

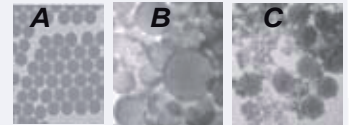
Bio-Adembeads Protein A/G or PAG for IP and co-IP

Immunoprecipitation (IP) is a child's play

- **Higher Efficiency** : effective (co) -IP and less background due to optimized magnetic nanoparticles
- **Simplify and speed the procedure** :
 - No centrifugation
 - Less incubation time and reduced washing steps
 - No preclearing step required
 - Clear visualization of the magnetic nanoparticles
 - No centrifugation
- **Gentle system** :
 - Minimal loss of protein
 - Minimal stress versus centrifugation
 - Isolation of large protein complexes.
- **Flexible system** : compatible with a wider range of antibodies
 - **Bio-Adembeads Protein A** coated with Protein A
 - **Bio-Adembeads Protein G** coated with Protein G
 - **Bio-Adembeads PAG** coated with Protein AG a genetically-engineered protein that combines the IgG binding domains of both Protein A and G.

Why Choose Adembeads?

Uniform size = less variability



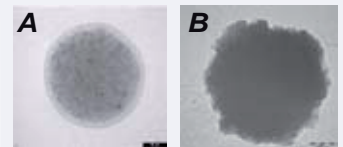
A - Calibrated Adembeads: unique size

B-C - Random size distribution of magnetic particles from competitors

Small size = large surface area

"The small size offers an increased surface area for binding and eliminates the sedimentation problems that affect larger particles."

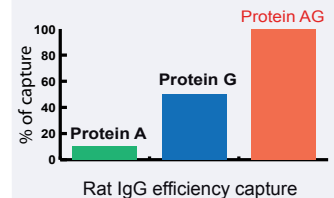
Homogeneous surface = less background



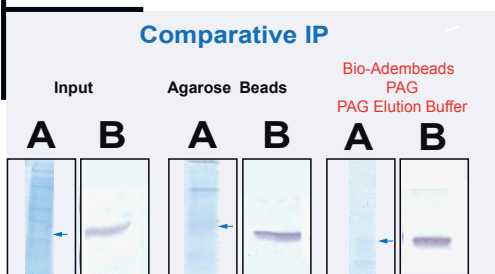
A - Core shell structure of Adembeads: defined surface area

B - Magnetic particle from competitor

Available with Protein A/G or AG

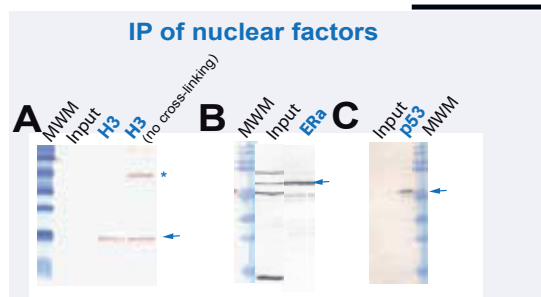


IP results obtained with Bio-Adembeads Protein A/G and PAG



Immunoprecipitation of acetyl-Histone H3 from HeLa cells with Bio-Adembeads PAG and Elution PAG Buffer.

A - Gel stained for total protein with coomassie blue
B - Gel was transferred for Western Blot



Western Blot analysis of immunoprecipitated nuclear proteins using Bio-Adembeads Protein A/G and cross-linking procedure and PAG Elution Buffer.

A - Acetyl Histone H3 (*Heavy IgG chain)
B - Estrogen receptor alpha (ERa)
C - p53 tumor suppressor protein