# **Millipore**

User Guide

# Amicon<sup>®</sup> Ultra-2 Centrifugal Filter Devices

for volumes up to 2 mL

For research use only; not for use in diagnostic procedures.



### Introduction

Amicon<sup>®</sup> Ultra-2 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 50-fold concentration. Typical processing time is 10 to 60 minutes depending on Molecular Weight Cut Off (MWCO). 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. The device can be spun in a swinging bucket or fixed angle rotor. Amicon<sup>®</sup> Ultra-2 devices are supplied non-sterile and are for single use only.

The Amicon<sup>®</sup> Ultra-2 product line includes 5 different cutoffs (Molecular Weight Cut Off, MWCO). These devices are for research use only and not for use in diagnostic procedures.

- Amicon<sup>®</sup> Ultra 3K device 3,000 MWCO
- Amicon<sup>®</sup> Ultra 10K device 10,000 MWCO
- Amicon<sup>®</sup> Ultra 30K device 30,000 MWCO
- Amicon<sup>®</sup> Ultra 50K device 50,000 MWCO
- Amicon<sup>®</sup> Ultra 100K device 100,000 MWCO

### **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<sup>®</sup> Ultra-2 device is supplied with two tubes. During operation, one tube is used to collect filtrate; the other to cap the device during concentration and subsequently to recover the concentrated sample.

Amicon® Ultra-2 Centrifugal Filter Devices

# **Required Equipment**

Centrifuge with swinging bucket or fixed angle rotor with wells/carriers that can accommodate  $17 \text{ mm} \times 100 \text{ mm}$  tubes (same well/carrier size as for Amicon<sup>®</sup> Ultra-4 devices and the former Centricon<sup>®</sup> device).

**CAUTION:** To avoid damage to the device during centrifugation, make sure it is properly assembled and seated at the bottom of the rotor. The rim of the concentrate collection tube should be inside the rotor well. 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<sup>®</sup> Ultra-2 devices contain trace amounts of glycerine. If this material interferes with analysis, pre-rinse the device with buffer or Milli-Q<sup>®</sup> water. If interference continues, rinse with 0.1 N NaOH followed by a second spin of buffer or Milli-Q<sup>®</sup> water.

**CAUTION:** Do not allow the membrane in Amicon<sup>®</sup> 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<sup>®</sup> Ultra-2 Centrifugal Filter Devices

- 1. Insert the Amicon<sup>®</sup> Ultra-2 device into the filtrate collection tube, making sure that the device is fully seated in the tube.
- 2. Add up to 2 mL of sample to the device and cover with concentrate collection tube. Push the tube firmly onto the device.

**WARNING:** Failure to fully seat the device in the filtrate collection tube and push the concentrate collection tube firmly onto the device may result in the device breaking during centrifugation. See figure below.



Add sample Make sure both tubes are **fully seated** onto device

3. Place filter device into the centrifuge rotor with one membrane panel facing the center of the rotor (one panel facing up and the other panel facing down). Make sure the device is seated on the bottom of the rotor well and that the rim of the concentrate collection tube is completely inside the well. See figures below. Counterbalance with a similar device.



Orient device correctly in rotor



- 4. Spin for approximately 10–60 minutes depending on the MWCO of the device used:
  - 4,000 × g maximum when using a swinging bucket rotor

7,500 × g maximum when using a fixed angle rotor

NOTE: When spinning viscous solutions such as undiluted serum or plasma, do not exceed 5,400 x g.

Refer to Figures 1 and 2 and Tables 2 and 3 for typical spin times.

- 5. Remove the assembled device from the centrifuge and separate the Amicon<sup>®</sup> Ultra filter device from the filtrate collection tube.
- 6. To recover the concentrated solute, invert the Amicon<sup>®</sup> Ultra filter device and concentrate collection tube. Place in centrifuge and counterbalance with a similar device. Spin for 2 minutes at 1,000 × g to transfer the concentrated sample from the device to the 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<sup>®</sup> Ultra-2 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<sup>®</sup> Ultra-2 30K device provides the best balance between PCR recovery and PCR primer removal for double-stranded DNA for base pairs ranging from 137 to 1159.

#### Table 1. Typical Recovery of Nucleotides from the Amicon® Ultra-2 30K Device

		Swingin 4,000 >	g Bucket R < g for 40	otor min	35° Fixed Angle Rotor 7,500 × g, for 15 min			
PCR Product (base pairs)	PCR Primer (bases)	PCR Recovery (%)	PCR Primer Removal (%)	Final Voume (µL)	PCR Recovery (%)	PCR Primer Removal (%)	Final Volume (µL)	
	10	83	92	44	78	93	27	
137	20	87	80	43	75	86	22	
	48	86	61	41	78	67	25	
	10	96	98	35	95	98	26	
1159	20	97	93	39	93	93	26	
	48	97	82	37	95	82	27	

100 µL PCR diluted to 2,000 µL starting volume, n=6

### **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. Figures 1 and 2 and Tables 2 and 3 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 2 mL sample in a fixed angle rotor is approximately 10 to 60 minutes (depending on device nominal molecular weight limit). While most of the sample is filtered in the first 10 to 20 minutes of centrifugation, the lowest concentrate volume ( $30-70 \mu$ L) is reached after spinning for 10 to 60 minutes.



#### Figure 1. Typical Filtrate Volume vs. Spin Time for Amicon® Ultra-2 Device, Swinging Bucket Rotor

Spin conditions: Swinging bucket rotor, 4,000  $\times$  g, room temperature, 2 mL starting volume. Protein markers used: Cytochrome c for 3K and 10K, BSA for 30K and 50K, and IgG for 100K, n=8.

#### Figure 2. Typical Filtrate Volume vs. Spin Time for Amicon® Ultra-2 Device, Fixed Angle Rotor



Spin conditions: 35° fixed angle rotor, 7,500  $\times$  g, room temperature, 2 mL 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, Swinging Bucket Rotor

	3K device		10K device		30K device		50K device		100K device	
Spin Time (min)	Conc. Volume (µL)	Conc. Factor (x)								
5					281	7	91	22	1070	2
10	880	2	190	11	71	27	47	42	523	4
15			96	21	52	38	44	47	167	12
20	317	7	65	31	43	46	38	52	65	31
30	147	30	48	42	39	51	38	53	37	53
40	102	20	44	45						
60	55	32								

Spin conditions: Swinging bucket rotor, 4,000  $\times$  g, room temperature, 2 mL starting volume. Protein markers used: Cytochrome c for 3K and 10K, BSA for 30K and 50K, and IgG for 100K, n=8. Shaded volumes were used for the calculation of protein recovery in Table 5.

#### Table 3. Typical Concentrate Volume/Concentration Factor v. Spin Time, Fixed Angle Rotor

	3K device		10K device		30K device		50K device		100K device	
Spin Time (min)	Conc. Volume (µL)	Conc. Factor (x)								
5					137	15	80	25	879	2
10	731	3	101	21	51	39	30	71	203	10
15			60	33	37	57	22	90	61	34
20	215	10	39	51	24	85	21	99	32	63
30	106	19	25	80	20	101	18	89	17	115
40	70	29	23	87						
60	45	45								

Spin conditions:  $35^{\circ}$  fixed angle rotor, 7,500 × g, room temperature, 2 mL starting volume. Protein markers used: Cytochrome c for 3K and 10K, BSA for 30K and 50K, and IgG for 100K, n=8. Shaded volumes were used for the calculation of protein recovery in Table 5.

### **Protein Retention and Concentrate Recovery**

The membranes used in Amicon<sup>®</sup> Ultra devices are characterized by a molecular weight cut off (MWCO); that is, their ability to retain molecules above a specified molecular weight. Solutes with molecular weights close to the MWCO 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. We recommend using a membrane with a MWCO at least two times smaller than the molecular weight of the protein solute that one intends to concentrate. Refer to Table 4.

#### **Table 4. Typical Retention of Protein Markers**

Marker/Concentration	Molecular Weight	Device MWCO	% Retention Swinging Bucket	Spin Time (min)	% Retention Fixed Angle	Spin Time (min)
a-Chymotrypsinogen (1 mg/mL)	25,000	3K	99	60	99	60
Cytochrome c (0.25 mg/mL)	12,400		100		100	
Vitamin B-12 (0.2 mg/mL)	1,350		6		8	
a-Chymotrypsinogen (1 mg/mL)	25,000	10K	99	30	99	20
Cytochrome c (0.25 mg/mL)	12,400		100		100	
Vitamin B-12 (0.2 mg/mL)	1,350		10		9	
BSA (1 mg/mL)	67,000	30K	100	20	100	15
Ovalbumin (1 mg/mL)	45,000		97		97	
Cytochrome c (0.25 mg/mL)	12,400		16		15	
BSA (1 mg/mL)	67,000	50K	97	15	100	10
Ovalbumin (1 mg/mL)	45,000		50		60	
Cytochrome c (0.25 mg/mL)	12,400		9		17	
Thyroglobulin (0.5 mg/mL)	677,000	100K	94	30	94	20
IgG (1 mg/mL)	156,000		95		95	
Ovalbumin (1 mg/mL)	45,000		12		13	

Spin Conditions: Swinging bucket rotor, 4,000  $\times$  g, or 35° fixed angle rotor, 7,500  $\times$  g, 2 mL starting volume, room temperature, n=12.

Factors that determine sample recovery include the nature of the protein solute relative to the device MWCO chosen, starting concentration, and concentration factor. Table 5 provides typical recoveries for Amicon<sup>®</sup> Ultra-2 devices.

#### Table 5. Typical Concentrate Recovery

		Spin Time (min)		Concentrate Volume (uL)		Concentration Factor (x)		Concentrate Recovery (%)	
Marker/ Concentration	Device MWCO	Swinging Bucket	Fixed Angle	Swinging Bucket	Fixed Angle	Swinging Bucket	Fixed Angle	Swinging Bucket	Fixed Angle
Cytochrome c (0.25 mg/mL)	3K	60	60	55	45	32	45	97	96
Cytochrome c (0.25 mg/mL)	10K	30	20	48	39	42	51	98	98
BSA (1 mg/mL)	30K	20	15	43	37	46	57	94	94
BSA (1 mg/mL)	50K	15	10	44	30	47	71	93	87
IgG (1 mg/mL)	100K	30	20	37	32	53	63	88	90

Spin Conditions: Swinging bucket rotor,  $4,000 \times g$ , or  $35^{\circ}$  fixed angle rotor,  $7,500 \times g$ , 2 mL starting volume, room temperature, n=8.

### **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 MWCO Amicon<sup>®</sup> Ultra-2 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.

- 1. 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.
- 5. Remove the device from the concentrate collection tube and weigh the filtrate and concentrate collection tubes.
- 6. 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 × 
$$\frac{W_{c} \times C_{c}}{W_{o} \times C_{c}}$$

% 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<sub>f</sub> = weight of filtrate
- C<sub>c</sub> = concentrate concentration
- $C_{o}$  = original starting material concentration
- $C_f$  = filtrate concentration

# **Specifications**

Maximum initial sample volume			2.0 mL				
Typical final concentrate volume			30-70 µL depending on MWCO				
Maximum relative centrifugal force							
Swinging bucket rotor		4,0 1,0	000 × g for c 000 × g for r	concentra ecovery	ation spii spin	n,	
Fixed angle rotor		7,5 1,0	500 × g for c 000 × g for r	concentra ecovery	ation spii spin	n,	
			<b>NOTE:</b> When spinning viscous solutions such as undiluted serum or plasma, do not exceed 5,400 x g.				
Active membrane area		1 0	cm <sup>2</sup>				
Hold-up volume			< 5 µL				
Dimensions							
Filter device and tube							
Length (concentra	tion mod	e; device i	in filtrate tub	e):		119.7 mm (4.71 in.)	
Length (recovery s	spin; devi	ce upside	down in con	centrate	tube):	95.3 mm (3.75 in.)	
Filter device Di	ameter:	15.9 mm	(0.63 in.)	L	ength:	70.7 mm (2.78 in.)	
Filtrate tube Di	ameter:	13.8 mm	(0.54 in.)	L	ength:	52.9 mm (2.08 in.)	
Concentrate tube Di	ameter:	13.7 mm	(0.54 in.)	L	ength:	34.5 mm (1.36 in.)	
Materials of Construction	n						
Filter device	Co	Copolymer styrene/butadiene					
Membrane	Ult	Ultracel <sup>®</sup> low-binding regenerated cellulose					
Collection tubes		Po	lypropylene				

# **Chemical Compatibility**

Amicon<sup>®</sup> 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 6. 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	≤10%	Sodium hydroxide	≤0.5 M
Alcohols			
n-Butanol	≤70%	Isopropanol	≤70%
Ethanol	≤70%	Methanol	≤60%
Detergents			-
Alconox <sup>®</sup> detergent	≤1%	Sodium dodecyl sulfate (SDS)	≤0.1%
CHAPS detergent	≤0.1%	Tergazyme <sup>®</sup> detergent	≤1%
Lubrol <sup>®</sup> PX detergent	≤0.1%	Triton <sup>®</sup> X-100 surfactant	≤0.1%
Nonidet <sup>™</sup> P-40 surfactant	≤2%	Tween <sup>®</sup> 20 surfactant	≤0.1%
Sodium deoxycholate	≤5%		
Organic solvents			
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			
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 HCl	≤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.

# **Product Ordering Information**

This section lists the catalogue numbers for Amicon<sup>®</sup> Ultra Ultrafiltration Devices. See the Technical Assistance section for contact information. You can purchase these products on-line at <u>www.sigmaaldrich.com/products</u>.

мwсо	Qty/ pk	Amicon® Ultra-0.5 device	Amicon® Ultra-2 device	Amicon® Ultra-4 device	Amicon® Ultra-15 device				
	8	UFC500308		UFC800308	UFC900308				
214	24	UFC500324	UFC200324	UFC800324	UFC900324				
3K	96	UFC500396		UFC800396	UFC900396				
	500	UFC5003BK							
	8	UFC501008		UFC801008	UFC901008				
101/	24	UFC501024	UFC201024	UFC801024	UFC901024				
IUK	96	UFC501096		UFC801096	UFC901096				
	500	UFC5010BK							
101/	8			UFC801008D	UFC901008D				
	24			UFC801024D	UFC901024D				
IVD	96			UFC801096D	UFC901096D				
	8	UFC503008		UFC803008	UFC903008				
304	24	UFC503024	UFC203024	UFC803024	UFC903024				
201	96	UFC503096		UFC803096	UFC903096				
	500	UFC5030BK							
	8	UFC505008		UFC805008	UFC905008				
FOR	24	UFC505024	UFC205024	UFC805024	UFC905024				
JUK	96	UFC505096		UFC805096	UFC905096				
	500	UFC5050BK							
	8	UFC510008		UFC810008	UFC910008				
1001	24	UFC510024	UFC210024	UFC810024	UFC910024				
TOOK	96	UFC510096		UFC810096	UFC910096				
	500	UFC5100BK							
* Amicon®	* Amicon <sup>®</sup> Ultra-4 and -15 10K devices are for in vitro diagnostic (IVD) use. All other devices are for research use only.								

### Notice

We provide information and advice to our customers on application technologies and regulatory matters to the best of our knowledge and ability, but without obligation or liability. Existing laws and regulations are to be observed in all cases by our customers. This also applies in respect to any rights of third parties. Our information and advice do not relieve our customers of their own responsibility for checking the suitability of our products for the envisaged purpose.

# **Contact Information**

For the location of the office nearest you, go to <u>www.sigmaaldrich.com/offices</u>.

### **Technical Assistance**

Visit the tech service page on our web site at <u>www.sigmaaldrich.com/techservice</u>.

### **Standard Warranty**

The applicable warranty for the products listed in this publication may be found at <u>www.sigmaaldrich.com/terms</u> ("Conditions of Sale").

The vibrant M, Millipore, Amicon, Milli-Q, and Ultracel are trademarks of Merck KGaA, Darmstadt, Germany, or its affiliates. All other trademarks are the property of their respective owners. Detailed information on trademarks is available via publicly accessible resources.

 ${\small ©}$  2019 Merck KGaA, Darmstadt, Germany and/or its affiliates. All rights reserved.

PR05484, Rev. 02/19



The life science business of Merck KGaA, Darmstadt, Germany operates as MilliporeSigma in the U.S. and Canada.