



**Revision: 1** 

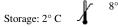
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## Tyrosinase; Clone T311 (Ready to Use)

Availability/Contents:	<u>Item #</u> A00132-0002 A00132-0007 A00132-0025	<u>Volume</u> 2 ml 7 ml 25ml	
Description:			
Species: Immunogen: Clone: Isotype: Format: Specificity:	T311 Mouse IgG1 This antibody is provided Tyrosinase has been sho	e protein was used as the immunogen for the T311 antibody. in a phosphate buffer saline containing 1% BSA. wn to be a very specific marker for melanomas. Cross reactivity with ssues tested has not been reported.	
Background:	pathway and is required f (Hu, 2011). Tyrosinase is biosynthesis, converting t DOPAquinone. The pigm cuticle formation in insect This melanocyte-specific melanosome. Tyrosinase sites, 17 cysteine residue 1 C-terminal TMD. It fold copper ions are incorpora hydroxylation of monophe diphenols to o-quinones ( synthesis. This protein is melanoma and an import	Tyrosinase (Tyr) is a trans-membrane glycoprotein, plays a key role in the melano synthetic pathway and is required for the synthesis of both types of melanin, eumelanin and pheomelanin (Hu, 2011). Tyrosinase is the rate limiting enzyme catalyzing the first two steps in the melanin biosynthesis, converting tyrosine to L-dihydroxy-phenylalanine (DOPA) and subsequently to DOPAquinone. The pigmentation of skin, the browning of vegetables, wound healing and cuticle formation in insects are some of the major responsibilities performed by tyrosinase. This melanocyte-specific enzyme is localized to the post-Golgi compartment termed the melanosome. Tyrosinase consists of 533 amino acids along with 7 occupied N-glycosylation sites, 17 cysteine residues grouped in 2 cysteine-rich domains, 2 copper binding domains, and 1 C-terminal TMD. It folds in the ER and is transported to the trans-Golgi network where two copper ions are incorporated and performs different catalytic reactions such as the hydroxylation of monophenols to o-diphenols (cresolase activity) and the oxidation of o-diphenols to o-quinones (catechol oxidase activity). This process initiates the melanin synthesis. This protein is related with severe skin diseases such as type 1 albinism and melanoma and an important target for anti-melanoma vaccine therapies Anti-tyrosinase antibodies may be applied for immunotherapy in patients with malignant melanoma .	
Species Reactivity: Positive Control:			
Cellular Localizatio	n: Cell Membrane / Membra	ine raft.	
Titer/ Working Dilu Microbiological Sta			





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Ordering Information and Current Pricing at www.scytek.com

## Instructions For Use A00132-IFU-IVD

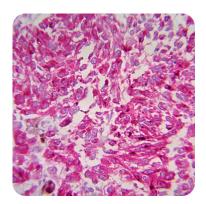
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**Uses/Limitations:** 

Not to be taken internally. For In Vitro Diagnostic Use. 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. Use caution when handling reagents. Non-Sterile.



Human Melanoma stained using UltraTek Alk-Phos and Permanent Red Chromogen.

## Procedure:

- 1. **Tissue Section Pretreatment (Highly Recommended):** Staining of formalin fixed, paraffin embedded tissue sections is significantly enhanced by pretreatment with Citrate Plus (ScyTek catalog# CPL500).
- 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 <u>reportable concentration</u> according to U.S. 29 CFR 1910.1200, OSHA Hazardous Communication Standard and EC Directive 91/155/EC.

## **References:**

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- Han HY, Lee JR, Xu WA, Hahn MJ, Yang JM, Park YD. Effect of Cl- on tyrosinase: complex inhibition kinetics and biochemical implication. J Biomol Struct Dyn. 2007 ;25(2):165-71.
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- Francis E, Wang N, Parag H, Halaban R, Hebert DN. Tyrosinase maturation and oligomerization in the endoplasmic reticulum require a melanocyte-specific factor. J Biol Chem. 2003 ;278(28):25607-17.
- 5. Hu HH, Guedj M, Descamps V, Jouary T, Bourillon A, Ezzedine K, Taieb A, Bagot M, Bensussan A, Saiag P, Grandchamp B, Basset-Seguin N, Soufir N. Assessment of tyrosinase variants and skin cancer risk in a large cohort of French subjects. J Dermatol Sci. 2011;64(2):127-33.
- 6. Yin SJ, Si YX, Wang ZJ, Wang SF, Oh S, Lee S, Sim SM, Yang JM, Qian GY, Lee J, Park YD. The effect of thiobarbituric acid on tyrosinase: inhibition kinetics and computational simulation. J Biomol Struct Dyn. 2011 Dec;29(3):463-70.
- 7. K B, Purohit R. Mutational analysis of TYR gene and its structural consequences in OCA1A. Gene. 2013;513(1):184-95.





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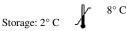
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