



EDTA BUFFER CONCENTRATE (20X)

(20 mM Ethylenediaminetetraacetic Acid)

Storage: 2-8°C

Catalog No: B04C-100 100ml

Kit Format:

100 ml liquid form;

Recommended Protocol:

Dilute EDTA solution 1:20 with deionized or distilled water before using; This will give a 1 mM ready-touse EDTA buffer, pH 8.0. Refer to HIER Procedure for details.

Precaution:

Wear gloves and take other necessary laboratory safety procedure.

Storage:

Store at 2-8°C. Do not freeze.

Remarks:

For research use only;

HIER Procedure for FFPE Tissue Sections:

- 1. Deparaffinize in xylene and rehydrate tissue in graded alcohols.
- 2. Block endogenous peroxidase with hydrogen peroxide solution for 10 minutes.
- 3. Rinse with distilled water.
- 4. Rinse with PBS.
- 5. Heat ready-to-use EDTA buffer in a beaker on a hot plate until the temperature of the buffer reaches 93°C-95°C. Place slides in heated buffer for 10-15 minutes.
- 6. Remove beaker with slides from the hot plate and allow it cool for 25-30 minutes.
- 7. Rinse slides with PBS 3 times, 2 minutes each time.
- 8. Apply avidin/biotin blocking or general blocking if required
- 9. Start immunostaining procedure.