

## Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

Weitere Information auf den folgenden Seiten! See the following pages for more information!



### Lieferung & Zahlungsart

siehe unsere Liefer- und Versandbedingungen

## Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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# Instructions For Use ABM-IFU

Rev. Date: August 1, 2019

**Revision: 3** 

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P.O. Box 3286 - Logan, Utah 84323, U.S.A. - Tel. (800) 729-8350 - Fax (435) 755-0015 - www.scytek.com

## UltraTek Alkaline Phosphatase

Uses/Limitations: A Ready-to-use streptavidin-Alkaline Phosphatase enzyme label designed for 3-step

immunohistochemistry protocols. Formulated to provide optimal staining with an incubation for 10 minutes. May be used with automated systems, reagent jars, and

manual dropping/pipetting.

Uses/Limitations: Not to be taken internally.

For In-Vitro Diagnostic use only.

Histological applications.

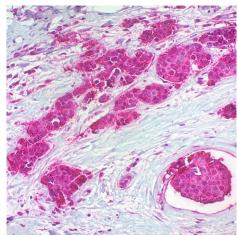
Do not use if reagent becomes cloudy. Do not use past expiration date.
Use caution when handling reagent.

Non-Sterile.

Control Tissue: Any well-fixed tissue.

#### **Availability/Contents:**

Item #	<u>Volume</u>
ABM008	8 ml
ABM015	15 ml
ABM125	125 ml
ABM500	500 ml
ABM999	1000 ml



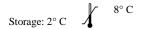
Human Breast stained with Permanent Red and UltraTek Alkaline Phosphatase within an IHC protocol.

**Storage:** Store at 2-8°Centigrade. Product is stable for 18 months from date of manufacture.

#### Procedure:

Allow reagents to come to room temperature before use.

- 1. Deparaffinize and rehydrate tissue section.
- 2. If required, incubate tissue in digestive enzyme.
- 3. Wash 4 times in buffer.
- 4. Place slide in protein block (ScyTek's "Superblock") and incubate 5-10 minutes at room temperature to block nonspecific background staining. Note: Do not exceed 10 minutes or there may be a reduction in desired stain.
- 5. Wash 1 time in buffer.
- Apply primary antibody and incubate according to manufacturer's protocol.
- 7. Wash 4 times in buffer.









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- 8. Place slide in biotinylated link secondary antibody, and incubate per instructions.
- 9. Wash 4 times in buffer.
- 10. Place slide in UltraTek Alkaline Phosphatase, and incubate for 10 minutes.
- 11. Rinse 4 times in buffer.
- 12. Place slide in appropriate chromogenic substrate and incubate until desired reaction is achieved.
- 13. Counterstain and coverslip.

#### **Troubleshooting Guide**

#### Overstaining:

- 1. Concentration of the primary antibody was too high or the incubation time was too long.
- 2. Temperature during incubation was too high.
- 3. Incubation time with link antibody or streptavidin/enzyme label was too long.

#### Nonspecific Background Staining:

- 1. Rinsing between steps was inadequate.
- 2. Tissue was allowed to dry with reagents on.
- 3. Folds in tissue trapped reagents.
- 4. Tissue contains endogenous enzyme.
- 5. Tissue contains endogenous biotin.
- 6. Antigen migrated in tissue.
- 7. Excessive tissue adhesive on slides.
- 8. Inadequate blocking with protein block.

#### Weak Staining:

- 1. Primary antibody concentration was too low or incubation time was too short.
- 2. Reagents are past their expiration date.
- 3. Reagent is reaching the end of its useful life.
- 4. Counterstain or mounting media were incompatible and dissolved the chromogen reaction product.
- 5. Room temperature was excessively cool.
- 6. The primary antibody does not recognize an antigen that survives fixation and embedding in high enough amounts.
- 7. Excessive incubation with protein block (Super Block or normal serum).

#### No Staining:

- 1. Steps were inadvertently left out.
- 2. There is no relevant antigen in the tissue.
- 3. Chromogenic substrate does not match enzyme label.
- 4. One or more components have been inactivated.

