

# MegaLong™ Protocol for Isolation of >100kb Genomic DNA (Yeast DNA)

## G-Biosciences

### Abstract

MegaLong™ isolates high molecular weight (>100kb) genomic DNA from a variety of samples, including animal tissues, cultured cells, whole blood, bacterial and yeast.

The protocol described here is for yeast samples. Please [refer to the appropriate protocol](#), depending on your application.

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## Guidelines

### INTRODUCTION

MegaLong™ isolates high molecular weight (>100kb) genomic DNA from a variety of samples, including animal tissues, cultured cells, whole blood, bacterial and yeast. MegaLong™ uses Genomic Tube-O-DIALYZER™, a unique micro dialysis device with a 0.45µm membrane, which minimizes sample manipulation, one of the main reasons for DNA breakage. MegaLong™ isolates nuclei under mild extraction conditions and releases genomic DNA by digestion of nuclear proteins with a highly active LongLife™ Proteinase K. The digestion is performed in the Tube-O-DIALYZER™ and after digestion the Tube-ODIALYZER™ is inverted to dialyze away digested protein and other impurities leaving behind highly pure and fully hydrated genomic DNA.

The fragile, high molecular weight genomic DNA can be stored in the Tube-O-DIALYZER™ to further minimize mechanical manipulation of the DNA. The DNA is suitable for Southern blot analysis, recovery of Lambda shuttle vectors from transgenic animals, PCR, analysis by pulsed-field electrophoresis or any application where genomic DNA is required.

### APPLICATIONS

MegaLong™ kit can be used for the isolation of genomic DNA from animal tissues, cultured cells, whole blood, bacterial and yeast. For samples unsuitable for the isolation of high molecular weight DNA with MegaLong™, G-Biosciences recommends using the OmniPrep™ Genomic DNA isolation kit (Cat. # 786-136).

The kit is supplied as a Micro or Large packs to process either 25 or 50 1-25mg samples.

## ITEM(S) SUPPLIED

Description	Cat. # 786-146	Cat. # 786-147
Nuclei Isolation Buffer	2 x 30ml	4 x 30ml
Suspension Buffer	1 x 10ml	2 x 10ml
Digestion Buffer	1 x 2ml	2 x 2ml
LongLife™ Proteinase K [5mg/ml]	2 x 0.5ml	4 x 0.5ml
Genomic Tube-O-Dialyzer™	25	50
Floats (Medi)	6	6
Caps (Medi)	25	50
Forceps	1	1

## STORAGE CONDITION

The kit is shipped at ambient temperature. Upon arrival, store LongLife™ Proteinase K at -20°C and remaining components at 4°C. LongLife™ Proteinase K solution is stable for 1 year, if stored properly.

## ADDITIONAL MATERIALS REQUIRED

- Microfuge tubes & pestles (Cat. # 786-138P)
- TE buffer

## Materials

MegaLong™ [786-146](#) by [G-Biosciences](#)

Pestles & Tubes [786-138P](#) by [G-Biosciences](#)

## Protocol

### Sample Preparation

#### Step 1.

On ice, prepare spheroplasts from a 1.5ml overnight culture.

### Tube-O-DIALYZER™ Preparation

#### Step 2.

Add 70µl Suspension Buffer to the nuclei and gently rock or tap the tube to dislodge the nuclei.

### Tube-O-DIALYZER™ Preparation

#### Step 3.

Vortex the LongLife™ Proteinase K and add 10µl to the nuclei.

### Tube-O-DIALYZER™ Preparation

#### Step 4.

Add 70µl Digestion Buffer and mix with gentle rocking.

### Tube-O-DIALYZER™ Preparation

#### Step 5.

Incubate at 55°C for 2-4 hours with periodic rocking. Do not vortex.



DURATION

00:30:00



NOTES

**Colin Heath** 23 Jun 2016

NOTE: For periodic rocking, gently invert the tube 2-3 times every 30 minutes.

### Tube-O-DIALYZER™ Preparation

#### Step 6.

After digestion is complete, centrifuge the tube for 20 seconds at 1,000g.



DURATION

00:00:20

### Tube-O-DIALYZER™ Preparation

#### Step 7.

Replace the cap with the dialysis cap. Do not discard the storage cap as this will be required for storage of DNA.

### Tube-O-DIALYZER™ Preparation

## Step 8.

Place the Tube-O-DIALYZER™ upside down in a 50ml centrifuge tube and centrifuge at 1000xg for 30 seconds to bring the sample onto the dialysis membrane.

### DURATION

00:00:30

### NOTES

**Colin Heath** 23 Jun 2016

NOTE: Do not centrifuge longer or faster than stated to prevent damage to membrane and sample loss.

## Tube-O-DIALYZER™ Preparation

### Step 9.

Remove the Tube-O-DIALYZER™ from the 50ml tube with forceps and keeping it inverted slide into the provided float and dialyze in 500ml 1X TE buffer at room temperature for 18-24 hours with 2-3 buffer changes. Gently swirl tube to mix contents at each buffer change.

### DURATION

18:00:00

### NOTES

**Colin Heath** 23 Jun 2016

NOTE: Cloudy DNA is an indication of incomplete dialysis, therefore dialyze for an additional 24 hours. Change dialysis buffer and mix the content of the Tube-ODIALYZER™ by gently swirling every few hours.

## Tube-O-DIALYZER™ Preparation

### Step 10.

Following dialysis the genomic DNA may be concentrated in the Tube-O-DIALYZER™ using either Tube-O-DIALYZER™ Concentrator (Cat. # 786-144) or Concentrator Solution (Cat. # 786-143). Simply prepare the Concentrator as per the instructions and invert the Tube-O-DIALYZER™ containing your DNA in the solution.

## Tube-O-DIALYZER™ Preparation

### Step 11.

If concentration is not required or following concentration, centrifuge the tube at 1000xg for 1 minute. Replace the dialysis cap with the normal cap. The genomic DNA is now ready for use.

### DURATION

00:01:00