

Polyclonal Antibody to *Candida albicans* - FITC

Alternate names:	C. albicans
Catalog No.:	BP1006F
Quantity:	1 ml
Concentration:	4-5 mg/ml (OD280 nm, E0.1% = 1.4)
Background:	<p><i>Candida albicans</i> is the most frequently isolated fungal pathogen of humans, affecting immunocompromised patients ranging from premature infants to AIDS sufferers. Systemic infections have an attributed mortality of 30-50%. <i>C. albicans</i> is a diploid organism which has eight sets of homologous chromosomes. It has a genome of approximately 16 Mb (haploid), about 30% greater than <i>S. cerevisiae</i> (baker's yeast).</p>
Host:	Rabbit
Immunogen:	<i>Candida albicans</i> , type A (ATCC#32354)
Format:	<p>State: Liquid purified Ig fraction (> 95% pure) Purification: Protein A Chromatography Buffer System: 0.01 M PBS, pH 7.2 containing 0.09% Sodium Azide as preservative and 10 mg/ml BSA as stabilizer Label: FITC – Highly purified Isomer I of Fluorescein Isothiocyanate Care is taken to ensure complete removal of any free fluorescein from the final product</p>
Applications:	<p>Suitable for use in Double-Diffusion and CIE, direct IFA, ELISA and Immunohistochemistry. Use neat in gel-precipitin reactions. Direct FA staining of target antigens in a permissive tissue culture system. Acetone fixation of the antigen source is recommended prior to staining. Other applications not tested. Optimal dilutions are dependent on conditions and should be determined by the user.</p>
Specificity:	<p>Recognizes numerous proteins in a soluble <i>C. albicans</i> extract (IEP). Has not been absorbed and does cross-react with other yeasts. Negative against human serum, urine and spinal fluid.</p>
Storage:	<p>Store the antibody undiluted at 2-8°C for one month or (in aliquots) at -20°C for longer. This product is photosensitive and should be protected from light. Avoid repeated freezing and thawing. Shelf life: one year from despatch.</p>
Product Citation:	1. Lies M. E. Vanhee, Wouter Meersseman, Katrien Lagrou, Johan Maertens, Hans J. Nelis, and Tom Coenye Rapid and direct quantification of viable <i>Candida</i> species in whole blood using immunomagnetic separation and solid-phase cytometry J. Clin. Microbiol., Feb 2010; 10.1128/JCM.00035-10.
General References:	1. Brand, A., et al., (2008), "An Internal Polarity Landmark is Important for Externally Induced Hyphal Behaviors in <i>Candida albicans</i> ", <i>Eukaryotic Cell</i> , 7(4): 712–720.

For research and in vitro use only. Not for diagnostic or therapeutic work.

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2. Fratti, R.A., et al., (1998), "Endothelial Cell Injury Caused by *Candida albicans* Is Dependent on Iron", *Infection and Immunity*, 66(1): 191–196.
3. Tsuchimori, N., et al., (2000), "Reduced Virulence of HWP1-Deficient Mutants of *Candida albicans* and Their Interactions with Host Cells", *Infection and Immunity*, 68(4): 1997–2002.
4. Phan, Q.T., et al., (2005), "N-cadherin Mediates Endocytosis of *Candida albicans* by Endothelial Cells", *The Journal of Biological Chemistry*, 280(11): 10455–10461.
5. Phan, Q.T., et al., (2000), "Role of Hyphal Formation in Interactions of *Candida albicans* with Endothelial Cells", *Infection and Immunity*, 68(6): 3485–3490.
6. Martinez-Lopez, R., et al., (2006), "*Candida albicans* Ecm33p is Important for Normal Cell Wall Architecture and Interactions with Host Cells", *Eukaryotic Cell.*, 5(1), 140–147.
7. Palmer, G.E., et al., (2005), "The *Candida albicans* Vacuole is Required for Differentiation and Efficient Macrophage Killing", *Eukaryotic Cell.*, 4(10), 1677–1686.

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