

Progesterone Receptor (Marker of Progestin Dependence); Clone PR484 (Concentrate)

Availability/Contents:

<u>Item #</u>	<u>Volume</u>
RA0263-C.5	0.5 ml

Description:

Species:	Mouse
Immunogen:	Recombinant human Progesterone Receptor protein
Clone:	PR484
Isotype:	IgG1, kappa
Entrez Gene ID:	5241 (Human)
Hu Chromosome Loc.:	11q22.1
Synonyms:	NR3C3; Nuclear receptor subfamily 3 group C member 3; PGR; PR; PRA; PRB; Progesterone receptor form A; Progesterone receptor form B.
Mol. Weight of Antigen:	PR-A (81kDa) and PR-B (116kDa)
Format:	200µg/ml of Ab purified from Bioreactor Concentrate by Protein A/G. Prepared in 10mM PBS with 0.05% BSA & 0.05% azide.
Specificity:	This antibody is specific to the progesterone receptor and shows minimal cross-reaction with other members of the family.
Background:	The progesterone receptor is expressed as two major isoforms: PR-A (81kDa) and PR-B (116kDa). Expression of the progesterone receptor has been suggested to reflect an intact estrogen regulatory machinery and therefore predict better clinical response to endocrine therapy than estrogen receptor expression alone. This antibody is excellent for immunohistochemical staining of formalin-fixed, paraffin-embedded tissues.
Species Reactivity:	Human. Others not known.
Positive Control:	T47-D cells or breast cancers.
Cellular Localization:	Nuclear
Titer/ Working Dilution:	Immunohistochemistry (Frozen and Formalin-fixed): 0.5-1 µg/ml Western Blotting: 0.5-1 µg/ml
Microbiological State:	This product is not sterile.

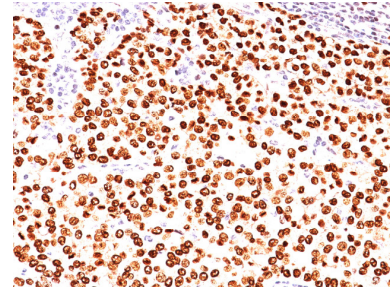
 Storage: 2° C  8° C


 ScyTek Laboratories, Inc.
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Uses/Limitations: Not to be taken internally.
 For Research Use Only.
 This product is intended for qualitative immunohistochemistry with normal and neoplastic formalin-fixed, paraffin-embedded tissue sections, to be viewed by light microscopy.
 Do not use if reagent becomes cloudy.
 Do not use past expiration date.
 Non-Sterile.



Ordering Information and Current Pricing at www.scytek.com

Formalin-fixed, paraffin-embedded human breast carcinoma stained with Progesterone Receptor; Clone PR484.

Procedure:

1. **Tissue Section Pretreatment (Required):** Staining of formalin fixed, paraffin embedded tissue sections is significantly enhanced by pretreatment with Citrate Plus (ScyTek catalog# CPL500).
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 “UltraTek HRP Anti-Polyvalent Lab Pack” (ScyTek catalog# UHP125, see IFU for instructions) combined with the “DAB Chromogen/Substrate Bulk Pack (High Contrast)” (ScyTek catalog# ACV500, see IFU for instructions).

Precautions: Contains Sodium Azide as a preservative (0.09% w/v).
 Do not pipette by mouth.
 Avoid contact of reagents and specimens with skin and mucous membranes.
 Avoid microbial contamination of reagents or increased nonspecific staining may occur.
 This product contains no hazardous material at a reportable concentration according to U.S. 29 CFR 1910.1200, OSHA Hazardous Communication Standard and EC Directive 91/155/EC.


References:

1. Press M, et al. Steroids. 2002 Aug; 67(9):799-813.
2. Mote P, et al. J Clin Pathol., 2001; 54: 624-630.

Warranty:

No products or “Instructions For Use (IFU)” are to be construed as a recommendation for use in violation of any patents. We make no representations, warranties or assurances as to the accuracy or completeness of information provided on our IFU or website. Our warranty is limited to the actual price paid for the product. ScyTek Laboratories, Inc. is not liable for any property damage, personal injury, time or effort or economic loss caused by our products. 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.

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