StripPRO™ 1 Min Stripping Buffer

V1.5

SP01-500/SP05-100

Store at room temperature
For Research Use Only
Caution: Including corrosive liquid

Introduction

StripPRO™ 1 Min Stripping Buffer effectively removes antibodies from Western blots in one minute. The unconjugated antigens on the stripped membrane are allowed to be reprobed and be detected with chemiluminescent substrates. **StripPRO™ 1 Min Stripping Buffer** is an ideal product for breaking antigen-antibody interaction, saving time and saving conserving samples.

Product Components

StripPRO™ 1 Min Stripping Buffer (SP01-500)

1 Min Stripping Buffer	500 mL	1 bottle
User's manual		

StripPRO™ 1 Min Stripping Buffer (5X) (SP05-100)

1 Min Stripping Buffer (5X)	100 mL	1 bottle
User's manual		

Safety Information

Please wear gloves, lab coat and goggles while operating. Prevent contact product directly. In case of contacting, wash with large amount of water.

Storage

StripPRO™ 1 Min Stripping Buffer could be stored at room temperature. Expiration date is labeled on the bottle or box.

VISUAL PROTEIN

Materials needed but not provided

1. Nitrocellulose or PVDF membrane probed by Western blotting procedure
2. Wash buffer such as phosphate-buffered saline (PBS) or Tris-buffered saline (TBS) with 0.05% Tween-20
3. Primary and secondary antibodies
4. Film or Image captured system

Instruction

NOTE: Prepare 1 Min Stripping Buffer by mixing 1 part of **1 Min Stripping Buffer (5X)** and 4 parts of ddH_2O **(For SP05-100 Only)**.

1. Wash nitrocellulose or PVDF membrane in wash buffer to remove the chemiluminescent substrate.
2. Incubate the membrane in 1 Min Stripping Buffer for 1-3 minutes at room temperature while shaking.
3. Discard 1 Min Stripping Buffer and wash the membrane 3 times in wash buffer.
4. Re-block the stripped membrane and perform immunodetection by Western Blot normal protocol.

Troubleshooting

Problem	Possible cause	Remedy
High background	Not sufficiently blocked after stripping	Optimize blocking conditions
Low signal or no signal	Antigen is not present or in low abundance	Load more protein in the gel
	Antibody concentrations are too low	Increase antibody concentrations
Previous signal obtained	High-affinity antigen-antibody interaction	Incubate at 37 °C for 10-15 minutes
	High sensitive detection reagents used	Detect weak signals first, then strip and detect strong signals in subsequent re-probing