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## Protocol specific for ARA9 (35-2) Antibody (NB100-127)

[[URL:https://www.novusbio.com/products/aip-ara9-antibody-35-2\_nb100-127]][[Caption:ARA9 (35-2) Antibody]]

1. Wash plate twice with cold PBS.

NB100-127 Protocol

- 2. Add 1 mL lysis buffer into P100 plate, sit on ice for 20 min with gentle shaking, centrifuge 14,000rpm/10min/4C, take supernatant.
- 3. Preclear the lysate with 50 uL protein G slurry, tumble, 45 min/4C, followed by a centrifuge 14,000rpm/15min/4C.
- 4. Add 1 ug antibody (3 uL anti-ARA9 antibody I used) to 50 uL protein G slurry, add 500 uL cold PBS, tumble, 1 hr/4C. Wash Ab/beads twice by adding PBS. Spin down beads by centrifuge at 1000g/1 min.
- 5. Add precleared lysate into pre-bond Ab/protein G complex, tumble O/N, 4C Spin down beads by centrifuge at 1000g/1 min.
- 6. Wash beads five times with lysis buffer.
- 7. Add SDS sample buffer to beads.