

# FREE PRODUCT OFFER!

Buy 2 Packs of Nanosep, Microsep or Macrosep Centrifugal Devices, Get 1 of the same size FREE!

## **Applications**

Centrifugal devices can replace traditional separation techniques, such as column chromatography, preparative electrophoresis, alcohol or salt precipitation, dialysis, and gradient centrifugation, when performing the following:

- Protein or nucleic acid concentration
- Desalting
- Buffer exchange
- Deproteination of biological samples
- Fractionation of protein mixtures
- Separation of primers from PCR products
- Separation of labeled nucleic acids or proteins from unincorporated nucleotides
- Virus concentration or removal
- Clarification of cell lysates and tissue homogenates
- Extracting, isolating and purifying nucleic acids

## **Benefits**

- Accelerate sample processing Concentrate and purify samples with starting volumes of < 50 µL to 60 mL.</li>
- Maximize sample recovery Obtain high flow rates and low non-specific protein and nucleic acid binding.
- Add versatility Available in various membrane types including low-binding
- Bio-Inert® (modified nylon), Supor® (polyethersulfone), and wwPTFE membranes, as well as Omega™ (modified polyethersulfone) ultrafiltration membrane in a variety of MWCOs.
- Prevent solution bypass Membrane seals stop solution leakage, minimizing sample loss.
- Easy visual identification Devices are color-coded for a wide variety of membranes, ranging from 1 kD to 0.45 μm.







#### **Nanosep Centrifugal Device**

Concentration selection guide for Nanosep Centrifugal Devices

Concentration Factor (Fold)	Starting Sample Volume (µL)	Volume Added to Collection Tube (µL)	Final Rententate Volume (µL)
2	200	572	100
3	200	530	67
4	200	508	50
5	200	496	40
6	200	487	33
10	200	470	20
20	200	470	10

## **Microsep Advance Centrifugal Device**

Concentration selection guide for Microsep Advance Centrifugal Devices

Concentration Factor (Fold)	Starting Sample Volume (mL)	Volume Added to Collection Tube (mL)	Final Rententate Volume (mL)
2	3.00	6.69	1.50
3	3.00	5.76	1.00
4	3.00	5.29	0.75
5	3.00	5.02	0.60
6	3.00	4.83	0.50
10	3.00	4.46	0.30
20	3.00	4.18	0.15
25	3.00	4.12	0.12

The above table shows what buffer volume should be added to the collection tube below the membrane to achieve desired concentration factors for the listed starting sample volume.

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For instance, for the concentration of 200 µL starting material by ten-fold (see highlight in table), the buffer volume to be added to the collection tube would be 470 µL, leaving 20 µL of concentrated material in the retentate.

For a complete Concentration Selection Guide visit: www.pall.com

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## **Ordering Information**

### **Qualifying Nanosep Centrifugal Devices**

200

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Part Number	Size	Pkg
PAL-OD003C33	3K, gray	24/pkg
PAL-OD010C33	10K, blue	24/pkg
PAL-OD030C33	30K, red	24/pkg
PAL-OD100C33	100K, clear	24/pkg
PAL-OD300C33	300K, orange	24/pkg

## **Qualifying Macrosep Advance Centrifugal Devices**

Part Number	Size	Pkg
PAL-MAP001C37	1K, yellow	24/pkg
PAL-MAP003C37	3K, gray	24/pkg
PAL-MAP010C37	10K, blue	24/pkg
PAL-MAP030C37	30K, red	24/pkg
PAL-MAP100C37	100K, clear	24/pkg

## **Qualifying NAB Nanosep Centrifugal Devices**

Part Number S	Size	Pkg
PAL-ODNABC33	NAB Nanosep Device	24/pkg

Promotion available until March 31, 2022.

Promotion assumes products are purchased at local list pricing. Contact your local ESBE representative with any questions.



#### **Qualifying Microsep Advance Centrifugal Devices**

Part Number	Size	Pkg
PAL-MCP001C41	1K, yellow	24/pkg
PAL-MCP003C41	3K, gray	24/pkg
PAL-MCP010C41	10K, blue	24/pkg
PAL-MCP030C41	30K, red	24/pkg
PAL-MCP100C41	100K, clear	24/pkg

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