

Adhesins and Ligands Involved in the Interaction of *Candida* spp. with Epithelial and Endothelial Surfaces

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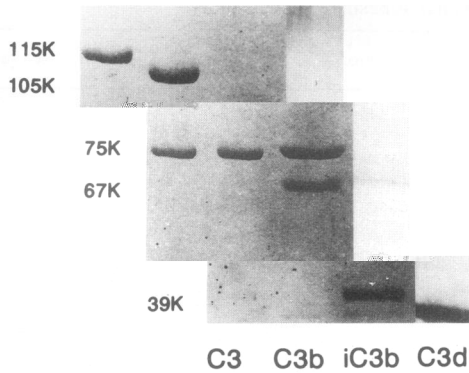
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appear to vary among different types of epithelial cells and

TABLE 1. Epithelial and endothelial adhesins

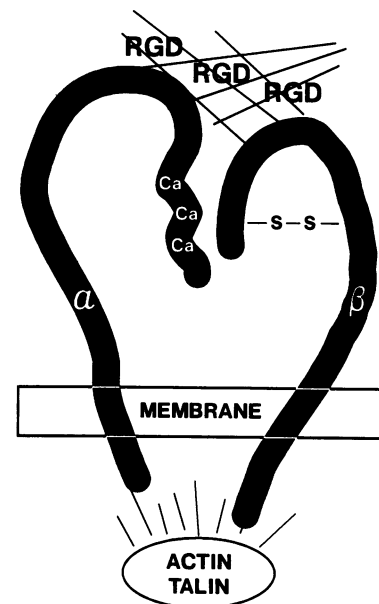
Adhesin	<i>M_r</i> (10 ³)	Epithelial ligand	Inhibitor(s)	Gene	Reference(s)
Epithelial					
Protein-protein					
Integrin analog (iC3b receptor)	130-165	iC3b	MAb, iC3b, RGD peptides	No	5, 27, 29, 37, 39, 45,
					49, 82
Fibronectin	60-68	Fibronectin	Fibronectin, proteases	No	100
Lectin-like					
Fucose-binding protein	ND ^a	Fucose	ND	No	22, 23, 25, 70, 72, 105



COMPLEMENT RECEPTORS

 β_2 INTEGRINS

Human Cells	Receptor	C3 Ligand	Mammal
Neutrophils, macrophages, monocytes, lymphocytes, glomerular mesangial cells, renal podocytes	CR1	C3b, C3d	PMNs, monocytes, RBCs
B lymphocytes	CR2	C3d	B lymphocytes
Neutrophils, macrophages, monocytes, NK cells	CR3	iC3b	PMNs, monocytes
Neutrophils, macrophages, monocytes, NK cells	CR4	iC3b	PMNs, monocytes

FIG. 3. Diagrammatic representation of α - and β -chains of the mammalian integrins. The amino termini are extracellular, the

transmembrane domains are represented by the shaded areas, and

the cytoplasmic tails are located at the carboxy termini, where they

TABLE 2. Binding of MAbs to *C. albicans*^a

Specificity	MAb	Isotype	with <i>C. albicans</i>	recognized (kDa)	EA rosette	Adhesion
α_M	OKM1	IgG2a	++++	42, 130, 165	Yes	No
	Anti-Mo1	IgM	++++	165	No	No
	MAb 17	IgM	+++	ND	ND	Yes
	MAb 44	IgG2a	+++	ND	ND	Yes
	Mn41	IgG1	+++	ND	ND	ND
	OKM10	IgG2a	++	ND	ND	ND
	M1/70	IgG2a	++	ND	ND	ND
	Mac-1	IgG2b	ND	ND	Yes	ND
α_X	BU-15	IgG1	++++	165 \pm 15	ND	ND
β_2	TS1/18	IgG	0	ND	ND	ND
	MHM23		0	ND	ND	ND
CR2	HB5	IgG2a	0	ND	Yes	ND
	Anti-gp140	Polyclonal	0	ND	Yes	ND
	Anti-B2	IgM	0	ND	No	ND

of binding of MAbs were not included. In ligand binding assays, OKM1, as can be seen from Fig. 4, there is a

assays, the 42-kDa protein bound C3(H₂O), C3b, and iC3b. immunofluorescence on yeast cells, germ tubes, and pseudo-

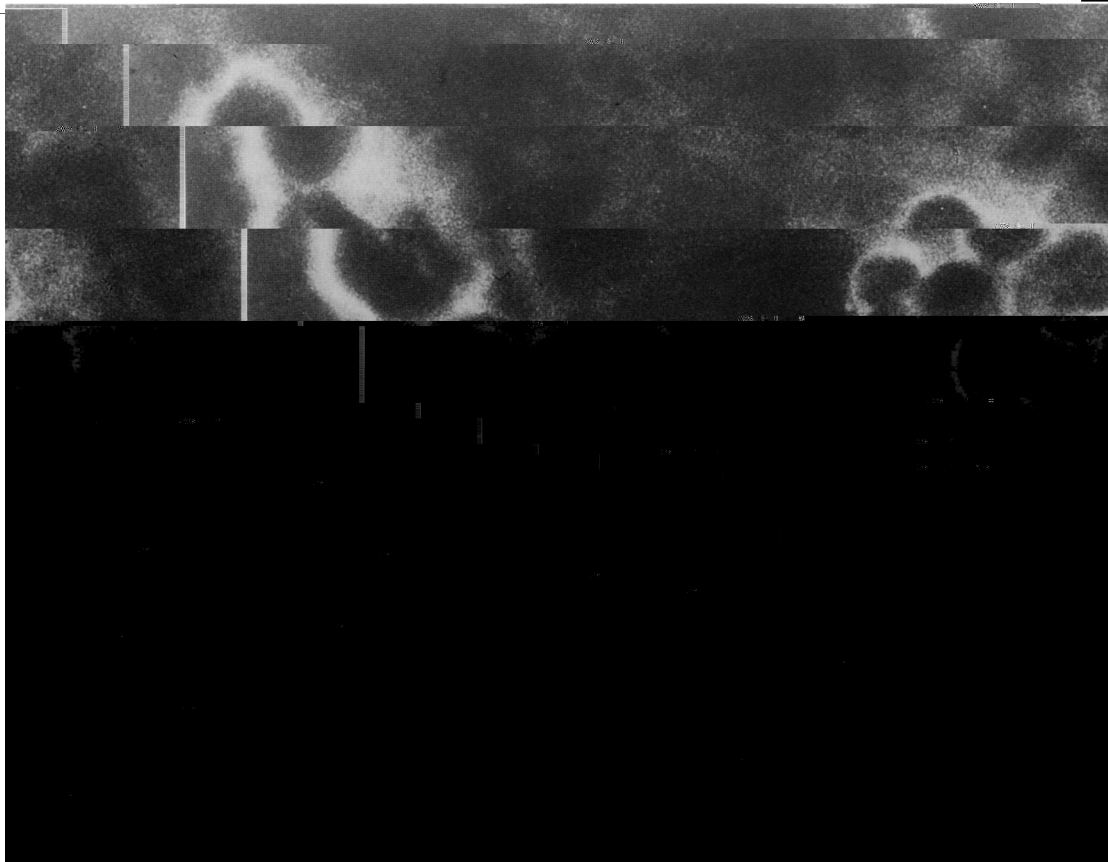


FIG. 4. Photomicrograph of *C. albicans* removed from the peritoneum of fungemic mice and incubated with MAb OKM1. There is visible immunofluorescence surrounding both blastospores and germ tubes (52).

O express the H antigen, a glycoside containing D-galactose, proteinase activity and from the employment of assays that

N-acetyl-D-glucosamine, *N*-acetyl-D-galactosamine, and an distinguish adhesion from invasion.

of the extracellular matrix, including type I and IV collagen, teins (62) but had no effect on hepatic uptake (96) empha-

fibronectin, and laminin. Interestingly, although fibronectin sizes again the tissue specificity of adhesion and the possi-

easily inhibited candidal binding to all four substrates, RGD bility that sequences surrounding the RGD site could

peptides did not reproducibly inhibit binding to collagen or regulate binding specificity.

antiserum against this protein has been used to stain kid- genesis, has just begun. Diligent investigation will provide

neys, cutis, or urethral epithelium and demonstrates the new insights into these mechanisms and ultimately improve

presence of the protein in both blastospores and pseudohy- treatment of candidal infections.

phal forms in vivo. However, the antibody did not signifi-

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albicans adherence to human vaginal epithelial cells in vitro. ties analogous to bovine conglutinin and functions as a recep-

Infect. Immun. 41:1024-1030. tor for zymosan and rabbit erythrocytes as well as a receptor

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the C3bi receptor of human monocytes and macrophages by