



# **Rapid Transfer Buffer (10X) (Powder)**

## **User Manual**

**Catalog # CRG1040**

Quick, efficient transfer of proteins from SDS-PAGE gels to membranes  
for Western blotting applications

**For research use only. Not for diagnostic or therapeutic procedures.**

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## I. INTRODUCTION

Cohesion Biosciences' Rapid Transfer Buffer is a simple one-component system for quick, efficient transfer of proteins from SDS-PAGE gels to membranes for Western blotting applications. Transfer is completed in 10 to 30 minutes using a standard semi-dry or wet transfer apparatus, respectively. Dedicated, expensive transfer equipment is not needed. The transfer efficiency is equivalent to that observed when using a Tris-Glycine-Methanol transfer buffer.

Rapid Transfer Buffer is a methanol-free, non-hazardous formulation that is compatible with both PVDF and nitrocellulose membranes. It works well with most gel types, including Laemmli, pre-cast, and others.

### **Product Characteristics**

- ※ Fast, efficient transfer in 10 - 30 minutes
- ※ Compatible with standard wet and semi-dry transfer equipment
- ※ Methanol-free, non-hazardous formulation
- ※ Transfers to PVDF and nitrocellulose from poured or pre-cast gels

## II. PREPARATION

### **Stock Solution:**

**1L size:** Dissolve the powder in 1L deionized water for Rapid Transfer Buffer (10X).

**500 ml size:** Dissolve the powder in 500 ml deionized water for Rapid Transfer Buffer (10X).

**100 ml size:** Dissolve the powder in 100 ml deionized water for Rapid Transfer Buffer (10X).

### **Working Solution:**

Prepare 1 L of Rapid Transfer Buffer (1X) by diluting 100 mL of Rapid Transfer Buffer (10X) with 800 mL of deionized water, then add 100 ml Ethanol.

### III. ASSAY PROCEDURE

#### Wet Transfer Protocol

1. Prepare membrane and filter paper for transfer:

a. A blotting membrane and 3 pieces of filter paper should be cut to fit dimensions of the gel.

Note: PVDF membranes must be pre-wetted according to the manufacturer's instructions in 100% methanol prior to equilibration in transfer buffer.

b. Equilibrate the membrane and filter paper in 1X Rapid Transfer Buffer for a minimum of 5 minutes.

2. Following protein electrophoresis, assemble the blotting sandwich following the manufacturer's instructions for the transfer apparatus.

3. Place the blotting sandwich in a wet transfer tank filled with 1X Rapid Transfer Buffer.

4. Transfer constant current 300 - 400 mA at room temperature.

| Current | Time         |
|---------|--------------|
| 300 mA  | 25 to 30 min |
| 350 mA  | 20 to 25 min |
| 400 mA  | 15 to 20 min |

#### Semi-Dry Transfer Protocol

1. Prepare membrane and filter paper for transfer:

a. A blotting membrane and 3 pieces of filter paper should be cut to fit dimensions of the gel.

Note: PVDF membranes must be pre-wetted according to the manufacturer's instructions in 100% methanol prior to equilibration in transfer buffer.

b. Equilibrate the membrane and filter paper in 1X Rapid Transfer Buffer for a minimum of 5 minutes.

2. Following electrophoresis, wash the gel for 2 minutes in deionized water.

3. Pre-equilibrate the gel in 1X Rapid Transfer Buffer for 5 minutes.
4. Assemble the blotting sandwich following the manufacturer's instructions for the semi-dry transfer apparatus.
5. Transfer constant current 300 - 400 mA at room temperature.

| Current | Time         |
|---------|--------------|
| 300 mA  | 25 to 30 min |
| 350 mA  | 20 to 25 min |
| 400 mA  | 15 to 20 min |

**Note:**

- a. For normal PAGE gel, if the molecular weight of the protein is greater than 150 kDa, the transfer time needs to be extended by 5-10 min.
- b. For the thicker PAGE gel, such as 1.5 mm thick gels, the transfer time can be extended by 5-10min
- c. If use higher current, it is recommended to add ice packs to cool down the buffer.

#### IV. TROUBLESHOOTING GUIDE

| Problem                  | Possible Cause  | Solution  |
|--------------------------|---|---|
| Low transfer efficiency  | • Insufficient transfer time  | Not all proteins transfer at the same rate and efficiency. If needed, transfer time may be increased; empirical testing will be required to determine non-standard protocol conditions. |
|                          | • Inadequate equilibration of membrane and/or filter paper in transfer buffer | Completely cover membrane with transfer buffer and incubate for 5 minutes with gentle agitation for best results.   |
|                          | • PVDF membrane not pre-wetted  | Use methanol to wet entire PVDF membrane. Slowly add deionized water (to avoid air bubbles on the membrane). Transfer membrane to Rapid Transfer Buffer for equilibration.              |
| Uneven transfer          | • Incomplete contact between membrane and gel                                 | Always take care to roll out air bubbles between the membrane and gel in the transfer sandwich.   |
|                          | • Incomplete hydration of membrane  | Pre-wet the membranes per the manufacturer's instructions.  |
| Poor transfer of protein | • Inefficient binding of some proteins to membrane                            | Reduce Transfer time to avoid low molecular weight proteins passing through the membrane. Increase transfer time for high molecular weight proteins.                                    |
| Equipment error          | • Incompatible transfer apparatus   | Rapid Transfer Buffer does not require, and is not compatible with specialized transfer devices. Use Rapid Transfer Buffer with a standard semi-dry or wet transfer apparatus.          |
|                          | • Incompatible power supply   | The optimal power supply should be capable of running at constant voltage with specifications of 5 - 250V, 0.01 - 3.0A, and 1 - 300W.   |

## **V. TECHNICAL SUPPORT**

For troubleshooting, information or assistance, please go online to [www.cohesionbio.com](http://www.cohesionbio.com) or contact us at [techsupport@cohesionbio.com](mailto:techsupport@cohesionbio.com)

## **VI. NOTES**