

SDS-OutTM Precipitation Kit

20308

Number Description

20308 SDS-Out Precipitation Kit, contains sufficient reagent for 200mL of sample

Kit Contents:

SDS-Out Precipitation Reagent, 10mL

Spin Cup Columns, 12 each, contains 0.45µm cellulose acetate filter

Microcentrifuge Tubes, 2mL, 12 each

Storage: Upon receipt store products at 4°C. These products are shipped at ambient temperature.

Introduction

The Thermo Scientific SDS-Out Precipitation Kit is for removing excess SDS from small sample volumes. Sodium dodecyl sulfate (SDS) is a detergent typically used for solublizing proteins. Several methods for removing excess SDS from protein samples include prolonged dialysis, anion exchange chromatography and acetone precipitation. These methods are often time-consuming, tedious and unsuitable for low-volume protein samples. The SDS-Out Precipitation Kit is easy to use and provides excellent protein recovery from low-volume samples (Table 1).

Note: SDS-Out Precipitation Reagent does not remove SDS that is bound to protein.

Table 1. Protein recovery after using the Thermo Scientific SDS-Out Precipitation Kit. Each sample contained 0.5mg of protein in 0.5mL. BSA = bovine serum albumin; STI = soybean trypsin inhibitor; OVS = ovalbumin.

Initial SDS	% Protein Recovery*						
Concentration	BSA	Cytochrome C	<u>STI</u>	<u>OVA</u>	Ribonuclease A	Myoglobin	Human IgG
1%	100	100	100	93.8	100	97.9	76.2
0.5%	98.4	100	98.8	98.1	100	100	88.7
0.25%	98.0	100	98.5	99.6	100	100	92.7
0.125%	97.5	100	99.0	99.8	100	100	100

^{*}Protein recovery was evaluated by measuring its absorbance at 280nm.

Procedure for SDS Removal

1. Add one volume of SDS-Out Precipitation Reagent to 20 volumes of protein solution (e.g., 19μL of reagent to 380μL of protein sample). Use the microcentrifuge tube provided in the kit for the precipitation.

Note: Maximum capacity for the Spin Cup Columns is 500µL.

- 2. Vortex to mix. Incubate tube in an ice bath for 20 minutes.
- 3. Centrifuge the tube at $10,000 \times g$ for 10 minutes.
- 4. Transfer supernatant to a spin cup column and centrifuge for 1 minute at $10,000 \times g$ to clarify the supernatant.



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