must be put into the product and the product must be inserted into a centrifuge. Due to the centrifugal forces, the particles or macromolecules that are sufficiently larger than the nominal pore size of the membrane are removed from the sample solution.

The product is supplied non-sterile. It is intended for single use and must be disposed of after one use.

The product is intended exclusively for use in accordance with these instructions. Any further use beyond this is considered improper.

Operating Conditions for the Product

The product is intended for general laboratory use.

The product may only be used with the equipment and under the operating conditions described in the Technical Data section of these instructions.

2.2 Personnel Qualification

Persons without sufficient knowledge in the safe use of the device can injure themselves and others.

If a specific qualification is required for an activity: The target group is indicated. If no qualification is specified: The activity can be carried out by the target group "operator".

2.3 Significance of these Instructions

Failure to follow the instructions might have serious consequences, e.g. danger to individuals.

- ▶ Read the instructions carefully and completely. The instructions for action build on each other.
- ► Ensure that the information contained in these instructions is available to all individuals working with the product.

2.4 Functionality of the Product

A damaged product or worn parts can lead to malfunctions or cause hazards which are difficult to identify.

Only operate the product when it is safe and in perfect working order.

3 Product Description

3.1 Product Overview

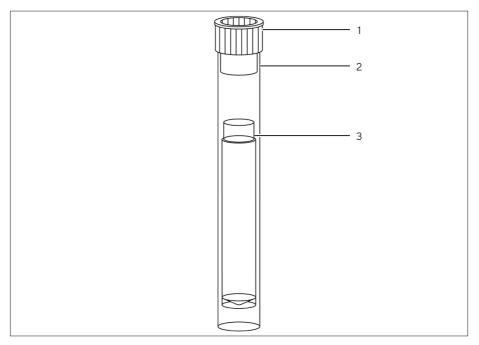


Fig. 1: Product overview (example)

Pos.	Description
1	Dust cap
2	Centrifuge tube
3	Filtrate vessel with membrane

3.2 Product Symbols

Symbol	Description
Qty	Quantity
REF	Catalogue number
LOT	Batch number
Σ	Use by
NON	Non-sterile product
<u> </u>	Do not re-use
<u> </u>	Consult instructions for use
*	Temperature limits

4 Process Preparation

4.1 Scope of Delivery

Item	Quantity
Product, packed in a cardboard box, according to packaging unit	12
Instructions for Use	1

4.2 Unpacking

- ▶ NOTICE Risk of product malfunctions due to exceeding the usability! Check the usability of the product (see specification on packaging). Dispose of products for which the usability has been exceeded.
- ▶ Unpack the product.

5 Operation

5.1 Pre-Rinsing the Product

The membrane in the product may contain traces of glycerin. If this substance can interfere with the analysis of the sample: Rinse the membrane before filtration.

- Remove the dust cap.
- ➤ Slide out the filtrate vessel and place it with the open end downwards. Do **not** touch the membrane.
- ▶ Dip the filtrate vessel with the membrane facing downwards in buffer solution or deionized water for 5 minutes.
- ► Alternatively, use a pipette to apply a filling volume of buffer solution or deionized water into the centrifuge tube.
- Slide the filtrate vessel into the centrifuge tube with the membrane facing downwards.
- ► Wash the buffer solution or deionized water through the membrane with the help of a suitable centrifuge.
- ▶ Decant the buffer solution or deionized water from the centrifuge tube and filtrate vessel.
- ▶ If the pre-rinsed product is not used immediately: Cover the surface of the membrane with buffer solution or water, replace the dust cap to avoid evaporation and store the product in the refrigerator. The membrane must not dry out.

5.2 Sanitizing the Product

The product can be sanitized before use. The sanitizing method must be suitable for the product (see Chapter "8.6 Sanitizing Methods", page 20).

Procedure

- Remove the dust cap.
- ▶ Sanitize the product using the desired sanitizing method.
- ► Empty the product.

5.3 Performing Filtration

5.3.1 Applying the Sample

It is recommended that a pipette is used to apply the sample into the product. The pipette must be compatible with the product (see Chapter "8.4.2 Pipettes", page 19).

Please ensure that the molecular weight cut-off (MWCO) of the product is suitable for the size of the target molecule(s) to be removed or concentrated. In order to ensure maximum removal or recovery of the target molecule, it is recommended to select a MWCO that is at least 50% below the size of the target molecule.

NOTICE

Risk of product malfunctions due to using unsuitable samples!

▶ Do not apply samples containing organic solvents into the product.

NOTICE

Risk of product malfunctions or damage to the centrifuge due to exceeding the maximum filling volume!

▶ Do not exceed the maximum filling volume (see Chapter "8.5 Operating Conditions", page 20).

Procedure

- ▶ Check whether the MWCO of the product is suitable for the application.
- Remove and discard the dust cap.
- ► Slide out the filtrate vessel and place it with the open end downwards. Do not touch the membrane.
- ▶ Apply the sample into the centrifuge tube using a pipette. Comply with the maximum filling volume. For non-coagulated blood samples, pre-centrifuge the sample for 2 minutes without the filtrate vessel.
- ➤ Slide the filtrate vessel into the centrifuge tube with the membrane facing downwards and leave for 5 minutes so that the membrane is wetted by the sample.

5.3.2 Performing Filtration

- ► Insert the product into the centrifuge. For samples containing lipids (e.g. milk), only use a fixed angle rotor.
- ▶ NOTICE Risk of product malfunctions or damage to the centrifuge. Comply with the approved centrifugation limit values (see Chapter "8.5 Operating Conditions", page 20).
- ➤ Centrifuge the product in the centrifuge until the desired filtrate volume or concentration level is achieved. For concentrated protein samples (e.g. blood, serum), start the filtration at half the maximum RCF for 5 minutes.

5.3.3 Removing the Sample

- ▶ If the filtration or concentration is complete: Remove the product from the centrifuge.
- ▶ NOTICE Risk of reduced sample volume due to diffusion. Remove the sample from the filtrate vessel with a pipette as soon as possible after performing filtration.
- ▶ If the concentrate is required: Remove the filtrate vessel from the centrifuge tube using opened forceps. Use a pipette to remove the concentrate from the centrifuge tube.
- ▶ If the membrane was pre-rinsed before filtration: Decant the filtrate and concentrate.

6 Storage

6.1 Storing the Product

If the product has been unpacked and the membrane has been pre-rinsed: The membrane must be protected against drying out. For this purpose, the membrane must be stored in a moist and cool condition.

NOTICE

Risk of damage to the product due to improper storage!

Comply with the storage specifications.

- ▶ If the product is packaged: Store the product in the packaging.
- ▶ If the product has been unpacked and membrane has been pre-rinsed:
 - ▶ Remove the dust cap.
 - ► Cover the membrane with buffer solution or water.
 - ► Replace the dust cap.
- ➤ Store the product according to the ambient conditions (see Chapter "8.3 Ambient Conditions", page 19).

7 Disposal

7.1 Decontaminating the Product

If the product has come into contact with hazardous substances: Steps must be taken to ensure proper decontamination and declaration. The operator of the product is responsible for adhering to local government regulations on the proper decontamination and declaration for transport and disposal.

Procedure

► If the product has come into contact with hazardous substances: Decontaminate the product.

7.2 Disposing of the Product

The product must be disposed of properly. The packaging is made of environmentally friendly materials that can be used as secondary raw materials.

Requirements

The product must be decontaminated.

- ▶ Dispose of the product in accordance with local government regulations.
- ▶ Dispose of the packaging in accordance with local government regulations.

8 Technical Specifications

8.1 Dimensions

	Unit	Value	
Dimensions			
Length × Diameter	mm	93 × 14	
Active membrane surface	cm²	0.79	
Weight	9	10	

8.2 Materials

	Materials
Filtrate vessel	Styrene Acrylonitrile (SAN)
Centrifuge tube	Polystyrene (PS)
Dust cap	Polyethylene (PE)
Membrane	Cellulose Triacetate (CTA) or Polyethersulfone (PES)

8.3 Ambient Conditions

	Unit	Value
Storage temperature		
When packed	°C	+15 - +30
When unpacked, with membrane kept moist	°C	+2 - +8

8.4 Equipment Required

8.4.1 Centrifuges

	Unit	Value
Swing bucket rotor or fixed angle rotor		
Minimum rotor angle for fixed angle rotor		25°
Rotor accepting centrifuge tubes with the following features		
Volume	mL	15
Diameter	mm	17
Base		Conical or flat

8.4.2 Pipettes

Pasteur pipette, variable volume or fixed volume pipette for sample application and concentrate or filtrate retrieval.

8.5 Operating Conditions

	Unit	Value
Filtration Volumes		
Filling volume, minimum	mL	0.5
Filling volume, maximum	mL	2.5
Membrane hold-up volume, minimum	μL	<5
Dead stop volume ¹	μL	100
Relative Centrifugal Force, maximum		
Centrifuge with swing bucket rotor	g	2,500
Centrifuge with fixed angle rotor	g	2,000

¹The dead stop volume may vary depending on the type and concentration of the sample, operating temperature and | or centrifuge rotor

8.6 Sanitizing Methods

Rinsing with 70% ethanol or with sanitizing gas mixture, e.g. ethylene oxide

Not suitable for autoclaving

8.7 Typical Performance Characteristics

2.5 mL start volume in centrifuge at 2,000 g	Time to filter 50 % of sample volume	Time to filter 90 % of sample volume	Passage of sample species	
BSA 1.0 mg/ml (66 kDa)			
5 kDa MWCO CTA	300 min	-	0 %	
10 kDa MWCO CTA	35 min	80 min	2 %	
30 kDa MWCO CTA	9 min	20 min	2 %	
Blue dextran 0.1 mg/ml (2,000 kDa)				
300 kDa MWCO PES	9 min	25 min	28 %	

Sartorius Stedim Lab Ltd. Sperry Way, Stonehouse GL10 3UT, UK

Phone: +44 1453 821972 www.sartorius.com

The information and figures contained in these instructions correspond to the version date specified below.

Sartorius reserves the right to make changes to the technology, features, specifications and design of the equipment without notice.

Masculine or feminine forms are used to facilitate legibility in these instructions and always simultaneously denote all genders.

Copyright notice:

These instructions, including all components, are protected by copyright.

Any use beyond the limits of the copyright law is not permitted without our approval.

This applies in particular to reprinting, translation and editing irrespective of the type of media used.

Last updated:

02 | 2023

© 2023 Sartorius Stedim Lab Ltd. Sperry Way, Stonehouse GL10 3UT, UK

KS | Publication No.: SE-6002-e230201