

## Immunoprecipitation

Typically, complete immunoprecipitation of radio-labeled antigen from extracts of transfected mammalian cells requires between 1-5  $\mu\text{L}$  of polyclonal antiserum, 5-100  $\mu\text{L}$  of hybridoma tissue culture medium, or 1-3  $\mu\text{L}$  of ascites. If more antibody is used than is necessary, non-specific background will increase. The concentration of immunoglobulins in antiserum is approximately 6-8 mg/mL. Supernatant from cultured hybridoma cells typically contains 10-100  $\mu\text{g}/\text{mL}$  immunoglobulin, and ascites fluid contains 5-7 mg/mL immunoglobulin.

- Step 1** Divide the preparation of antigen into two equally sized aliquots and place in microfuge tubes. Adjust the volume of each aliquot to 0.5 mL with I.P. Buffer\*. To one aliquot, add antibody directed against the target protein. To the other aliquot, add the same volume of a control antibody. Gently rock both aliquots for 1 hour at 4°C.
- Step 2** Add Protein-G Sepharose to the antigen-antibody mixture, and incubate for 1 hour at 4°C on a rocking platform.
- Step 3** Centrifuge the Protein-G Sepharose antibody-antigen complex at 10,000xG for 20 seconds at 4°C in a microfuge. Remove the supernatant by gentle aspiration. Add 1 ml of Wash Buffer\*\* and re-suspend the Sepharose beads.
- Step 4** Incubate the re-suspended beads for 20 minutes at 4°C on a rocking platform. This allows time for the I.P. buffer to equilibrate with the fluid trapped between the beads.
- Step 5** Repeat steps 3 and 4 three times. Collect the final washed Protein-G Sepharose antibody-antigen complex by centrifugation at 10,000xG for 20 seconds at 4°C in a microfuge. Take care to remove the last traces of the final wash.
- Step 6** Add reducing gel-loading buffer and boil for 3 minutes.
- Step 7** Remove the Protein-G Sepharose from the complex by centrifugation at 10,000xG for 20 seconds at room temperature in a microfuge. Transfer the supernatant to a fresh tube and separate the sample by method of electrophoresis.

### \*IP Buffer

- 50 mm Tris HCl (pH 7.5)
- 150 mM NaCl
- 0.1% Tween 20 (or 0.1% Nonidet P 40 (NP 40))
- 1 mM EDTA (pH 8.0)
- 0.25% Gelatin
- 0.02% Sodium Azide

### \*Wash Buffer

- 50 mm Tris HCl (pH 7.5)
- 150 mM NaCl
- 0.1% Tween 20 (or 0.1% Nonidet P 40 (NP 40))
- 1 mM EDTA (pH 8.0)