Murine Laminin Binds to *Histoplasma capsulatum*
A Possible Mechanism of Dissemination

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Abstract

Histoplasmosis, an increasingly important opportunistic infection in immunosuppressed subjects, is characterized by hematogenous dissemination of the yeast from the lung. The mechanism of this dissemination is not fully understood. Laminin, the major glycoprotein of the extracellular matrix, is known to mediate the attachment of various invasive pathogens to host tissues. In the current study, laminin is demonstrated to bind to *Histoplasma capsulatum* in a rapid, specific, and saturable manner. Scatchard analysis with ¹²⁵I-labeled laminin revealed an estimated 3.0 × 10⁴ binding sites per yeast with an apparent Kᵣ for laminin binding of 1.6 × 10⁻⁹ M. Laminin binding to *H. capsulatum* was decreased from 62.⁵ ± 1.⁷ ng (P < 0.001) in the presence of 3,000 nM of Ile-Lys-Val-Ala-Val, a pentapeptide within one major cell attachment site of laminin. A 50-kD *H. capsulatum* laminin-binding protein was demonstrated using an ¹²⁵I-Ln blot of *H. capsulatum* cell wall proteins. The 50-kD protein is also recognized by antibodies directed at the 67-kD laminin receptor, suggesting they are related. This study proposes a possible mechanism for *H. capsulatum* attachment to laminin, an important first step required for the yeast to recognize and traverse the basement membrane. *(J. Clin. Invest. 1995. 96:1010–1017.)*

Key words: attachment sites · yeasts · extracellular matrix · glycoproteins · immunosuppression

Introduction

*Histoplasma capsulatum*, a dimorphic fungal pathogen with worldwide distribution, is endemic to the United States. Disseminated histoplasmosis has emerged as an important opportunistic infection in immunosuppressed subjects, especially those with AIDS. Infection is acquired by inhaling airborne microconidia that elude pulmonary defense mechanisms and lodge in the alveoli and interstitium *(1).* Transformation of the infectious microconidia to the pathogenic yeast is an absolute requirement for progression to histoplasmosis *(2).*

The initial events following microconidia deposition in human alveoli are unknown. Experimental evidence with cultured human alveolar macrophages suggest that the microconidia are rapidly ingested by alveolar macrophages *(3).* Experimental infections in animals demonstrated transformation of the microconidia phase to the yeast phase, which proliferate within the parasitized macrophages. The host macrophage ruptures and the released yeast parasitize other macrophages, producing a local exudate composed of infected macrophages and lymphocytes *(1).* Evidence from a human lung biopsy specimen demonstrated that extracellular yeast forms existed within the alveolar exudate and throughout the pulmonary interstitium *(4).* Current concepts of histoplasmosis in the normal host suggest there is spread of intracellular yeast to adjacent alveoli, to regional lymph nodes, and to distant metastatic sites. The mechanism of metastasis to distant sites is based on indirect evidence. The process of tissue invasion and dissemination is virtually unstudied, particularly during severe immune suppression, when extracellular *H. capsulatum* organisms may be involved in dissemination and pathogenesis of the disease.

The lung contains an alveolar basement membrane that lies between type I alveolar epithelial cells and capillary endothelial cells as well as an endothelial basement membrane that separates endothelial cells from interstitial connective tissue cells along the thick portion of the alveolar wall *(5).* Laminin, an 850-kD glycoprotein, is the most abundant glycoprotein of the basement membrane and is composed of three polypeptide chains designated A, B₁, B₂, each containing six domains *(6, 7).* The multiple domains confer upon laminin many biological properties, including promoting cellular adhesion, growth, differentiation, migration, and matrix assembly. Laminin receptors have been demonstrated on metastatic tumor cells, granulocytes, lymphocytes, and macrophages *(8).* Recent studies have demonstrated that the invasive pathogenic organisms *Candida albicans* *(8, 9), Aspergillus fumigatus* *(10), Paracoccidioides brasiliensis* *(11),* staphylococci *(12–14), Helioacter pylori* *(15, 16), Treponema pallidum* *(17), streptococci* *(18–20), Toxoplasma gondii* *(21),* and *Escherichia coli* *(22, 23)* possess surface proteins with strong laminin-binding activity. The mechanism of laminin binding to the microorganism’s laminin receptor has only recently been investigated in *A. fumigatus,* enterobacteria, and *H. pylori* *(10, 24, 25).*

In this study, laminin is demonstrated to bind *H. capsulatum* in a rapid, specific, and saturable fashion. The major cell-binding region of the laminin A chain appears to be the molecular site of laminin binding to *H. capsulatum.* ¹²⁵I-labeled laminin *(¹²⁵I-Ln)₃* blotting with size-fractionated *H. capsulatum* cell wall proteins demonstrates a 50-kD protein that binds laminin.

1. Abbreviations used in this paper: ⁵²Cr- *H. capsulatum,* ⁵¹Cr-labeled *H. capsulatum*; ECM, extracellular matrix; ¹²⁵I-Ln, ¹²⁵I-labeled laminin; IKVAV, isoleucine-lysine-valine-alanine-valine; YIGSR, tyrosine-isoleucine-glycine-serine-arginine.
Further, antibodies directed at a known 67-kD laminin receptor recognize the 50-kD protein on *H. capsulatum*. These studies provide a biochemical mechanism for the interaction of *H. capsulatum* with a key basement membrane protein and suggest a specific protein on the surface of *H. capsulatum* as the site of laminin binding.

**Methods**

*H. capsulatum* isolation. The yeast phase of *H. capsulatum* strain G217b was maintained using brain–heart infusion agar (Difco Laboratories Inc., Detroit, MI) supplemented with 0.1% cysteine (Sigma Chemical Co., St. Louis, MO). Yeasts were inoculated onto brain–heart infusion agar slants, incubated for 24 h at 37°C, streaked over the remainder of the slant, and allowed to grow for 3–4 days. Yeasts were then harvested from the slant, washed four times in PBS and centrifuged at 1,200 g. *H. capsulatum* were counted using a hemacytometer and adjusted to proper concentrations. A fresh preparation of *H. capsulatum* was used for each experiment.

In vivo pulmonary *H. capsulatum* infection. In selected studies, *H. capsulatum* organisms (10^7/100 μl) were tracheally inoculated into CD4 lymphocyte-depleted mice using a minor adaptation of a previously described method (26). Mice were sacrificed 10 days after using Beuthanasia-d (0.1 ml, injected intraperitoneally) and exsanguination. The thoracic cavity was opened, the lungs were removed, and impression smears were made from the cut surface of the lung tissue and stained with Wright Giemsa stain. The lung tissue was paraffin embedded, sectioned (6 μm), and stained with hematoxylin-eosin.

In vitro *H. capsulatum* attachment assay. THP-1 *H. capsulatum* were incubated in 2 ml of RPMI 1640 (Biofluid Corp., Rockville, MD) supplemented with glutamine (0.6 μg/ml), penicillin (100 U/ml), streptomycin (0.5 μg/ml), and 300 μCi 35S-sodium chloride (specific activity 607.7 μCi/mg; New England Nuclear, Boston, MA) for 90 min at 37°C. After incubation, the 35S-labeled *H. capsulatum* (35S-Cr–*H. capsulatum*) suspension was centrifuged (1,200 g for 10 min), the supernatant was saved, and the pellet was resuspended in RPMI 1640. The *H. capsulatum* suspension was washed four times in RPMI 1640 to remove unincorporated 35S-Cr. The pellet was resuspended in RPMI 1640 at a concentration of 1 × 10^6/ml.

*H. capsulatum* attachment to immobilized basement-membrane proteins was performed using laminin or Matrigel® (Becton Dickinson, Bedford, MA) coated wells (Costar Corp., Cambridge, MA). Laminin-coated wells were prepared according to the manufacturer’s recommendation. Briefly, 50 μl of Matrigel® was added to square centimeter of well surface area and left to adhere at 37°C for 30 min. The wells were washed with PBS until the desired concentration was reached. The wells were washed three times with PBS. The attached 35S-Cr–*H. capsulatum* were disrupted with 10% Triton X-100 (Sigma Chemical Co.) and the wells washed with PBS. 35S-Cr–*H. capsulatum* was quantified in each fraction (5500 gamma counter; Beckman Instruments, Inc., Fullerton, CA) with the percentage of attachment expressed as follows: percentage of attachment = (A/ (A + B)) × 100, where A = 35S-Cr–*H. capsulatum* bound to the ECM and B = 35S-Cr–*H. capsulatum* free in the media.

Radioiodination of laminin. Laminin was labeled with 1 μCi of 125I (specific activity 17.4 mCi/μg; Amersham, Arlington Heights, IL) for 15 min at room temperature using Iodo-Beads (Pierce, Rockford, IL) (27). 125I-Ln was separated from free 125I by gel filtration on a Sephadex G-25 M PD-10 column (Pharmacia, Uppsala, Sweden) previously washed with equilibration buffer (0.05 M Tris Cl, 0.15 M NaCl, 0.02% sodium azide, pH 7.4) and saturated with 2 ml of 1% BSA in equilibration buffer (8). The specific activity of the 125I-Ln was 8.7 μCi/μg.

Production of polyclonal antibody to the *H. capsulatum* 50-kD protein. Polyclonal antibodies to the 50-kD *H. capsulatum* protein were developed according to previously described protocols (28) with a few modifications. Briefly, *H. capsulatum* cell wall proteins were separated by SDS-PAGE as outlined later. The gel was rinsed with ddH2O and stained for 15 min with ice-cold 0.25 M KCl and 1 mM DTT. The 50-kD band was identified and separated from the gel and destained for 10 min with cold ddH2O containing 1 mM DTT. The gel band was homogenized with a minimal amount of resuspended, and four volumes of cold acetone was added to the gel elute, and the samples were allowed to precipitate at −70°C. The samples were centrifuged at 1,200 g for 10 min, the acetone supernatants removed, and the protein was reconstituted in a minimal volume of 1 mM DTT. The presence of cell wall protein was confirmed by the presence of a single band on SDS-PAGE. 50–100 μg of the antigen was mixed with an equal volume of Hunter’s TiterMax adjuvant (Cryrix Corp., Norcross, GA) (1:1, vol/vol) and injected intraperitoneally into BALB/c mice. Mice were boosted at 4 weeks, and the primary injections were performed on days 1 and 14. The booster injection, without adjuvant, was performed on day 28. After the booster injection, the mice were exsanguinated, the serum was collected, and murine polyclonal antibodies to the 50-kD protein (PAB 50) were assayed. PAB 50 identified the 50-kD protein band in a Western blot of *H. capsulatum* cell wall proteins (Fig. 9, lane C).

Immunofluorescence. To confirm that the *H. capsulatum* 50-kD cell wall protein, identified in this study as the putative laminin-binding site, is a cell surface protein, the following immunofluorescence assay was employed. 2 × 10^7 yeast were suspended in 1 ml of attachment buffer (50 mM Hapes, 5 mM CaCl2, 0.1% BSA, pH 7.4) and incubated for 60 min at 37°C with murine polyclonal antibodies to the 50-kD cell wall protein at a dilution of 1:100. The yeast were then washed two times in 1 ml of attachment buffer, and four volumes of fluorescein-conjugated goat anti–mouse IgG (Calbiochem-Novabiochem Corp., La Jolla, CA) at a dilution of 1:40. The yeast were washed in attachment buffer, and cyto-preparation smears were prepared by centrifugation at 1,200 g (Cytospin 2; Shandon Inc., Pittsburgh, PA). The yeast were viewed using a Zeiss Axioplan microscope equipped with epifluorescence (Carl Zeiss Instruments Inc., Thornwood, NY). Control experiments were performed to substitute preimmune serum for PAB 50.

*H. capsulatum* binding to *H. capsulatum*. *H. capsulatum* was quantified using minor modifications of a previously described method (29). Reaction mixtures containing 2 × 10^6 *H. capsulatum*, 125I-Ln (concentration from 125 ng/ml to 3,000 ng/ml), and attachment buffer (50 mM Hapes, 5 mM CaCl2, 0.1% BSA, pH 7.4) in a volume of 1 ml were incubated and rotated at 37°C for 0.5–120 min. The mixture was centrifuged at 1,200 g for 5 min, the supernatant was removed and saved, and the pellet was resuspended in 1 ml of attachment buffer. After a second centrifugation, the supernatant was saved and added to the previous supernatant. The *H. capsulatum* pellet was resuspended in 1 ml of attachment buffer and was removed from the incubation tube and placed in a 12 × 75 mm tube for counting. The pellet with the bound 125I-Ln and the remaining supernatant in the incubation tube containing unbound 125I-Ln were quantified. Specific binding was determined by the addition of 40 μg of unlabeled laminin to the reaction mixture. Bound 125I-Ln was calculated as follows: Bound 125I-Ln = (CPMpellet/CPMpellet + supernatant) × 125I-Ln added.

There are high-affinity laminin-binding proteins which bind to oligosaccharide-binding sites within laminin. These sites include galactose and poly-N-acetylated-lactosamine oligosaccharides of laminin (30) as well as the oligomannoside chains of laminin (24). To determine if the carbohydrate structures of laminin were involved in the binding of laminin to *H. capsulatum*, the binding of 125I-Ln to *H. capsulatum* was conducted in the presence or absence of α-methyl-D-mannoside and N-
acetyl-lactosamine. Binding assays were performed using 2 × 10^7 H. capsulatum in attachment buffer, 250 ng ^125^I-Ln (0.3 nM), and increasing concentrations of α-methyl-mannopyranoside or N-acetyl-lactosamine (3–3,000 nM) (Sigma Chemical Co) in a 1-ml volume. After 30 min, the amount of bound ^125^I-Ln was calculated as earlier.

Major cell attachment sites on laminin include the peptide sequences Tyr-Ile-Gly-Ser-Arg (YIGSR) present on the B1 chain (31) and Ile-Lys-Val-Ala-Val (IKVAV) on the A chain (32). To determine if defined laminin cell–binding domains were involved in laminin binding to H. capsulatum, the binding of ^125^I-Ln to H. capsulatum was conducted in the presence or absence of the peptide sequences YIGSR and IKVAV. Binding assays were performed using 2 × 10^7 H. capsulatum in sample buffer, 250 ng ^125^I-Ln (0.3 nM), and increasing concentrations (750–3,000 nM) of amino acid sequence 925–933 of domain III of the B1 chain (includes YIGSR) and amino acid sequence 2,091–2,108 of domain I of the A chain (includes IKVAV) (Sigma Chemical Co.). The final volume of the mixture was 1 ml, which was incubated for 30 min, and the amount of bound ^125^I-Ln was calculated as discussed previously.

To confirm that the 30-kD H. capsulatum surface protein served as a receptor for laminin binding, the binding of ^125^I-Ln to H. capsulatum was performed in the presence or absence of PAb 50. Binding assays were performed using 2 × 10^7 H. capsulatum in sample buffer, 250 ng ^125^I-Ln, and increasing concentrations of PAb 50 (1:500, 1:50) in a 1-ml volume. After 30 min, the amount of bound ^125^I-Ln was calculated as earlier.

**Binding of ^125^I-Ln to H. capsulatum cell wall proteins.** A cell wall preparation of H. capsulatum was prepared as previously described with minor modifications (33). The yeasts were harvested from BHI broth (Difco Laboratories, Detroit, MI), washed in PBS, and centrifuged at 1,200 g for 10 min. The yeasts were resuspended in PBS and were disrupted with glass beads in a homogenizer (B. Braun Biotech Intl. GmbH, Melsungen, Germany) at 4°C using 30-s alternating cycles of homogenizing and cooling for a total of 4–6 min. Disruption of cells was verified by light microscopy. The homogenate was centrifuged at 20,000 g for 20 min (Sorvall RC 28S; DuPont, Wilmington, DE), and the cell-free supernatant was concentrated in PBS to a concentration of 2 mg/ml as determined by bichinchoninic acid protein assay (Pierce).

To separate cell wall proteins, the cell wall preparation was solubilized in SDS-PAGE sample buffer (0.375 M Tris, 20% glycerol, 4% SDS, 0.002% bromophenol blue, and 4% 2-mercaptopethanol) and placed on a discontinuous 7.5–20% gradient sodium dodecyl sulfate polyacrylamide gel. The gel was electrophoresed at 20 mA for 4 h and stained with Coomassie blue. Alternatively, the proteins on the gel were transferred to polyvinyl difluoride (PVDF) transfer membrane (Immobilon-P; Millipore Corp., Bedford, MA) in transfer buffer at 200 mA for 2 h (34). The PVDF membrane was incubated with blocking solution (5% dried milk in 150 mM NaCl, 50 mM Tris-Cl, pH 7.6, Tris-buffered saline [TBS]) for 1 h, washed with TBS-T (TBS-T) three times and then blotted with 1 ml ^125^I-Ln in TBS-T for 90 min. The membrane was washed in TBS-T and air dried. Autoradiography was performed at −70°C for 24 h with Kodak XAR-5 film (Eastman Kodak Co., Rochester, NY).

Western blot with antibodies to laminin receptor proteins. The H. capsulatum cell wall preparation was separated by SDS-PAGE (10% gel) and transferred to a PVDF membrane as described earlier. The PVDF membrane was incubated with blocking solution for 1 h, washed with TBS-T three times, and blotted with PAb 50 (1:1,000) and PAb 4099 (1:100) in TBS-T for 1 h. The membranes were washed and then incubated with anti-murine (PAb 50) and anti-rabbit (PAb 4099) alkaline phosphatase conjugate (BioRad, Hercules, CA) for 1 h in TBS-T. The membranes were washed and developed with a buffer (AP Conjugate Buffer Kit; BioRad), according to the manufacturer's specifications.

PAb 50 was described earlier; PAb 4099 recognizes the 67-kD laminin receptor of murine and human endothelial cells and has been described previously (35–37). PAb 4099 is affinity purified and diluted into PBS containing carrier ovalbumin. Its optimal dilution was empirically established to be specific and sufficiently sensitive at 1:100.

Statistical analysis. All results are expressed as mean±SEM. Statistical comparisons were performed using Student's t test for unpaired data or the one-way analysis of variance with paired comparisons performed using Fisher's least significant difference method. Statistical significance was accepted for P < 0.05 (38).

**Results**

An exuberant pulmonary infection developed in the CD4 lymphocyte-depleted mice (Fig. 1). Free H. capsulatum organisms were easily demonstrated in the lung tissue and in lung impression smears of the infected mice. In profoundly immunosuppressed animals, H. capsulatum pulmonary infection develops rapidly, with evidence for both intracellular and extracellular organisms present in the lung tissue.

^51^Cr–H. capsulatum attached to immobilized laminin and the basement membrane preparation Matrigel® (Fig. 2). The percentage of attachment of ^51^Cr–H. capsulatum was in direct proportion to the amount of immobilized laminin adherent to the well (data not shown). The percentage of attachment of
Figure 2. \(^{51}Cr-H.\) capsulatum attachment to extracellular matrix proteins. \(1 \times 10^8\) \(^{51}Cr-H.\) capsulatum were incubated with wells coated with laminin or the reconstituted basement-membrane preparation Matrigel\(^R\) for 30 min. The assay was performed in the presence or absence of anti-laminin IgG (1:5). The presence of anti-laminin IgG significantly reduced the percentage of attachment of \(^{51}Cr-H.\) capsulatum to laminin and Matrigel\(^R\) (59±4 to 21±3 and 67±2 to 39±5, respectively). Values are mean±SEM; \(n = 9.\)

\(^{51}Cr-H.\) capsulatum to laminin and Matrigel\(^R\) was significantly diminished in the presence of polyclonal anti-laminin IgG (59±4 to 21±3 and 67±2 to 39±5, respectively; \(P < 0.01,\) both comparisons). Matrigel\(^R\) is a solubilized basement-membrane preparation extracted from the Engelbreth-Holm-Swarm mouse sarcoma tumor. Its major component is laminin, followed by collagen IV, heparan sulfate proteoglycans, enactin, and nidogen.

Laminin binds \(H.\) capsulatum in vitro in a time-dependent manner and achieves apparent equilibrium by 10 min (Fig. 3). At 0.5 min, 21±2 ng \(^{125}I\)-Ln per \(2 \times 10^7\) yeast is bound, whereas beyond 10 min, binding of \(^{125}I\)-Ln reaches a plateau at approximately 73±4 ng per \(2 \times 10^7\) yeast.

The specificity and saturability of \(^{125}I\)-Ln binding to \(H.\) capsulatum were demonstrated using \(^{125}I\)-Ln (125–3,000 ng) in the presence or absence of 40 \(\mu\)g excess unlabeled laminin (Fig. 4 A). The addition of increasing amounts of \(^{125}I\)-Ln to the binding assay resulted in an increase in both total and nonspecific binding, with saturation of specific binding evident at concentrations in excess of 2,000 ng \(^{125}I\)-Ln added. The Scatchard plot using the specific binding data is linear (\(r = 0.88\)), indicating a probable homogenous population of binding sites for laminin on \(H.\) capsulatum organisms (Fig. 4 B). The binding dissociation constant \((K_d)\), obtained from the slope of the linear plot, is 1.6 \(\times 10^{-9}\) M. Each \(H.\) capsulatum yeast is estimated to have 3.0 \(\times 10^9\) binding sites for laminin, as calculated from the abscissa intercept.

To determine if the carbohydrate structures of laminin were involved in the binding of laminin to \(H.\) capsulatum, \(^{125}I\)-Ln binding to \(H.\) capsulatum was conducted in the presence or absence of \(\alpha\)-methyl-d-mannoside or \(\alpha\)-acetyl-lactosamine. The amount of \(^{125}I\)-Ln bound (ng) to \(H.\) capsulatum in the presence of 0 nM and 3,000 nM of \(\alpha\)-methyl-d-mannoside was 53±2 ng and 51±2 ng, respectively (\(P > 0.05\)); similarly, the binding in the presence of 0 nM and 3,000 nM of \(N\)-acetyl-lactosamine was 57±3 ng and 58±3 ng, respectively (\(P > 0.05\)). Therefore,
there was no evidence that laminin oligosaccharide side chains were involved in laminin binding to *H. capsulatum* (Fig. 5 A).

The major epithelial, neuronal, and malignant cell attachment sites of laminin have been localized to domain III of the B1 chain and domain I of the A chain which contain the peptide sequences YIGSR and IKVAV, respectively (31, 32). Therefore, to determine if these laminin cell-binding domains were involved in laminin binding to *H. capsulatum*, the binding of 125I-Ln to *H. capsulatum* was conducted in the presence or absence of YIGSR or IKVAV containing peptide sequences. The peptide sequence containing YIGSR had no significant effect on 125I-Ln binding to *H. capsulatum* (Fig. 5 B); whereas, the A chain peptide containing IKVAV demonstrated a concentration-dependent inhibition of 125I-Ln binding to *H. capsulatum* (Fig 5 B). As an example, laminin binding to *H. capsulatum* in the absence of IKVAV was 62±1 ng, compared with 17±1 ng in the presence of 3,000 nM of IKVAV (P < 0.001). This suggests that laminin binds to *H. capsulatum* via the major cell attachment site located within domain I of the laminin A chain.

The *H. capsulatum* cell wall preparation, separated by SDS-PAGE and stained with Coomassie blue, revealed numerous protein bands whose sizes ranged from 18 kDa to greater than 116 kDa (Fig. 6). The *H. capsulatum* laminin-binding protein was demonstrated following the transfer of *H. capsulatum* proteins to a PVDF membrane and the subsequent incubation of these proteins with 125I-Ln. Autoradiography of the 125I-Ln blot revealed that laminin binds to an *H. capsulatum* protein with an estimated mass of 50 kD. There is no visible band when the *H. capsulatum* cell wall proteins are blotted with 125I-Ln in the presence of excess unlabeled laminin, indicating that the band is the result of laminin specifically binding to the *H. capsulatum* cell wall protein.

Evidence that the 50-kD *H. capsulatum* protein is a cell surface protein is provided using immunofluorescence of *H. capsulatum* with PAb 50. 2 × 10^7 *H. capsulatum* were incubated with PAb 50 or preimmune serum (1:40), washed, and subsequently incubated with fluorescein-conjugated goat anti-mouse IgG (1:100). The immunofluorescent pattern (Fig. 7) by the yeast was prominent and was relatively similar for all yeast with respect to distribution and intensity. The yeast incubated with preimmune serum did not demonstrate immunofluorescence. These data indicate that the 50-kD protein is a cell surface protein.

To define further the role of the 50-kD protein as the *H. capsulatum* laminin-binding protein, the 125I-Ln binding studies were performed in the presence or absence of PAb 50. 2 × 10^7 yeast were incubated with 250 ng of 125I-Ln in the presence or absence of PAb 50. PAb 50 demonstrated a significant inhibition of 125I-Ln binding to *H. capsulatum* (Fig. 8). The amount of 125I-Ln bound to *H. capsulatum* was 93±9 ng and decreased to 40±5 ng (P < 0.05) in the presence of PAb 50 (1:50).

Western blot analysis of *H. capsulatum* cell wall proteins used PAb 4099, which recognizes the 67-kD laminin receptor. The *H. capsulatum* cell wall proteins were separated by SDS-PAGE, transferred to a PVDF membrane, and incubated with PAb 4099. This antibody recognized the 50-kD *H. capsulatum* cell wall protein (Fig. 9), indicating that the protein cross-reacts with an antibody that recognizes a known laminin receptor, suggesting that they are related.

**Discussion**

Histoplasmosis is characterized by early hematogenous dissemination of the yeast form from the initial site of infection. The mechanism of this dissemination is not completely understood.
Figure 7. Immunofluorescence pattern demonstrating the *H. capsulatum* 50-kD cell surface protein. 2 × 10^5 *H. capsulatum* were incubated with PAb 50 or with preimmune serum (1:100), followed by incubation with FITC-labeled goat anti-murine IgG (1:100). Phase and epifluorescence microscopy (A and B) of *H. capsulatum* incubated with PAb 50 demonstrate immunofluorescence, suggesting that the 50-kD protein is a surface protein. There was no immunofluorescence noted in the control experiment using preimmune serum (C and D).

and has been speculated to occur within blood-borne macrophages. The documentation of a patient with disseminated disease and interstitial and alveolar extracellular *H. capsulatum* yeast suggests that not all *H. capsulatum* are intracellular and, therefore, that alternative methods of dissemination are possible (4). In the current study, *H. capsulatum* yeast in CD4 lymphocyte-depleted mice exhibited exuberant growth, with both intracellular and extracellular organisms easily detected. Because dissemination during immunosuppression invariably occurs (39), these findings suggest that *H. capsulatum*, as an extracellular organism, may play a role in the pathogenesis of the disease in the immune-compromised host. Thus, the ability of *H. capsulatum* to interact with key constituents of the alveolar capillary barrier was examined.

*H. capsulatum* is capable of attaching to the basement-membrane preparation Matrigel®, and this attachment is significantly reduced by blocking laminin, the major glycoprotein of the basement membrane. This study demonstrates that laminin binds *H. capsulatum* in a rapid, specific, and saturable fashion. In addition, this study suggests that an *H. capsulatum* 50-kD surface protein is the laminin-binding protein and that antibodies to this 50-kD surface protein inhibit laminin binding to the yeast. Western blot analysis demonstrates the recognition of this surface receptor by 67-kD laminin receptor antibodies. The Scatchard plot estimates the number of laminin binding sites per *H. capsulatum* organism to be 3.0 × 10^7. Furthermore, the linearity of the plot suggests that a homogenous group of binding sites exists with a similar affinity for laminin.

Several classes of laminin-binding proteins exist, the function of which remains to be fully defined (40). However, the expression of laminin-binding proteins by pathogenic microorganisms has been demonstrated to enhance the organism's ability to attach to host cells and invade host tissues. These pathogenic microorganisms possess laminin-binding proteins which vary in molecular size from 32 (T. gonodii) to 145 kD (*Streptococcus gordonii*), with the majority of pathogens possessing laminin-binding proteins from 44 to 67 kD (8, 11, 14, 18, 21, 23). Invasive *S. aureus* has been demonstrated to possess a 52-kD surface protein which functions as a laminin-binding protein. Monoclonal antibodies to this 52-kD protein were demonstrated to bind to the 67-kD laminin receptor of murine melanoma cells, suggesting that conservation or convergence among laminin receptors exists (41). In this study, we demonstrate that the *H. capsulatum* 50-kD cell surface protein is involved in laminin binding and that this protein is recognized by antibody to the

Figure 8. The effect of polyclonal antibody to *H. capsulatum* 50-kD surface protein on ^125^I-Ln binding to *H. capsulatum*. 2 × 10^7 *H. capsulatum* were incubated with 250 ng of ^125^I-Ln in the presence of PAb 50 or murine IgG. PAb 50 demonstrated significant inhibition of ^125^I-Ln binding to *H. capsulatum*. Values are mean ± SEM; n = 9.

Figure 9. Western blot analysis of *H. capsulatum* cell wall proteins with PAb 4099. The cell wall proteins were separated by means of SDS-PAGE (10% gel), transferred to a PVDF membrane, and incubated with PAb 4099 (1:100), an antibody directed toward the 67-kD laminin receptor. Lane A demonstrates the molecular weight markers. Lane B is a Coomassie blue stain of the *H. capsulatum* cell wall proteins. Lane C demonstrates the 50-kD cell wall protein recognized by PAb 50. Lane D demonstrates the 50-kD cell wall protein recognized by PAb 4099. This suggests that the 50-kD cell protein of *H. capsulatum* shares epitopes with the known 67-kD laminin receptor. Lanes E and F are control lanes. In lane E, rabbit IgG is the primary antibody and in lane F there is no primary antibody, only the conjugate antiserum.
67-kD laminin receptor, lending further support to the concept of conservation or convergence among laminin receptors. Western blot analysis, autoradiography, and the linear Scatchard plot all indicate a single laminin-binding protein, which suggests that a predominant receptor class on the surface of *H. capsulatum* is involved in the majority of laminin binding.

Laminin exists in vivo as a large multidomain glycoprotein of the extracellular matrix. It is composed of globular and rod like domains arranged in a cruciform shape and is well suited for mediating attachment between distant sites on cells and other components of the extracellular matrix (7, 40, 42). To date, the domain within laminin recognized by laminin-binding proteins has been described for only three invasive pathogens. The laminin-oligomannoside chains, terminal sialic acid residues, and the PI fragment are the target for laminin-binding proteins of type I enterobacterial fimbriae, *H. pylori* and *A. fumigatus*, respectively (10, 24, 25). In this study, we have demonstrated that laminin binding to *H. capsulatum* is localized to domain I of the A chain and that this binding is dependent upon the IKVAV peptide sequence. The peptide sequence IKVAV has been demonstrated to mimic some of the biological activity of the intact laminin molecule (32). The IKVAV sequence is located in the carboxyl region of the α-helical domain of the laminin A chain, which is one the regions involved in promoting cell adhesion, spreading, and migration. The A chain serves as a ligand for the 67-kD laminin receptor group, among others (40).

In summary, we present evidence that the glycoprotein laminin binds to *H. capsulatum* yeast in a rapid, specific, and satura-

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