YOU ARE HERE

Protein Purification > Immunoprecipitation

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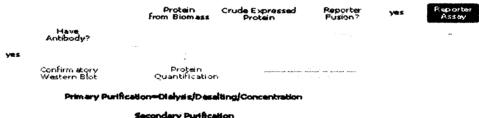
Immunoprecipitation is often used to purify a target protein from solution. Two separate approaches are commonly employed to perform immunoprecipitations. In the first, an antibody specific to the target protein is added to the protein mixture. The antibody-antigen complex is then precipitated from the solution using an insoluble resin that binds to the antibody complex (such as Protein A or Protein G immobilized on a solid support). Unbound proteins are removed by washing the resin, and the protein is eluted from the resin for further analysis. A drawback to this method is that when the antigen is eluted from the resin, it is contaminated with the antibody. This may complicate analysis of the purified

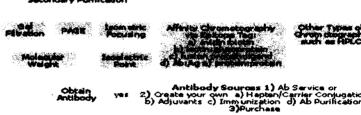
Pierce developed Seize Primary and Seize X Immunoprecipitation Kits to purify a protein by immunoprecipitation without contaminating the preparation with antibody. Using the Seize Kits, an antibody is covalently linked to the resin. Then the antibody coupled to resin is added to a crude protein mixture to capture and precipitate the antigen. The antigen is eluted from the resin while the antibody remains covalently bound to the resin. This simplifies the analysis of the purified protein by eliminating the large amount of antibody that would normally interfere with analyses such as SDS-PAGE or protein assays.

For more information on immunoprecipitation, see Current Protocols in Protein Science, Chapter 9, John Wiley & Sons, Inc.

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· Go to Primary Purification





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To Protein Function



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