

P.O. Box 8522 Portland, Maine 04104 207-856-6620 207-856-6864 (fax) Immunochemicals for Infectious Disease Research

www.ViroStat-Inc.com

MONOTOPE™

product information for

## Candida albicans

## I. Monoclonal Antibody (Mouse) Specificity

Product No.'s Ig class

6401 NA IgG2a ELISA titer > 1:25,000 vs. all strains examined. Marginal IFA staining.

6402 NA IgM Reactive with all C. albicans strains tested (9) + C. glabrata & C. tropicales. ELISA & IFA\*

**II. Purified Preparations** 

Product No.'s

6401 6402

ns Product No.'s 6

MONOTOPE™ purified preparations consist of >90% pure mouse monoclonal antibody which has been purified from ascites fluid or culture medium by protein A chromatography or sequential differential precipitations. The final preparation is formulated to a protein concentration of 100 µg/ml in 0.01 M phosphate buffered saline, pH 7.2 and contains 0.1% sodium azide. Each vial contains 1.0 ml. This product contains no stabilizing proteins and should be stored at 2-8°C until ready for use.

Working dilution must be determined by the user. Suggested starting ranges are 1:10-1:50 for IFA and 1:20-1:200 for ELISA.

## THESE PRODUCTS ARE FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES

III. Fluorescein Conjugates

Product No.'s

NA

NA

These MONOTOPE™ products consist of purified monoclonal antibody conjugated with high purity isomer I of fluorescein isothiocyanate. Care is taken to ensure complete removal of any free fluorescein from the final product. The final preparation is formulated to an antibody concentration of of 100 µg/ml in 0.01 M phosphate buffered saline, pH 7.2 containing 0.1% sodium azide plus bovine serum albumin at 10 mg/ml. Each vial contains 1.0 ml. This product should be stored at 2-8°C until ready for use. Avoid repeated freeze-thawing by storing multiple aliquots at -20°C. Applications for these products include direct FA staining of target antigen in a permissive tissue culture system. Working strength must be determined by the user for each specific application but a starting range of 1:5 - 1:20 is recommended. Acetone fixation of the antigen source is recommended prior to staining.

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Comments: \* Negative vs. C. parapsilosis and C. krusei