## <u>FLEXTUBES</u>

#### Dialysis

IBI FlexTubes are innovative dialysis tubes providing cutting edge dialysis and buffer exchange technology. This revolutionary design allows for easy loading and unloading with a standard pipette tip. IBI FlexTubes are the only dialysis tool available today that can be used for high throughput applications, thus accommodating a large number of samples in a single buffer tank. IBI FlexTubes provide rapid, high performance extraction of macro-molecules without contamination, with yields up to 98%! They are ideal for protein purification, automated protein sequencing, peptide mapping, and amino acid analysis. IBI FlexTubes are available in numerous sample sizes and membrane cutoffs. From MINI to MEGA, from 1K MWCO to 50K MWCO choose the tube that's right for your application. All IBI FlexTubes can be used for dialysis applications.

#### Gel Extraction - DNA, RNA, Large DNA

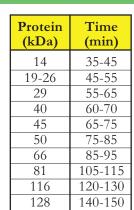
IBI FlexTubes are also ideal for gel extraction of molecules such as small nucleic acid fragments (DNA Oligonucleotides down to 15nt) and RNA. They are also ideal for extraction of large protein nucleic acid complexes and large size DNA fragments (>100kb) from agarose or polyacrylamide gels. DNA recovery from either polyacrylamide or agarose gels is typically greater than 95%! IBI FlexTubes easily fit inside any electrophoresis tank assembly for extraction applications. When used for electro-elution/extractions the IBI FlexTubes are an excellent high throughput device. Typcial extraction times for a 50kDa molecule in a MIDI FlexTubes are a single-use product.

#### Protein Extractions from Polyacrylamide Gels Compatible with MALDI-MS

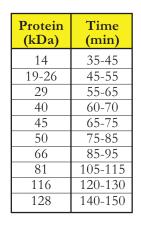
SDS-PAGE is a widely used technique for the separation and estimation of the molecular weight of individual proteins. Accuracy of molecular weight determination and characterization is limited by low recovery of proteins. Recently, MALDI-TOF-MS has found widespread use for determination of molecular mass of intact proteins isolated from gels. Isolation of proteins from a gel using IBI FlexTube electro-elution systems deliver up to 80% recovery yields! Now the combination of SDS-PAGE, IBI FlexTubes, and SDS removal buffer can give you a more accurate determination of a protein's molecular mass than ever before. Even difficult proteins; such as, integral membrane proteins (hydrophobic) and high molecular mass proteins can effectively be separated. This unique method provides a powerful means for characterizing endogenous proteins of a wide molecular weight range, separated by SDS-PAGE. The combination of the three methods provide significantly improved protein yield and SDS free samples. The result is a MALDI-MS analysis with greater sensitivity. MINI, MIDI, and MAXI IBI FlexTubes can all be used in these applications, select the one that fits your sample.

#### Oligo PAGE Electro-Elution and De-Salting

Oligo PAGE purification is made easy with IBI FlexTubes! This method is a vast improvement over the Diffusion Method, which is a lengthy over-night process that typically produces low yields. The IBI Flextubes will allow the extraction of the protein from the polyacrylamide gel, followed by de-salting of the sample by dialysis using the same FlexTube. IBI FlexTubes are ideal for high throughput applications, and use an easy to follow protocol.









**PURIFICATION** 

PRODUCTS

## FLEXTUBES MEGA

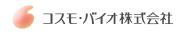
IBI MEGA FlexTubes are typically used for dialysis or buffer exchange with volumes of 3-20ml. IBI FlexTubes allow rapid, secure, simple loading and recovery, with very high performance as the most convenient user friendly dialysis system on the market today! IBI MEGA FlexTubes can also be used for applications such as; preparation of protein samples for MALDI-MS, sample concentration, tissue culture extraction purification, removal of salts, surfactants, solvents, and detergents, as well as peptide dialysis and virus-particle purification. The performance of IBI MEGA FlexTubes are regularly monitored and tested by simple dialysis and assays commonly used for proteins and/or nucleic acids.

#### **Key Features**

MWCO:	3.5K, 6-8K, or 12-14K	NEW
TUBE VOLUME:	10ml, 15ml, or 20ml	NET
DIALYSIS VOLUME:	3-20ml	
Membrane:	Ultra-clean, sulfur, and heavy metal free. EDTA treated.	<i>,</i> – ,

### **O**RDERING **I**NFORMATION

CATALOG#	DESCRIPTION	VOLUME	QUANTITY
IB48007	MEGA FLEXTUBE FLOAT - 1 PLACE		
IB48300	MEGA 10ML FLEXTUBE KIT, 1K MWCO, FLOAT	10мг	2 PACK
IB48301	MEGA 15ml FlexTube Kit, 1K MWCO, Float	15ML	2 PACK
IB48302	MEGA 20ML FLEXTUBE KIT, 1K MWCO, FLOAT	20ml	2 PACK
IB48310	MEGA 10ml FlexTube Kit, 1K MWCO	10мг	5 PACK
IB48311	MEGA 10ML FLEXTUBE KIT, 1K MWCO	10мг	15 <b>Р</b> АСК
IB48320	MEGA 15ml FlexTube Kit, 1K MWCO	15м∟	5 PACK
IB48321	MEGA 15ml FlexTube Kit, 1K MWCO	15мг	15 <b>Р</b> АСК
IB48330	MEGA 20ML FLEXTUBE KIT, 1K MWCO	20ml	5 Pack
IB48331	MEGA 20ML FLEXTUBE KIT, 1K MWCO	20мг	15 PACK
IB48340	MEGA 10ML FLEXTUBE KIT, 3.5K MWCO, FLOAT	10мг	2 PACK
IB48341	MEGA 15ML FLEXTUBE KIT, 3.5K MWCO, FLOAT	15м∟	2 PACK
IB48342	MEGA 20ML FLEXTUBE KIT, 3.5K MWCO, FLOAT	20мг	2 PACK
IB48350	MEGA 10ML FLEXTUBE KIT, 3.5K MWCO	10мг	5 Pack
IB48351	MEGA 10ML FLEXTUBE KIT, 3.5K MWCO	10мг	15 <b>Р</b> АСК
IB48360	MEGA 15ML FLEXTUBE KIT, 3.5K MWCO	15мг	5 Pack
IB48361	MEGA 15ML FLEXTUBE KIT, 3.5K MWCO	15мг	15 Pack
IB48370	MEGA 20ML FLEXTUBE KIT, 3.5K MWCO	20ml	5 Pack
IB48371	MEGA 20ML FLEXTUBE KIT, 3.5K MWCO	20мг	15 Pack
IB48380	MEGA 10ML FLEXTUBE KIT, 6K MWCO, FLOAT	10ml	2 PACK
IB48381	MEGA 15ml FlexTube Kit, 6K MWCO, Float	15м∟	2 PACK
IB48382	MEGA 20ML FLEXTUBE KIT, 6K MWCO, FLOAT	20мг	2 PACK
IB48390	MEGA 10ML FLEXTUBE KIT, 6K MWCO	10мг	5 Pack
IB48391	MEGA 10ML FLEXTUBE KIT, 6K MWCO	10ml	15 Pack
IB48400	MEGA 15ML FLEXTUBE KIT, 6K MWCO	15ML	5 PACK
IB48401	MEGA 15ML FLEXTUBE KIT, 6K MWCO	15мL	15 Pack
IB48410	MEGA 20ML FLEXTUBE KIT, 6K MWCO	20ml	5 PACK
IB48411	MEGA 20ML FLEXTUBE KIT, 6K MWCO	20мг	15 Pack
IB48420	MEGA 10ML FLEXTUBE KIT, 12K MWCO, FLOAT	10ml	2 PACK
IB48421	MEGA 15ML FLEXTUBE KIT, 12K MWCO, FLOAT	15мL	2 PACK
IB48422	MEGA 20ML FLEXTUBE KIT, 12K MWCO, FLOAT	20ml	2 PACK
IB48430	MEGA 10ML FLEXTUBE KIT, 12K MWCO	10ML	5 Pack
IB48431	MEGA 10ML FLEXTUBE KIT, 12K MWCO	10ml	15 PACK
IB48440	MEGA 15ML FLEXTUBE KIT, 12K MWCO	15ML	5 PACK
IB48441	MEGA 15ML FLEXTUBE KIT, 12K MWCO	15ML	15 PACK
IB48450	MEGA 20ML FLEXTUBE KIT, 12K MWCO	20ML	5 PACK
IB48451	MEGA 20ML FLEXTUBE KIT, 12K MWCO	20мг	15 Pack



# FLEXTUBE HANDBOOK

## **MEGA**



**IBI FLEXTUBES ARE MANUFACTURED BY GENE-BIO-APPLICATION LTD. - ISRAEL** 

**IBI/GEBA FLEXTUBES ARE COVERED BY PATENT APPLICATION WO0190731** 



## **IBI SCIENTIFIC**

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## A. PACKAGING AND HANDLING

- Wearing gloves is highly recommended when handling the kit contents.
- IBI FlexTubes are autoclaved and bacterial free.
- IBI FlexTube membranes are ultra-clean, sulfur and heavy metal free, and EDTA treated.

## **B.** APPLICATIONS

- Dialysis or buffer exchange with volumes between 3 and 20ml.
- Preparation of protein samples for MALDI-MS,
- Sample concentration.
- Large-scale protein dialysis, such as antibodies and recombinant protein purification.
- Removal of contaminating micro-molecules.
- Tissue culture extraction purification.
- Removal of salts, surfactants, solvents, and detergents.
- Complex formation studies (protein-protein, protein-DNA, and protein-RNA).
- pH and buffer adjustment of sample solutions, protein extraction or cell extraction.
- High throughput dialysis.
- Peptide dialysis, as small as 10 amino acids.
- Virus-particles purification.

## C. KIT CONTENTS

- IBI FlexTubes
- Supporting tray (for electro elution protocol)
- Floating rack (for dialysis protocol)
- Information and Protocol Manual

2 / 10 / 30 / 50 / 100 pieces 1ea. (select kits) 1ea. (select kits) 1ea.

### **D. STORAGE CONDITIONS**

IBI FlexTube kits should be stored in a dry place at room temperature (15-25°C). Under these conditions, IBI FlexTube kits can be stored for up to 12 months without any deterioration in performance or quality. For longer storage times, it is recommended that IBI FlexTube kits be stored in a cool dry place, like a refrigerator.

### **E. PRODUCT LIMITATIONS**

IBI FlexTube kits are developed, designed, and sold for research purposes only. They are not to be used for human diagnostic purposes or drug production, nor for producing any substance intended to be administered to humans unless expressly cleared for that purpose by the Food and Drug Administration (USA) or the appropriate regulatory authorities in the country of use. All due care and attention should be exercised in the handling of materials described in this text.

## F. QUALITY CONTROL

The performance of IBI FlexTube kits are regularly monitored. IBI FlexTube kits are tested by using them for extraction of Proteins, DNA and RNA fragments of various sizes from either agarose or Polyacrylamide gels. IBI FlexTube kits are also tested for simple dialysis of salts or buffer exchange. The quality of the isolated Protein, DNA and RNA fragments, or of the sample after dialysis is checked by several assays commonly used for proteins, nucleic acids, and dialysis. Determining the recovery from a specific amount of loaded samples will test the quality of the IBI FlexTube membranes.

### G. IBI FLEXTUBES

These devices can be used for dialysis or buffer exchange at volume samples between 3 and 20ml. IBI FlexTubes allow rapid, secure, simple loading and recovery, with high performance as the most convenient, user friendly dialysis system on the market.

#### **SPECIFICATIONS**

Membrane cut-off Tube volume capacity Volume of sample for dialysis Membrane 1K, 3.5K, 6-8K, 12-14K MWCO
10, 15, or 20ml - depending on kit
3 - 20ml
Ultra-clean, sulfur and heavy metal free. EDTA treated

## H. DIALYSIS WITH MEGA FLEXTUBE KITS

**IMPORTANT:** To perform dialysis with a range of 3 - 10ml use IBI MEGA 10ml FlexTubes. To perform dialysis with a rnage of 10 - 15ml us IBI MEGA 15ml FlexTubes. To perform dialysis with a range of 15 - 20ml use IBI MEGA 20ml FlexTubes.



Figure 1: Dialysis with MEGA FlexTube

#### **PROCEDURE**

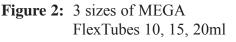
**1.** Fill the IBI MEGA FlexTube with 20ml of dH<sub>2</sub>O, incubate for at least 5 minutes, and empty the tube.

**IMPORTANT:** Check carefully that there is no dH<sub>2</sub>O leaking from the tube. Absorption of water by the dry membrane may cause a decrease in water level. Avoid positioning the MEGA FlexTubes on their cap when containing samples.

2. Load sample into IBI MEGA FlexTube. Close the tube with provided cap. For sample volume up to 10ml use MEGA 10ml FlexTubes. For sample volume between 10 and 15ml use MEGA 15ml FlexTubes. For sample volumes between 15 and 20ml use MEGA 20ml FlexTubes.

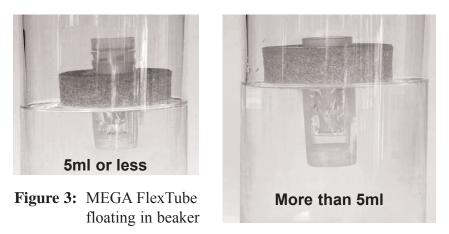
**IMPORTANT:** Sample volume should be in the range of 3 - 20ml. If smaller volumes (3-5ml) are used, load the samples close to the inner membrane.





**3.** Load IBI MEGA FlexTube into float. Position in beaker and stir. Volumes of the desired buffer are typically 100 to 1000 times that of the sample.

**IMPORTANT:** Adjust height of IBI MEGA FlexTube in the floating balance rack for upright floating during process. For less 5ml samples, place the floating rack up to the middle of the tube. For dialysis of more than 5ml, adjust the floating rack at the top of the MEGA FlexTube.



**4.** Adjust the stir bat speed. Low-molecular weight salts and buffers equilibrate within 3 hours. Equilibration time for viscous samples will be slightly longer.

**IMPORTANT:** User must determine exact equilibration times for the dialysis.

- 5. Change the dialysis buffer as necessary.
- 6. Pipette out the sample carefully from the IBI MEGA FlexTube into a clean tube.

If sample volume has increased during dialysis, let your sample evaporate on the bench top (increasing airflow across the membrane will help speed up the process). Check the sample approximately every 10 minutes or less to prevent full evaporation and/or dryness.

#### SAMPLE CONCENTRATION BY EVAPORATION

IBI MEGA FlexTubes are ideally suited for sample concentration via evaporation because of their dual membranes and large surface area. Dialysis and concentration in the same device reduces protein loss. Unlike closed-system centrifuge type devices, sample concentration can be easily monitored in the MEGA FlexTubes as well.

- 1. Place sample in the MEGA FlexTube or use an already dialyzed sample.
- 2. Let the sample evaporate on the bench top, using a fan to increase airflow across the membranes and speed up the process. Check the sample every 10 minutes of less to prevent full evaporation and dry sample condition. Concentrating by evaporation water with sample, small molecules (buffer salts, reducing agents, etc.) will also get concentrated because no diffusion occurs.

**IMPORTANT:** When evaporating water from your sample, small molecules (buffer salts, reducing agents, etc.) will also be concentrated.

## I. PROTEIN PRECIPITATION PROTOCOLS

#### TRICHLOROACETIC ACID (TCA) PRECIPITATION PROCEDURE

**1.** Add equal volume of 20% TCA to the microcentrifuge tube containing the extracted protein solution and mix properly.

For example, add 20ml of 20% TCA to a 20ml sample.

- **2.** Incubate for 1 hour at 4°C.
- 3. Spin in a microcentrifuge at 4°C for 30 minutes at 14,000RPM.
- 4. Discard supernatant carefully.
- 5. Add 14ml of cold acetone.
- **6.** Incubate at -20°C for 30 minutes and centrifuge the sample at 4°C for 30 minutes at 14,000RPM.

To increase protein precipitation yield incubate the samples over night at -20°C.

- 7. Discard supernatant and air-dry the pellet.
- 8. Resuspend the pellet using 0.1M NaOH or dH<sub>2</sub>O (use at least 0.7ml to perform resuspension).

#### MS PRECIPITATION PROCEDURE

- **1.** Add 1:10 by volume of MS buffer to the protein containing solution and mix properly. For example, add 0.3ml of MS buffer to a 3ml sample.
- 2. Incubate for 15 minutes at room temperature.
- **3.** Add 1:2 by volume of 20% TCA and mix properly. For example, add 11ml of 20% TCA to a 22ml sample.
- **4.** Incubate for 1 hour at 4°C.
- 5. Centrifuge the sample at 4°C for 30 minutes at 14,000RPM.
- 6. Carefully descent the supernatant without disturbing the pellet.

- 7. Add 14ml of cold acetone.
- **8.** Incubate at -20°C for 30 minutes and centrifuge the sample at 4°C for 30 minutes at 14,000RPM.

To increase protein precipitation yield incubate the samples over night at -20°C.

- 9. Carefully descent the supernatant without disturbing the pellet. Air-dry the pellet.
- **10.** Resuspend the pellet in a suitable buffer solution or 0.1M NaOH (use at least 0.7ml to perform resuspension).

## J. DNA OR RNA PRECIPITATION

#### **PROCEDURE**

**1.** Add 0.1 by volume of 3M KAc pH-5.2 and 0.7-1 by volume of isopropanol to the solution. Mix gently by inverting the tube several times.

For example, add 2ml of 3M KAc and 2.31 - 3.3ml isopropanol to a 14ml sample.

**Note:** Addition of carrier (e.g. 80µg tRNA or 80µg glycogen) to the solution will increase the efficiency of precipitation.

- 2. Incubate at -20°C for 10 minutes.
- 3. Centrifuge the sample at 4°C for 30 minutes at 14,000RPM.
- 4. Carefully discard the supernatant without disturbing the pellet.
- 5. Wash the pellet with cooled 70% ethanol.
- 6. Air-dry the pellet for 5-20 minutes

Do not overdry the pellet (e.g., by using a vacuum evaporator), as this will make the DNA, especially if it is of high molecular weight, difficult to redissolve.

7. Redissolve the DNA or RNA in a suitable buffer.

Use a buffer with pH >8.0 for redissolving, as DNA does not dissolve readily in acidic buffers.





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