

Rapid Transfer Buffer

<u>Code</u>	<u>Description</u>	<u>Size</u>
N789-1L	Rapid Transfer Buffer	1 liter
N789-4L		4 liters

General Information:

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. Most transfers are complete in 10 to 20 minutes with standard semi-dry or wet transfer apparatus, respectively. Dedicated, expensive transfer equipment is not needed.

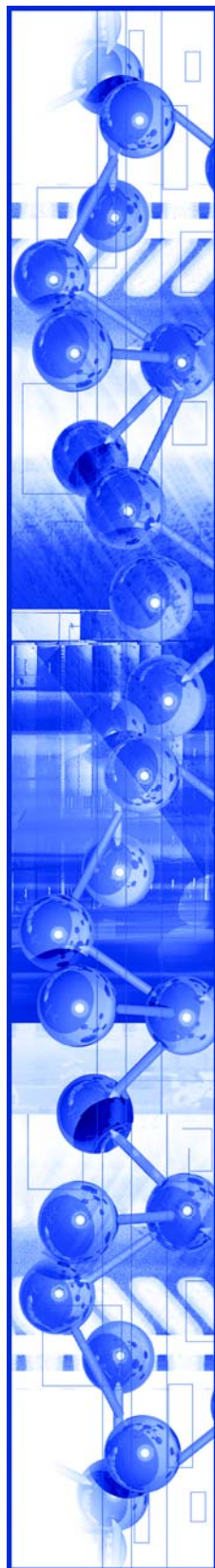
Rapid Transfer Buffer is a methanol-free, non-hazardous formulation that is compatible with both PVDF and nitrocellulose membranes.

Storage/Stability:

Rapid Transfer Buffer is stable at 18 – 26 °C for 6 months.

Application Disclaimer

*For Research Use Only.
Not for Therapeutic or Diagnostic Use.*



Procedure:**Wet Transfer Protocol**

1. Prepare 1 liter of 1X Rapid Transfer Buffer by diluting 100 ml 10X Rapid Transfer with 900 mL of deionized water.
 - a. Prepare membrane and filter paper for transfer: Membrane and 2 pieces of Whatman filter paper should be cut to fit dimensions of the gel.
 - b. → PVDF membranes must be pre-wetted according to manufacturer's instructions in 100% methanol prior to equilibration in transfer buffer.
 - c. Equilibrate membrane and filter paper in 1X Rapid Transfer Buffer for a minimum of 5 minutes.
2. Following electrophoresis, assemble the blotting sandwich following manufacturer's instructions.
3. Place the blotting sandwich in a wet transfer tank filled with 1X Rapid Transfer Buffer.
4. Transfer for 20 minutes at 75V at room temperature.

Semi-Dry Transfer Protocol

1. Prepare 100 ml of 1X Rapid Transfer Buffer by diluting 10 ml 10X Rapid Transfer with 90 mL of deionized water.
 - a. Prepare membrane and filter paper for transfer: Membranes and 2 pieces of Whatman filter paper should be cut to the dimensions of the gel.
 - b. → PVDF membranes must be pre-wetted in 100% methanol prior to equilibration in transfer buffer.
 - c. Equilibrate membrane in 1X Rapid Transfer Buffer for a minimum of 5 minutes.
2. Following electrophoresis, wash the gel for 5 minutes in deionized water.
3. Pre-equilibrate the gel in 1X Rapid Transfer Buffer for 5 minutes.
4. Assemble the blotting sandwich following manufacturer's instructions.
5. Transfer for 10 minutes at 25V at room temperature.

Related Products*Electrophoresis*

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| M317 | Fluorescent Sprint NEXT GEL™, 10% Solution |
| M318 | Fluorescent Sprint NEXT GEL™, 12.5% Solution |

Membrane stains

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| M282 | Proact™ Membrane Stain |
| K793 | Ponceau S Stain |

Blocking Solution

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| M325 | RapidBlock™ Solution |
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Chemiluminescent Substrates

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|------|--|
| N218 | VisiGlo™ Chemiluminescent Substrate |
| N219 | VisiGlo™ Plus Chemiluminescent Substrate |

Stripping Buffer

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| N552 | Gentle Review™ Stripping Buffer |
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X-ray film background reducer

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| N656 | UnDo™ X-Ray Film Background Reducer |
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