

User Guide

Amicon® Ultra-0.5 Centrifugal Filter Devices

for volumes up to 500 μL

For research use only; not for use in diagnostic procedures



Introduction

Amicon® Ultra-0.5 centrifugal filter devices provide fast ultrafiltration, with the capability for high concentration factors and easy concentrate recovery from dilute and complex sample matrices. The vertical design and available membrane surface area provide fast sample processing, high sample recovery (typically greater than 90% of dilute starting solution), and the capability for 30-fold concentration. Typical processing time is 10 to 30 minutes depending on Nominal Molecular Weight Limit (NMWL). Solute polarization and subsequent fouling of the membrane are minimized by the vertical design, and a physical deadstop in the filter device prevents spinning to dryness and potential sample loss. Efficient recovery of the concentrated sample (retained species) is achieved by a convenient reverse spin step after collecting the filtrate. Amicon® Ultra-0.5 devices are supplied non-sterile and are for single use only.

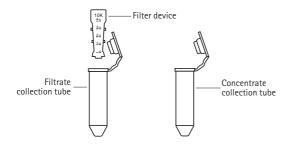
The Amicon® Ultra-0.5 product line includes 5 different cutoffs (Nominal Molecular Weight Limit, NMWL). These devices are for research use only and not for use in diagnostic procedures.

- Amicon® Ultra 3K device 3,000 NMWL
- Amicon® Ultra 10K device 10,000 NMWL
- Amicon® Ultra 30K device 30,000 NMWL
- Amicon® Ultra 50K device 50,000 NMWL
- Amicon® Ultra 100K device 100,000 NMWL

Applications

- Concentration of biological samples containing antigens, antibodies, enzymes, nucleic acids (DNA/RNA samples, either single- or double-stranded), microorganisms, column eluates, and purified samples
- Purification of macromolecular components found in tissue culture extracts and cell lysates, removal
 of primer, linkers, or molecular labels from a reaction mix, and protein removal prior to HPLC
- Desalting, buffer exchange, or diafiltration

Materials Supplied



The Amicon® Ultra-0.5 device is supplied with two microcentrifuge tubes. During operation, one tube is used to collect filtrate; the other to recover the concentrated sample.

Required Equipment

Centrifuge with fixed angle rotor that can accommodate 1.5 mL microcentrifuge tubes

CAUTION: To avoid damage to the device during centrifugation, check clearance before spinning.

Suitability

Preliminary recovery and retention studies are suggested to ensure suitability for intended use. See the "How to Quantify Recoveries" section.

Device Storage

Store at room temperature.

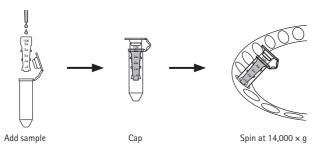
Prerinsing

The ultrafiltration membranes in Amicon® Ultra-0.5 devices contain trace amounts of glycerine. If this material interferes with analysis, pre-rinse the device with buffer or Milli-Q® water. If interference continues, rinse with 0.1 N NaOH followed by a second spin of buffer or Milli-Q® water.

CAUTION: Do not allow the membrane in Amicon® Ultra filter devices to dry out once wet. If you are not using the device immediately after pre-rinsing, leave fluid on the membrane until the device is used

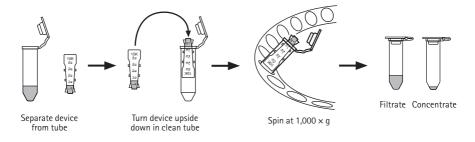
How to Use Amicon® Ultra-0.5 Centrifugal Filter Devices

- Insert the Amicon® Ultra-0.5 device into one of the provided microcentrifuge tubes.
- 2. Add up to 500 μL of sample to the Amicon® Ultra filter device and cap it.
- Place capped filter device into the centrifuge rotor, aligning the cap strap toward the center of the rotor; counterbalance with a similar device.
- 4. Spin the device at 14,000 x g for approximately 10–30 minutes depending on the NMWL of the device used. Refer to Figure 1 and Table 2 for typical spin times.



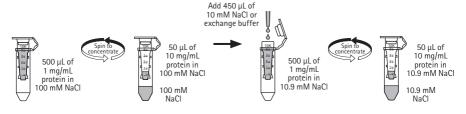
- Remove the assembled device from the centrifuge and separate the Amicon® Ultra filter device from the microcentrifuge tube.
- 6. To recover the concentrated solute, place the Amicon® Ultra filter device upside down in a clean microcentrifuge tube. Place in centrifuge, aligning open cap towards the center of the rotor; counterbalance with a similar device. Spin for 2 minutes at 1,000 × g to transfer the concentrated sample from the device to the tube. The ultrafiltrate can be stored in the centrifuge tube.

NOTE: For optimal recovery, perform the reverse spin immediately.



Desalting or Diafiltration

Desalting, buffer exchange, or diafiltration are important methods for removing salts or solvents in solutions containing biomolecules. The removal of salts or the exchange of buffers can be accomplished in the Amicon® Ultra-0.5 device by concentrating the sample, discarding the filtrate, then reconstituting the concentrate to the original sample volume with any desired solvent. The process of "washing out" can be repeated until the concentration of the contaminating microsolute has been sufficiently reduced. See example below.



Performance - DNA Concentration

The Amicon® Ultra-0.5 30K device provides the best balance between recovery and spin time for double-stranded DNA for base pairs ranging from 137 to 1159. To achieve maximum PCR product recovery and primer removal with primers greater than 20 bases, one or two additional spins with Tris-EDTA (TE) buffer are recommended.

Table 1. Typical recovery of nucleotides from Amicon® Ultra-0.5 devices

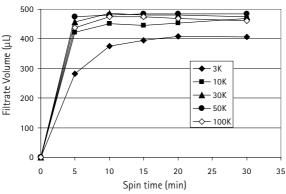
PCR Product (base pairs)	PCR Primer (bases)	PCR Recovery (%)	PCR Primer Removal (%)	TE Washes (number)
137	10	≥ 95	≥ 90	0
	20	≥ 90	≥ 85	1
	48	≥ 90	≥ 75	2
301	10	≥ 90	≥ 90	0
	20	≥ 85	≥ 90	1
	48	≥ 90	≥ 80	2
657	10	≥ 95	≥ 90	0
	20	≥ 90	≥ 90	1
	48	≥ 95	≥ 90	2
1159	10	≥ 90	≥ 90	0
	20	≥ 90	≥ 95	1
	48	≥ 95	≥ 95	2

Performance - Protein Concentration

Flow Rate

Factors affecting flow rate include sample concentration, starting volume, chemical nature of solute, relative centrifugal force, centrifuge rotor angle, membrane type, and temperature. Figure 1 and Table 2 can be used to estimate the time required to achieve a given volume of filtrate or concentrate for a variety of protein markers. A typical spin time for a 500 μ L sample is approximately 10 to 30 minutes (depending on device nominal molecular weight limit). While most of the sample is filtered in the first 5 to 10 minutes of centrifugation, the lowest concentrate volume (15–20 μ L) is reached after spinning for 10 to 30 minutes.

Figure 1. Typical Filtrate Volume vs. Spin Time



Spin conditions: 40° fixed angle rotor, $14,000\times g$, room temperature, $500~\mu L$ starting volume. Protein markers used: Cytochrome c for 3K and 10K, BSA for 30K and 50K, and IgG for 100K, n=8.

Table 2. Typical Concentrate Volume / Concentration Factor vs. Spin Time

	3K de	evice	10K d	evice	30K d	evice	50K d	evice	100K d	levice
Spin Time (min)	Conc. Volume (µL)	Conc. Factor (x)								
5	215	2	74	7	42	12	28	18	58	9
10	114	4	42	12	23	22	20	25	19	26
15	80	6	27	18	19	27	17	30	15	33
20	62	8	20	25	17	30	15	33	13	36
30	48	10	17	30	15	32	15	36	11	41

Spin conditions: 40° fixed angle rotor, $14,000 \times g$, room temperature, $500 \mu L$ starting volume. Protein markers used: Cytochrome c for 3K and 10K, BSA for 30K and 50K, and IgG for 100K, n=12. Shaded volumes were used for the calculation of protein recovery in Table 4.

Protein Retention and Concentrate Recovery

The membranes used in Amicon® Ultra devices are characterized by a nominal molecular weight limit (NMWL); that is, their ability to retain molecules above a specified molecular weight. Solutes with molecular weights close to the NMWL may be only partially retained. Membrane retention depends on the solute's molecular size and shape. For most applications, molecular weight is a convenient parameter to use in assessing retention characteristics. Merck Millipore Ltd. (Millipore) recommends using a membrane with a NMWL at least two times smaller than the molecular weight of the protein solute that one intends to concentrate. Refer to the table below.

Table 3. Typical Retention of Protein Markers

Marker/Concentration	Molecular Weight	Device NMWL	% Retention	Spin Time (min)
α-Chymotrypsinogen (1 mg/mL)	25,000	3K	>95	30
Cytochrome c (0.25 mg/mL)	12,400		>95	30
Vitamin B-12 (0.2 mg/mL)	1,350		< 42	30
α-Chymotrypsinogen (1 mg/mL)	25,000	10K	>95	15
Cytochrome c (0.25 mg/mL)	12,400		>95	15
Vitamin B-12 (0.2 mg/mL)	1,350		<23	15
BSA (1 mg/mL)	67,000	30K	>95	10
Ovalbumin (1 mg/mL)	45,000		>95	10
Cytochrome c (0.25 mg/mL)	12,400		<35	10
BSA (1 mg/mL)	67,000	50K	>95	10
Ovalbumin (1 mg/mL)	45,000		~ 40	10
Cytochrome c (0.25 mg/mL)	12,400		< 20	10
Thyroglobulin (0.5 mg/mL)	677,000	100K	>95	10
lgG (1 mg/mL)	156,000		>95	10
Ovalbumin (1 mg/mL)	45,000		<30	10

Spin Conditions: 40° fixed angle rotor, 14,000 \times g, room temperature, 500 μL starting volume, n=12.

Protein Retention and Concentrate Recovery, continued

Factors that determine sample recovery include the nature of the protein solute relative to the device NMWL chosen, starting concentration, and concentration factor. Table 4 provides typical recoveries for Amicon® Ultra-0.5 devices.

Table 4. Typical Concentrate Recovery

Marker/ Concentration	Molecular Weight	Device NMWL	Spin Time (min)	Concentrate Volume (µL)	Concentration Factor (x)	Concentrate Recovery (%)
Cytochrome c (0.25 mg/mL)	12,400	3K	30	48	10	98
Cytochrome c (0.25 mg/mL)	12,400	10K	15	27	18	95
BSA (1 mg/mL)	67,000	30K	10	23	22	97
BSA (1 mg/mL)	67,000	50K	10	20	25	92
IgG (1 mg/mL)	156,000	100K	10	19	26	92

Spin Conditions: 40° fixed angle rotor, 14,000 \times g, room temperature, 500 μ L starting volume, n=12. The shaded volumes were taken from Table 2.

Maximizing Sample Recovery

Low sample recovery in the concentrate may be due to adsorptive losses, over-concentration, or passage of sample through the membrane.

- Adsorptive losses depend upon solute concentration, its hydrophobic nature, temperature and time
 of contact with filter device surfaces, sample composition, and pH. To minimize losses, remove
 concentrated samples immediately after centrifugal spin.
- If starting sample concentration is high, monitor the centrifugation process in order to avoid overconcentration of the sample. Over-concentration can lead to precipitation and potential sample loss.
- If the sample appears to be passing through the membrane, choose a lower NMWL Amicon® Ultra-0.5
 device.

How to Quantify Recoveries

Calculate total recovery, percent concentrate recovery, and percent filtrate recovery using the method below. The procedure provides a close approximation of recoveries for solutions having concentrations up to roughly 20 mg/mL.

NOTE: Appropriate assay techniques include absorption spectrophotometry, radioimmunoassay, refractive index, and conductivity.

Direct Weighing Procedure

The density of most dilute proteins is nearly equal to the density of water (i.e., 1 g/mL). Using this property, the concentrate and filtrate volumes can be quantified by weighing them and converting the units from grams to milliliters. This technique is valid only for solutions with concentrations of approximately $20 \ mg/mL$ or less.

- Separately weigh the empty filter device, filtrate collection tube, and concentrate collection tube before use.
- 2. Fill filter device with solution and reweigh.
- 3. Assemble device in filtrate collection tube and centrifuge per instructions.
- 4. Collect the concentrate by reverse spin into the pre-weighed concentrate collection tube.
- Remove the device from the concentrate collection tube and weigh the filtrate and concentrate collection tubes.
- Subtract weight of empty device/tubes to calculate weights of starting material, filtrate, and concentrate.
- 7. Assay the starting material, filtrate, and concentrate to determine solute concentration.
- 8. Calculate recoveries using the weight/volume data and the measured concentrations as follows:

% concentrate recovery = 100
$$\times \frac{W_c \times C_c}{W_o \times C_o}$$

% filtrate recovery = 100
$$\times \frac{W_f \times C_f}{W_o \times C_o}$$

% total recovery = % concentrate recovery + % filtrate recovery

W_c = total weight of concentrate before assay

W_o = weight of original starting material

 $W_f = weight of filtrate$

 C_c = concentrate concentration

 $C_o =$ original starting material concentration

 C_f = filtrate concentration

Specifications

 $\begin{tabular}{lll} Maximum initial sample volume & 500 μL \\ \hline Typical final concentrate volume & 15-20 μL \\ \end{tabular}$

Recommended relative centrifugal force $14,000 \times g$ for concentration spin

1,000 × g for reverse spin

 $\begin{tabular}{lll} \mbox{Maximum relative centrifugal force} & 15,000 \times g \\ \mbox{Active membrane area} & 1~\mbox{cm}^2 \\ \mbox{Hold-up volume} & <5~\mbox{μL} \\ \end{tabular}$

Dimensions

Filter device and tube

Length (concentration mode; device in tube): 49.9 mm (1.96 in.) Length (reverse spin; device upside down in tube): 47.4 mm (1.87 in.)

 Tube (cap closed)
 Diameter: 10.8 mm (0.43 in.)
 Length: 42.1 mm (1.66 in.)

 Filter device
 Diameter: 9.4 mm (0.37 in.)
 Length: 29.5 mm (1.16 in.)

Materials of Construction

Filter device Copolymer styrene/butadiene

Membrane Ultracel® low binding regenerated cellulose

Collection tubes Polypropylene

Chemical Compatibility

Amicon® Ultra centrifugal devices are intended for use with biological fluids and aqueous solutions. Before use, check the sample for chemical compatibility with the device.

Table 5. Chemical Compatibility of Amicon® Ultra Filter Devices

Acids	Concentration		Concentration
Acetic acid	≤ 50%*	Phosphoric acid	≤30%
Formic acid	≤ 5%*	Sulfamic acid	≤ 3%
Hydrochloric acid	≤ 1.0 M	Sulfuric acid	≤ 3%
Lactic acid	≤50%	Trichloroacetic acid (TCA)	≤10%*
Nitric acid	≤ 10%	Trifluoroacetic acid (TFA)	≤30%*

Alkalis

Ammonium hydroxide	/droxide ≤ 10% Sodium hydroxide		≤0.5 M
Alcohols			
n-Butanol	≤ 70%	Isopropanol	≤ 70%
Ethanol	≤ 70%	Methanol	≤ 60%

Detergents

Alconox® detergent	≤ 1%	≤ 1% Lubrol® PX detergent	
CHAPS detergent	≤ 0.1%	Nonidet™ P-40 surfactant	≤ 2%
Sodium deoxycholate	≤ 5%	Triton® X-100 surfactant	≤ 0.1%
Sodium dodecyl sulfate (SDS)	≤ 0.1%	Tween® 20 surfactant	≤ 0.1%
Tergazyme® detergent	≤ 1%		

Organic solvents

Organic solvenes			
Acetone	not recommended	Ethyl acetate	not recommended
Acetonitrile	≤20%	Formaldehyde	≤ 5%
Benzene	not recommended	Pyridine	not recommended
Carbon tetrachloride	not recommended	Tetrahydrofuran	not recommended
Chloroform	not recommended	Toluene	not recommended
Dimethyl sulfoxide (DMSO)	≤ 5%*		

Miscellaneous

MISCEITATICOUS			
Ammonium sulfate	Saturated	Phenol	≤ 1%
Diethyl pyrocarbonate	≤ 0.2%	Phosphate buffer (pH 8.2)	≤1 M
Dithiothreitol (DTT)	≤0.1 M	Polyethylene glycol	≤ 10%
Glycerine	≤ 70%	Sodium carbonate	≤ 20%
Guanidine HCI	≤6 M	Tris buffer (pH 8.2)	≤1 M
Imidazole	≤ 100 mM	Urea	≤8 M
Mercaptoethanol	≤0.1 M		

^{*} Contact with this chemical may cause materials to leach out of the component parts. Solvent blanks are recommended to determine whether leachables represent potential assay interferences.

Centrifugal Product Ordering Information

Amicon® Ultra Devices

Initial volume (mL)	Final concentrate volume (µL)	Product	Qty/ pk	3K	10K	30K	50K	100K
0.5	15–20	Amicon® Ultra-0.5 device	8 24 96 500	UFC500308 UFC500324 UFC500396 UFC5003BK	UFC501008 UFC501024 UFC501096 UFC5010BK	UFC503008 UFC503024 UFC503096 UFC5030BK	UFC505008 UFC505024 UFC505096 UFC5050BK	UFC510008 UFC510024 UFC510096 UFC5100BK
Amicon®	Amicon® Ultra-0.5 Collection Tubes		96	UFC50VL96				
2	15–70	Amicon® Ultra-2 pre-launch device	24	UFC200324PL	UFC201024PL	UFC203024PL	UFC205024PL	UFC210024PL
4	50-100	Amicon® Ultra-4 device	8 24 96	UFC800308 UFC800324 UFC800396	UFC801008* UFC801024* UFC801096*	UFC803008 UFC803024 UFC803096	UFC805008 UFC805024 UFC805096	UFC810008 UFC810024 UFC810096
15	150-300	Amicon® Ultra-15 device	8 24 96	UFC900308 UFC900324 UFC900396	UFC901008* UFC901024* UFC901096*	UFC903008 UFC903024 UFC903096	UFC905008 UFC905024 UFC905096	UFC910008 UFC910024 UFC910096

^{*} Amicon® Ultra-4 and -15 10K devices are for in vitro diagnostic use. All other devices are for research use only.

Technical Assistance

For more information, contact the office nearest you. In the U.S., call 1-800-MILLIPORE (1-800-645-5476). Outside the U.S., go to our web site at www.millipore.com/offices for up-to-date worldwide contact information. You can also visit the tech service page on our web site at www.millipore.com/techservice.

Standard Warranty

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