## **Streptavidin Magnetic Particles**



Product No. 544050 (2 mL), 544051 (5 mL)

## 1. General Description

The Abraxis' superparamagnetic nanoparticles are coupled with a biomolecule, such as Streptavidin, and are utilized in the magnetic separation and isolation of biotin-labeled proteins and nucleic acids used in protein interaction studies, DNA-protein pulldowns, and purification of biotin-labeled proteins and nucleic acids. The particles have a large surface area with high capture efficiencies.

2. Safety Instructions

Reagents contain 0.05-0.1% sodium azide as a preservative. Sodium azide may react with lead or copper plumbing to produce metal azides which might cause explosion. To prevent azide accumulation in plumbing, flush with copious amounts of water immediately after disposal.

3. Storage and Stability

The Streptavidin Magnetic Particles should be stored in the refrigerator (4-8°C). The reagent must be allowed to reach room temperature (20-25°C) before use and may be used until the expiration date on the box. Do not freeze, dry, or centrifuge the particles as they may result in loss of binding activity and aggregation.

4. Test Principle

Streptavidin magnetic particles are incubated with the solutions containing biotin-labeled compounds and then separated by magnets. After the unbound particulates are washed from the particles, the bound biotin are eluted from the particles using the elution buffer. The particles are then magnetically separated from the eluted solution, which is removed manually.

5. Warning and Precautions

-This product is for in vitro research use only, do not use in vivo.
-Do not freeze the reagent.
-Prior to use, ensure that the product has not expired by verifying that the date of use is prior to the expiration date on the label.
-Ensure that reagent bottle caps are tight after each use to prevent drying of reagents.
-Mistakes in handling the test can also cause errors. Possible sources for such errors can be:

Inadequate storage conditions of the test kit (or reagents), incorrect pipetting sequence or inaccurate volumes of the reagents, too long or too short incubation times, and/or short magnetic separation times.

6. Characteristics

Particle mean diameter: ~0.5 µm Particle concentration: 5 mg/mL Binding capacity: ≥60 µg biotin/mg of particles

7. Antibody Isolation	Ordering Information			
A. Materials Provided	Description	Size (mL)	Part Number	
1. Streptavidin magnetic particles, 5 mg/mL				
	Streptavidin Particles	2, 5	544050/51	
B. Additional Materials (not provided with the kit)	Other related preducts			
1. Binding/Wash Buffer: TBS - 0.05% Tween 20 detergent	Other related products:			
2. Elution Buffer: 0.1 M Glycine pH 2.0, 5 mL	Biotin Magnetic Particles	2,5	544000/01	
3. Neutralization Buffer: 1M Tris pH 8.0, 1 mL	Multi-6 Microcentrifuge Separato	or	472260	
4. Micro-pipettes with disposable plastic tips (10-200 and 200-1000 μL)	Solo Tube Separator		472270	
5. 1.5 mL or 2.0 mL Eppendorf or microcentrifuge vials	15/50 mL Tube Separator		472250	
6. Timer	Multi Purpose Magnetic Separat	tor (15/50/microcentrifuc		
7. Rotator			jc) 472200	
8. Distilled or deionized water				
9. Vortex mixer				
10. Solo or Multi-6 Microcentrifuge Separator (PN 472270; PN 472260)				
C. Procedures				
1. Add 100 $\mu$ L (0.5 mg) of particles to 1 mL of binding buffer in each tube to wash particles.				
2. Magnetically separate using a magnetic separator for 2 minutes or when the supernatant is clear.				
3. Remove the supernatant and wash once more by adding 1 mL of binding buffer.				
4. Repeat step 2 and remove the supernatant.				
5. Resuspend particles by adding 450 uL of binding buffer.		General Limited Warranty: Abraxis LLC warrants the products manufactured by the Company, against defects and workmanship when used in accordance with the applicable instructions for a period not to extend beyond the product's printed expiration date. Abraxis LLC makes no other warranty, expressed or implied. There is no warranty of merchantability or fitness for a particular purpose.		
6. Add 50 μL of serum or cell culture supernatant to the particles.				
Note: Sample volume can be modified according to user preference. If the sample volume is < 500µL, dilute it to a final volume of 500µL with Binding/Wash Buffer.				
7. Gently mix using vortex or rotator for 30 minutes.				
8. Magnetically separate using a magnetic separator for 2 minutes or when the supernatant is clear.				
9. Remove supernatant and wash with 0.5 mL Binding/Wash buffer to remove unbound proteins.	For ordering or technical assistance con	tact: Abraxis		
10. Repeat steps 8 and 9 once more. Remove supernatant.	For ordening or technical assistance con		road Drive	
11. Add 100 $\mu$ L of elution buffer to particles and mix well.			ster, PA 18974 5) 357-3911	
12. Incubate at room temperature for 10 minutes with occasional gentle mixing or vortex.			5) 357-3911 5) 357-5232	
13. Separate for 2 minutes and remove the eluent to a new tube containing 15 $\mu$ L of neutralization buffer.			info@abraxiskits.com	
		WEB:	<u>www.abraxiskits.com</u> 101215	