

Cell Lysis Buffer for ELISA

Catalog Number EA-0001

(For Research Use Only)

Preparation of Cell Lysate from Tissue

- 1. Weigh tissue sample and add 1ml of cell lysis buffer to 1mg of tissue.
- 2. Homogenized tissues with (PowerGen 125 or equivalent) on ice.
- 3. Sonicate lysates briefly on ice.
- 4. Centrifuge the sample at 10,000 RPM for 5 minutes to pellet the tissue debris.
- 5. Collect supernatant, measure the protein concentration of supernatant. The supernatant then can be used for assay. The supernatant also can be aliquoted, and frozen at -80°C. Avoid multiple freeze/thaws.
- 6. Dilute the cell lysate in the provided Diluent Buffer to the final 10ug/100ul per well.