

Instructions For Use A00058-IFU-IVD

Rev. Date: June 10, 2022

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P.O. Box 3286 - Logan, Utah 84323, U.S.A. - Tel. (800) 729-8350 - Tel. (435) 755-9848 - Fax (435) 755-0015 - www.ScyTek.com

Alpha-1-Fetoprotein (AFP); Polyclonal

Catalog Number	<u>Format</u>
A00058-0002	(Ready-To-Use)
A00058-0007	(Ready-To-Use)
A00058-0025	(Ready-To-Use)

<u>Volume</u> 2 ml 7 ml 25 ml

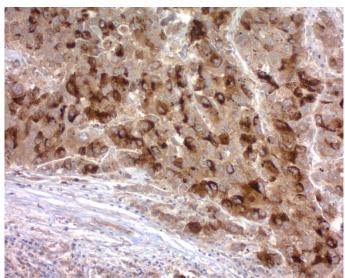
Intended Use

For In Vitro Diagnostic use. This antibody is intended for the qualitative visualization of the anatomical elements listed in the Specificity section. It is intended to be used within an Immunohistochemistry (IHC) procedure on formalin-fixed paraffin-embedded (FFPE) human tissue followed by visualization by light microscopy. Any diagnostic interpretation of the results of this antibody is to be complemented by morphological studies using proper controls and should be evaluated within the context of the patient's clinical history and other diagnostic tests by a qualified pathologist.

Dilution Deads to Lless No forther dilution required

Description

Titer/Working Dilution: Ready-to-Use: No further dilution required.	
•	Concentrate: N/A
Species:	Rabbit
İmmunogen:	Alpha-1-Fetoprotein isolated from human cord serum.
Clone:	Polyclonal
Isotype:	N/A
Entrez Gene ID:	174 (Human)
Hu Chromosome Loc.: 4q13.3	
Synonyms:	Alpha fetoglobulin, FETA, HPAFP
Mol. Wt. of Antigen:	70kDa
Format:	Ready-To-Use antibody has been pretitered and quality
	controlled to work on formalin-fixed paraffin-embedded tissue
	sections. No further titration is required.
	Concentrate: N/A
Specificity:	This antibody reacts with a 70 kD protein identified as alpha-1-
	fetoprotein.
Background:	Alpha-1-Fetoprotein is normally synthesized in the liver, intestinal
	tract, and yolk sac of the fetus. Antibody to AFP has been found
	to be useful in detecting hepatocellular carcinomas (HCC) and
	germ cell neoplasms, such as yolk sac tumors.
Species Reactivity:	Human, Others-not known
Positive Control:	Fetal liver, Hepatocellular Carcinoma.
Cellular Localization:	Cytoplasmic
Microbiological State:	Nonsterile.
-	



Human Liver Ca stained using α-Fetoprotein; Polyclonal. Pretreatment with Tris-EDTA HIER Solution (10x) pH 9.0 for 5 minutes, PolyTek Anti-Rabbit Polymerized HRP and DAB Chromogen/Substrate (High Contrast). Counterstained with Hematoxylin, Mayer's (Lillie's Modification). Final magnification 200X.

Materials and Reagents Required but not Provided

- 1. Control tissue and reagents
- 2. Xylene, graded alcohols, and deionized/distilled water
- 3. Antibody Diluent.
- 4. IHC detection system. Suggested: ScyTek Cat# ABZ125 "CRF Anti-Polyvalent HRP
- Polymer" and ScyTek Cat# ACV500 "DAB Chromogen/Substrate Kit (High Contrast)".
- 5. Wash buffer for rinses (ScyTek Cat# TBT500)
- 6. HIER Retrieval Solution
- 7. Hematoxylin counterstain and bluing reagent (ScyTek Cat# HMM500 and BRT500) 8. Mounting medium and coverslips

Note: ScyTek Laboratories has a wide range of IHC reagents and ancillaries that can be found at scytek.com.

Procedure

1. **Tissue Section Pretreatment (Recommended):** Staining of formalin fixed, paraffin embedded tissue sections is enhanced by pretreatment with pH 8-9 HIER Solution (see ScyTek catalog# ETA or TES for instructions).

2. **Primary Antibody Incubation Time:** We suggest an incubation period of 30 minutes at room temperature. However, depending upon the fixation conditions and the staining system employed, optimal incubation should be determined by the user.

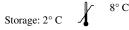
3. Visualization: For maximum staining intensity we recommend the "CRF Anti-Polyvalent HRP Polymer" (ScyTek catalog# ABZ125, see IFU for instructions) combined with the "DAB Chromogen/Substrate Bulk Pack (High Contrast)" (ScyTek catalog# ACV500, see IFU for instructions).

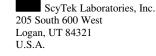
Storage and Stability

Do not Freeze. Store at 2-8°C. Return to 2-8° immediately after use. Do not use after expiration date printed on label. Verify visually that antibody has not been contaminated before use. Do not use if reagent becomes cloudy or precipitates.



Ec REP Emergo Europe Prinsessegracht 20 2514 AP The Hague, The Netherlands







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Limitations

Immunohistochemistry is a complex technique involving both histological and immunological detection methods. Tissue processing and handling prior to immunostaining can cause inconsistent results. Variations in fixation and embedding or the inherent nature of the tissue specimen may cause variations in results. Endogenous peroxidase activity or pseudoperoxidase activity in erythrocytes and endogenous biotin may cause non-specific staining depending on detection system used. This data sheet's recommendations and procedures were validated using ScyTek IHC reagents and may not be suitable for other detection systems.

Precautions

1. Contains Sodium Azide as a preservative (0.09% w/v), do not ingest. Sodium Azide may react with lead and copper plumbing to form highly explosive metal azides. Upon disposal, flush with large volumes of water to prevent azide build-up in plumbing. This product contains no hazardous material at a <u>reportable concentration</u> according to U.S. 29 CFR 1910.1200, OSHA Hazardous Communication Standard and EC Directive 91/155/EC.

2. Do not pipette by mouth.

3. Avoid contact of reagents and specimens with skin and mucous membranes.

Avoid microbial contamination of reagents or increased nonspecific staining may occur.
The user must validate any procedures and recommendations that differ from this data

sheet.

6. The SDS may be found at scytek.com

References

1. MacDonald et al. Clin Chim Acta 87: 367, 1978.

2. Jacobson et al. Am J Surg Pathol 5: 257, 1981.

Warranty

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ScyTek Laboratories, Inc. 205 South 600 West Logan, UT 84321 U.S.A.



Ec REP Emergo Europe Prinsessegracht 20 2514 AP The Hague, The Netherlands