Members of the Candida parapsilosis Complex and Candida albicans are Differentially Recognized by Human Peripheral Blood Mononuclear Cells

Eine Estrada-Mata1, María J. Navarro-Arias1, Luis A. Pérez-García1, Erika Mellado-Mojica2, Mercedes G. López2, Katalin Csonka3, Attila Gacser3 and Héctor M. Mora-Montes1*

1 Departamento de Biología, División de Ciencias Naturales y Exactas, Campus Guanajuato, Universidad de Guanajuato, Guanajuato, México, 2 Centro de Investigación y de Estudios Avanzados del Instituto Politécnico Nacional, Irapuato, México, 3 Department of Microbiology, University of Szeged, Szeged, Hungary

The systemic infections caused by members of the Candida parapsilosis complex are currently associated with high morbidity and mortality rates, and are considered as relevant as those caused by Candida albicans. Since the fungal cell wall is the first point of contact with the host cells, here we performed a comparison of this organelle in members of the C. parapsilosis complex, and its relevance during interaction with human peripheral blood mononuclear cells (PBMCs). We found that the wall of the C. parapsilosis complex members is similar in composition, but differs from C. albicans, with less mannan content and more β-glucan and porosity levels. Furthermore, lectin-based analysis showed increased chitin and β1,3-glucan exposure at the surface of C. parapsilosis sensu lato when compared to C. albicans. Yeast cells of members of the C. parapsilosis complex stimulated more cytokine production by human PBMCs than C. albicans cells; and this significantly changed upon removal of O-linked mannans, indicating this wall component plays a significant role in cytokine stimulation by C. parapsilosis sensu lato. When inner wall components were exposed on the wall surface, C. parapsilosis sensu stricto and C. metapsilosis, but not C. orthopsilosis, stimulated higher cytokine production. Moreover, we found a strong dependency on β1,3-glucan recognition for the members of the C. parapsilosis complex, but not for live C. albicans cells; whereas TLR4 was required for TNFα production by the three members of the complex, and stimulation of IL-6 by C. orthopsilosis. Mannose receptor had a significant role during TNFα and IL-1β stimulation by members of the complex. Finally, we demonstrated that purified N- and O-mannans from either C. parapsilosis sensu lato or C. albicans are capable to block the recognition of these pathogens by human PBMCs. Together, our results suggest that the innate immune recognition of the members of the C. parapsilosis complex is differential of that reported for C. albicans. In addition, we propose that purified cell wall mannans can be used as antagonist to block specific receptors on innate immune cells.

Keywords: Candida parapsilosis, Candida albicans, host-fungus interaction, cell wall, cytokine, mononuclear cells
INTRODUCTION

Fungal infections are currently a burden for most of the health systems worldwide, and among them, superficial and invasive candidiasis are of special interest, since the latter has a mortality rate higher than 45% in infected patients (Brown et al., 2012). *Candida albicans* is the most frequent causative agent of candidiasis, being responsible of about 50% of total invasive candidiasis, while other members of the *Candida* genus, named emerging species, contribute together to the rest of the reported cases (Trofa et al., 2008). *Candida parapsilosis sensu lato* is a species that is mostly found in neonate patients, and is a versatile yeast-like organism that, at difference of other pathogenic *Candida* species, can be found colonizing non-human organisms and inert material from the environment (Trofa et al., 2008). This organism is in fact a complex of three closely related species: *C. parapsilosis sensu stricto*, *Candida orthopsilosis* and *Candida metapsilosis* (Tavanti et al., 2005), which have subtle, but key differences in terms of virulence (Nemeth et al., 2013; Gago et al., 2014), drug sensitivity (Spreghini et al., 2012; Szenzenstein et al., 2013), and secretion of hydrolytic enzymes (Trevino-Rangel Rde et al., 2013).

The establishment of a protective anti-*Candida* immune response in the host relays on a proper activation of the innate immune branch, and significant efforts have been done to understand this host–pathogen interaction, using *C. albicans* as a model (Netea et al., 2015). In *C. parapsilosis sensu lato*, there is currently limited information about the mechanisms underlying its interaction with components of the innate immunity. Thus far, it has been demonstrated that galectin 3 expressed by human neutrophils drives an increased *C. parapsilosis* phagocytosis, but not when challenged against *C. albicans* yeast cells (Linden et al., 2013). In addition, human peripheral blood mononuclear cells (PBMCs) stimulated with heat-killed (HK) *C. parapsilosis sensu stricto* yeast cells produced lower Interleukin (IL) 18, interferon γ, IL-17 and IL-22, but higher levels of IL-10, when compared to cells confronted with *C. albicans* (Toth et al., 2013). Despite this progress, there are not reports dealing with the interaction of immune cells with members of the *C. parapsilosis* complex.

The fungal cell wall contains most of the pathogen-associated molecular patterns recognized by pattern recognition receptors (PRRs) on innate immune cells, and again, the *C. albicans* cell wall is the best model thus far characterized (Diaz-Jiménez et al., 2012). This structure is composed of four main polysaccharides arranged in two well-defined layers: the outermost layer composed of glycoproteins, bearing N- and O-linked mannans, and the components of the inner layer, chitin, β1,6- and β1,3-glucans (Diaz-Jiménez et al., 2012). The participation of this organelle in the activation of the innate immune response has been thoroughly studied. It is now well established that β1,3-glucan is normally hidden from the recognition of dectin-1 and TLR2, and if accessible, plays a major role in the induction of pro-inflammatory cytokines and phagocytosis by macrophages (Ganttner et al., 2005; Gow et al., 2007; Heinsbroek et al., 2008). The N-linked mannans play also a significant role in both cytokine stimulation and macrophage uptake, via the mannose receptor (MR), dectin-2, and DC-SIGN (Netea et al., 2006; Mora-Montes et al., 2007, 2010; Cambi et al., 2008; McKenzie et al., 2010; Sajo and Iwakura, 2011). On the contrary, O-linked mannans play a dispensable role in both immunological processes, although it engages to TLR4 (Netea et al., 2006), a potent PRRs that plays a significant role in controlling bacteria (Miyake, 2007). Thus far, there is limited information about the cell wall of members of the *C. parapsilosis* complex, and the particular contribution of PRRs in the activation of cytokine production. Here, we performed a comparative study of the cell wall composition of *C. albicans*, *C. parapsilosis sensu stricto*, *C. orthopsilosis*, and *C. metapsilosis* and found that, although the composition is similar, the arrangement of the components has significant differences that impact their ability to activate human PBMCs. Moreover, we demonstrated that purified N- and O-linked mannans from either *C. albicans*, *C. parapsilosis sensu stricto*, *C. orthopsilosis* or *C. metapsilosis* are capable to block the recognitions of these pathogens by human PBMCs.

MATERIALS AND METHODS

Strains and Culturing Conditions

*Candida albicans* SC5314 (Gillum et al., 1984), *C. parapsilosis sensu stricto* SZMC 8110, *C. orthopsilosis* SZMC 1545, and *C. metapsilosis* SZMC 1548 (Szenzenstein et al., 2013) were used in this study. Cells were propagated at 30°C in Sabouraud broth [1% (w/v) mycological peptone, 4% (w/v) glucose], and maintained in plates containing medium added with 2% (w/v) agar. For all the experiments here reported, 500 μL of overnight-grown cells were used to inoculated 100 mL of fresh medium and incubated at 30°C with shaking at 200 rpm, until reach the mid-log growth phase (typically 5–6 h). Cells were incubated at 56°C for 1 h for heat inactivation as described (Mora-Montes et al., 2007). For all the cases, inactivation was confirmed by loss of fungal growth in Sabouraud medium at 30°C for 72 h. To remove O-linked mannans, cells were incubated overnight with 100 mM NaOH as described (Diaz-Jimenez et al., 2012). Under these conditions, more than 94% cells kept viability, as tested by CFU/mL before and after treatment with NaOH.

Cell Wall Analysis

Cell homogenates were obtained in a Braun homogenizer, with 5 cycles of 30 s and cooling on ice in-between (Mora-Montes et al., 2007). The cell walls were pelleted by centrifugation and thoroughly washed with deionized water. Further wall cleansing was performed with hot 2% (v/v) SDS, 0.3 M β-mercaptoethanol, and 1 M NaCl as described (Mora-Montes et al., 2007). Freeze-dried cell walls were hydrolyzed by adding 2 M trifluoroacetic acid and boiling for 3 h; then, the acid was evaporated, and samples were suspended in deionized water. The hydrolysates were analyzed by High Performance Anion Exchange Chromatography coupled to Pulsed Amperometric Detection (HPAEC-PAD), using a gradient of sodium acetate in 150 mM NaOH (flow 0.5 mL/min) as follows: 0–5 min = 45–75 mM NaOH, 5.1–15.0 min = 90 mM NaOH,
When required, PBMCs were pre-incubated for 60 min at 37 °C. Applied potentials for detection by the amperometric pulse were: E1 (400 ms), E2 (20 ms), E3 (20 ms), and E4 (60 ms) of +0.1, −2.0, +0.6, and −0.1 V, respectively. Protein content was determined upon lyophilized cell walls were alkali-hydrolyzed (Mora-Montes et al., 2007), using the Bradford protein assay.

The cell wall porosity was assessed using the polycation method previously reported (De Nobel et al., 1990). Aliquots containing 1 × 10⁸ cells were pelleted, the supernatant discarded and cells suspended in either 10 mM Tris-HCl, pH 7.4 (buffer A), buffer A plus 30 µg/mL poly-L-lysine (MW 30–70 kDa, Sigma Cat. No. P-2636) or buffer A plus 30 µg/mL DEAE-dextran (MW 500 kDa, Sigma Cat. No. D-9885). Then, cells were incubated at 30 °C for 30 min, and shaking (200 rpm), pelleted by centrifuging, the supernatant saved, and further centrifuged before reading the absorbance at 260 nm. The relative cell wall porosity to DEAE-dextran was calculated as described (De Nobel et al., 1990).

The level of cell wall phosphomannan was determined by the ability of cells to bind the cationic dye Alcian Blue as described (Hobson et al., 2004).

Analysis of Chitin and β1,3-Glucan Exposure at the Cell Surface

Cells were labeled with 1 mg/mL fluorescein isothiocyanate-wheat germ agglutinin conjugate (WGA-FITC; Sigma) (Mora-Montes et al., 2011) for chitin staining; while β1,3-glucan was labeled with 5 µg/mL IgG Fc-Dectin-1 chimera (Graham et al., 2006) for 40 min at room temperature, followed by incubating with 1 µg/mL donkey anti IgG Fc-FITC for 40 min at room temperature (Marakalala et al., 2013). In both cases, samples were examined by fluorescence microscopy using a Zeiss AxioScope-40 microscope and an Axiocam MRc camera. Pixels associated to 100 fluorescent cells were obtained with Adobe Photoshop™ CS6 and the following formula: [(total of green pixels-background green pixels) × 100]/total pixels.

Ethics Statement

Universidad de Guanajuato, though the Ethics Committee, approved the use of human cells in this study (permission number 17082011). Blood samples from healthy adult volunteers were obtained upon information about the study was provided and written informed consent was signed.

Human PBMCs-Candida Interaction

Human PBMCs were isolated by density centrifugation using Histopaque-1077 (Sigma) as reported (Endres et al., 1988). The immune cell-fungus interaction was performed in 96-well microplates with 5 × 10⁵ PBMCs, in 100-µL RPMI 1640 Dutch modification (Sigma), and 100 µL with 1 × 10⁵ fungal cells. When required, PBMCs were pre-incubated for 60 min at 37°C with either 200 µg/mL purified mannan, laminarin (200 µg/mL), anti-MR (10 µg/mL, Invitrogen, Cat. No. Mab-Hmr) or anti-TLR4 (10 µg/mL, Santa Cruz Biotechnology, Cat. No. sc-293072) prior to stimulation with yeast cells. Isotype matched, irrelevant IgG₁ antibodies (10 µg/mL, Santa Cruz Biotechnology, Cat. No. sc-52003) were used as controls for experiments assessing MR and TLR4. Despite all the reagents used for the pre-incubation experiments were negative to contamination by LPS (tested with the Limulus amebocyte lysate from Sigma) all reactions were performed in presence of 5 µg/mL polymyxin B (Sigma) (Schwartz et al., 1972). In all cases, the interactions were incubated for 24 h at 37°C with 5% (v/v) CO₂, plates were centrifuged for 10 min at 3000 × g and 4°C and supernatants saved and kept at −20°C until used. ELISA kits from Peprotech were used to measure the concentration of TNFα, IL-6, and IL-10; while IL-1β levels were measured using an ELISA kit from R&D Systems. Mock interactions with only human PBMCs incubated with RPMI 1640 Dutch modification were included as negative controls. For all the cases, the amount of cytokine quantified in the negative controls was subtracted before data analysis.

N- and O-Linked Mannan Isolation

The N- and O-linked mannans were isolated, and purified using the endoglycosidase H and β-elimination strategy previously reported by our group (Mora-Montes et al., 2012). Absence of protein and other wall polysaccharides was confirmed by the Bradford method and HPAEC-PAD, respectively (Mora-Montes et al., 2012); while lack of bacterial and fungal contamination was assessed by negative detection of LPS and no growth in LB and Sabouraud broth for 72 h at 37 and 28°C, respectively.

Statistical Analysis

Statistical analysis was performed using GraphPad Prism 6 software. Cytokine stimulation using human PMBCs was performed in duplicate with six healthy donors, whereas the rest of the experiments were performed at least thrice in duplicate. Data represent cumulative results of all experiments performed and are showed as mean ± SD. The Mann–Whitney U test or unpaired t-test was used to establish statistical significance (see figure legends for details), with a significance level set at P < 0.05.

RESULTS

Members of the C. parapsilosis Complex have Similar Composition in the Cell Wall but Differ from that Present in C. albicans

In order to assess the proportions of the main cell wall components in the three members of the C. parapsilosis complex, cell walls where isolated, acid-hydrolyzed and the proportion of N-acetylglucosamine, mannose and glucose, the basic unit of chitin, mannan and β-glucan, respectively (Mora-Montes et al., 2007, 2010, 2012), were quantified by HPAEC-PAD. C. parapsilosis sensu stricto, C. orthopsilosis and C. metapsilosis showed similar levels of the three polysaccharides analyzed (Table 1), but when compared with C. albicans, the species belonging to the complex showed significantly less mannan and increased β-glucan content (Table 1). In terms of chitin content, only C. metapsilosis wall displayed higher content than
the **C. albicans** cell wall (Table 1). No differences were observed for protein content in the four yeast species analyzed (Table 1). Next, in order to confirm the lower mannan content in the members of the **C. parapsilosis** complex, we assessed the level of phosphomannan content and the wall porosity, parameters that suffer alterations depending on the status of the protein mannosylation pathways (De Nobel et al., 1990; Bates et al., 2005, 2006; Mora-Montes et al., 2007, 2010; West et al., 2013; Lopes-Bezerra et al., 2015). Interestingly, the cell wall from **C. metapsilosis** showed lower phosphomannan content and higher porosity, when compared with **C. albicans** cells (Table 1). To assess whether the β1,3-glucan and chitin fibers in the members of the complex are underneath the mannan layer, as reported in **C. albicans** (Gow et al., 2007; Mora-Montes et al., 2011), yeast cells were labeled with the IgG Fc-Dectin-1 chimera (Graham et al., 2006), and the lectin-β1,3-glucan interaction revealed with FITC-conjugated IgG; whereas chitin was labeled with WGA-FITC. The analysis of the fluorescence associated to 100 cells indicated that, **C. parapsilosis sensu stricto**, **C. orthopsilosis**, and **C. metapsilosis** have β1,3-glucan and chitin more exposed at the cell surface than **C. albicans** cells (Figure 1). However, when compared the three members of the complex, **C. parapsilosis** displayed significantly higher levels of both polysaccharides at the wall surface than the other two species (Figure 1). As control, a similar experiment was conducted with HK yeast cells, where the components of the inner wall layer were artificially exposed at the cell wall surface and as consequence, higher labeling with the lectins was observed, with exception of **C. orthopsilosis**, where HK and live cells had similar content of both polysaccharides at the surface (Figure 1). Overall, these data indicate that the cell wall composition and organization among members of the **C. parapsilosis** complex differ to that described in **C. albicans**.

**C. albicans and C. parapsilosis sensu lato Differentially Stimulate Cytokine Production by Human PBMCs**

The differences in composition and organization of the cell wall above described may suggest that the fungus-immune cell interaction differs between member of the **C. parapsilosis** complex and **C. albicans**. Thus, in order to get some insights about such interaction, yeast cells were co-incubated with human PBMCs and secreted cytokines levels were measured as a read out of such interaction. Live **C. albicans** cells barely stimulated the production of either TNFα, IL-1β, IL-6 or IL-10 (Figure 2); however, **C. parapsilosis sensu stricto**, **C. orthopsilosis**, and **C. metapsilosis** stimulated higher and similar levels of TNFα, IL-1β, and IL-10 (Figure 2). Although IL-6 stimulation was significantly higher during the interaction with any of the three members of the **C. parapsilosis** complex, **C. orthopsilosis** stimulated the highest IL-6 production, when compared with the other species analyzed (Figure 2). Next, we compared the cytokine profile stimulated with either live or HK cells with or without β-elimination treatment to remove O-linked mannans from the cell wall (Diaz-Jimenez et al., 2012). As previously reported for **C. albicans** (Gow et al., 2007), HK yeast cells stimulated significant levels of TNFα, IL-1β, IL-6, and IL-10, when compared to live cells (Figure 3). A subtle, not significant reduction was observed in the production of the cytokines upon stimulation of human PBMCs with β-eliminated HK **C. albicans** cells (Figure 3). Upon heat-inactivation, **C. parapsilosis sensu stricto** and **C. metapsilosis** stimulated increased levels of TNFα, IL-6 and IL-10, and these cytokine levels were similar when live yeast cells were O-deglycosylated, i.e., β-eliminated (Figure 3). No further changes in the cytokine stimulation were observed when this treatment was applied to HK cells (Figure 3). For stimulation of IL-1β, modest levels of this cytokine were obtained, being not as abundant as those stimulated with HK **C. albicans** cells, as reported (Toth et al., 2013), and displaying no significant changes even though cells were both HK and β-eliminated (Figure 3). For **C. orthopsilosis**, the heat killing and β-elimination did not affect production of TNFα, IL-1β, IL-6 and IL-10 by human PBMCs. Therefore, **C. parapsilosis sensu lato** is capable to stimulate a different cytokine production than that observed in **C. albicans**, when incubated with human PBMCs. Furthermore, these data indicate that **C. parapsilosis sensu stricto** and **C. metapsilosis** but not **C. orthopsilosis**, have a similar interaction with these immune cells.

**Role of PRRs in the Sensing of Members of the **C. parapsilosis** Complex**

Next, in order to assess the importance of some PRRs during the fungus-immune cell interaction, human PBMCs were pre-incubated with antagonist of specific PRRs before the interaction with fungal cells. To evaluate the relevance of dectin-1 during this interaction, human PBMCs were pre-incubated with the specific antagonist laminarin (Maneu et al., 2011; Huang et al., 2012; Cohen-Kedar et al., 2014). Results showed in Figure 4 indicate that stimulation of TNFα, IL-1β, IL-6, and IL-10

![Table 1: Cell wall analysis of Candida albicans and members of the C. parapsilosis complex.](image)

| Organism            | Chitin (%) | Mannan (%) | Glucan (%) | Phosphomannan content (μg g⁻¹) | Porosity (%) | Protein (μg g⁻¹) |
|---------------------|------------|------------|------------|--------------------------------|--------------|-----------------|
| C. albicans         | 2.6 ± 1.2  | 33.0 ± 5.0 | 65.0 ± 3.3 | 127.2 ± 9.4                    | 28.2 ± 8.7   | 125.7 ± 21.1    |
| C. parapsilosis s. str. | 5.2 ± 2.7  | 13.2 ± 6.8* | 81.6 ± 9.4* | 72.5 ± 7.5*                    | 60.9 ± 3.9*  | 143.6 ± 47.3    |
| C. orthopsilosis    | 4.9 ± 1.4  | 20.5 ± 1.3* | 74.6 ± 0.9* | 58.5 ± 12.6*                   | 57.2 ± 1.7*  | 107.0 ± 20.7    |
| C. metapsilosis     | 6.6 ± 1.3* | 21.7 ± 1.5* | 71.7 ± 2.7* | 85.4 ± 17.9*                   | 71.8 ± 13.1* | 97.3 ± 25.0     |

*μg of Alcian Blue bound/OD₆₀₀ = 1; bRelative to DEAE-Dextran; cμg of protein/mg of cell wall; * P < 0.05, when compared to **C. albicans**.
C. orthopsilosis PBMCs were co-incubated with IL-6 stimulation was significantly diminished only when human C. parapsilosis by the three members of the cytokine production (not shown). TNF HK cells did not show significant differences in the levels of the same isotype gave no significant differences in the protocols (Mora-Montes et al., 2012). Purified mannans from C. albicans have been previously used to block its proper recognition by human PBMCs (Mora-Montes et al., 2011), thus we aimed to isolate

by HK C. albicans cells was significantly dependent on the engagement of dectin-1 with β1,3-glucan, but not for live cells. When similar experiments were conducted with members of the C. parapsilosis complex, levels of TNFα, IL-6, and IL-10 stimulated with the three species were significantly reduced upon pre-incubation with laminarin, indicating a strong dependence on this receptor for their stimulation (Figure 4). At difference of C. albicans, stimulation of cytokines by live C. parapsilosis sensu stricto cells was significantly sensitive to the presence of the dectin-1 antagonist. Interestingly, when human PBMCs were stimulated with C. parapsilosis sensu lato the IL-1β production was insensitive to the dectin-1 blocking, suggesting this receptor does not play a significant role for IL-1β stimulation by members of the C. parapsilosis complex.

To assess the relevance of TLR4 receptor during recognition of C. parapsilosis sensu lato, human PBMCs were pre-incubated with blocking anti-TLR4 antibodies before challenged with yeast cells. Results shown in Figure 5 indicate that blocking of TLR4 has no impact on the cytokine production stimulated by live C. albicans cells. Similar experiments conducted with HK cells did not show significant differences in the levels of cytokine production (not shown). TNFα production stimulated by the three members of the C. parapsilosis complex was significantly reduced when TLR4 was blocked (Figure 5); while IL-6 stimulation was significantly diminished only when human PBMCs were co-incubated with C. orthopsilosis (Figure 5). Experiments conducted in presence of an irrelevant antibody of the same isotype gave no significant differences in the cytokine level (Figure 5). No significant changes where observed when IL-10 and IL-1β were quantified after pre-incubation with the TLR4 blocking antibodies (Figure 5). Therefore, TLR4 is required for TNFα production but not for IL-10 nor IL-1β, when human PBMCs are challenged with C. parapsilosis sensu lato. Furthermore, production of IL-6 was specifically affected when TLR4 engagement was disrupted in cells stimulated with C. orthopsilosis.

When the yeast-PBMC interaction was performed with human cells pre-treated with anti-MR antibodies, there were no significant changes in the cytokine production stimulated by C. albicans cells (Figure 6). However, TNFα, and IL-1β stimulation by C. parapsilosis sensu lato was significantly reduced upon MR blocking (Figure 6). IL-6 levels were only significantly reduced in anti-MR-pre-incubated human PBMCs stimulated with either C. orthopsilosis or C. metapsilosis, but not with C. parapsilosis sensu stricto (Figure 6); while blocking of MR only affected IL-10 production in cells stimulated with C. orthopsilosis (Figure 6). Control interactions with an irrelevant antibody did not show changes in the cytokine production stimulated by all tested yeast cells (Figure 6). Taken together, these results indicate the role of dectin-1, TLR4 and MR in the recognition of members of the C. parapsilosis complex and C. albicans is different.

**Purified Mannans from C. albicans and C. parapsilosis sensu lato are Capable to Mimic the Effect of Anti-TLR4 and Anti-MR Antibodies on Human PMBCs**

Purified cell wall components from C. albicans have been previously used to block its proper recognition by human PBMCs (Mora-Montes et al., 2011), thus we aimed to isolate either N-linked or O-linked mannan using well-established protocols (Mora-Montes et al., 2012). Purified mannans from

![Figure 1](image1.png)

**FIGURE 1 | Analysis of β1,3-glucan and chitin exposure at the cell wall surface.** Live (L) yeast cells were incubated with IgG Fc-Dectin-1 and then with FITC-conjugated IgG to label β1,3-glucan (closed circles). Alternatively, chitin was labeled with WGA-FITC (open circles). The fluorescence associated to each cell is plotted. In addition, the experiment was conducted with cells pre-incubation with laminarin, indicating a strong dependence of the cell wall β1,3-glucan and chitin exposure at the cell wall surface.

![Figure 2](image2.png)

**FIGURE 2 | Candida albicans and members of the C. parapsilosis complex differentially stimulate the cytokine production by human PBMCs.** Human PBMCs were co-incubated 24 h with live yeast cells, the supernatant collected and used to quantify the cytokine levels. Strains used are C. albicans SC5314, C. parapsilosis sensu stricto SZMC 8110, C. orthopsilosis SZMC 1545, and C. metapsilosis SZMC 1548. *P < 0.05, when compared with the cytokine level stimulated by other species.
any of the three members of the C. parapsilosis complex or C. albicans were negative to fungal, bacteria and protein or other wall polysaccharide contamination, and did not stimulate any cytokine production when used at concentration up to 400 μg/mL (not shown). When human PBMCs were pre-incubated with 200 μg/mL N- or O-linked mannans from C. albicans and then stimulated with C. parapsilosis sensu stricto, TNFα stimulation was significantly reduced, but not IL-6 production (Table 2). Similar results were observed when mannans from either C. parapsilosis sensu lato, C. orthopsilosis or C. metapsilosis were used in the pre-incubation step (Table 2), or when mannans were used to block the proper recognition of C. orthopsilosis or C. metapsilosis (not shown). Therefore, N- and O-linked mannans are able to block the recognition of C. parapsilosis sensu lato as the anti-MR and anti TLR4 antibodies, respectively.

DISCUSSION

Members of the Candida genus are frequently regarded as similar to C. albicans; however, it is possible to demonstrate they have both genetic and metabolic differences that can affect the interaction with the host (Butler et al., 2009). Here, our comparative analysis of cell wall composition showed that members of the C. parapsilosis complex displayed lower mannan quantity, with similar protein level, when compared to C. albicans cells. In order to explain these data, we hypothesize that the mannans displayed on the surface have lower content of mannose per oligosaccharide unit, i.e., they are shorter in C. parapsilosis sensu lato. Indeed, it has been demonstrated C. parapsilosis has shorter N-linked mannans than those from the C. albicans cell wall (Shibata et al., 1995). The presence of short mannans is likely to affect the cell wall porosity, offering higher ability to DEAE-dextran, a bulky polycation, to reach the plasma membrane with the consequent increment in nucleic acid leakage (De Nobel et al., 1990). Moreover, these shorter mannans could be responsible of the increased content of both chitin and β1,3-glucan exposed at the wall surface. Our hypothesis of shorter mannans on the surface of C. parapsilosis sensu lato cells is also supported by the reduced phosphomannan content, which is an indirect measurement of both N- and O-linked mannan length (Bates et al., 2005, 2006; Mora-Montes et al., 2007, 2010; West et al., 2013; Lopes-Bezerra et al., 2015). This lower content of cell wall mannan seems to be naturally compensated by increasing the β-glucan content. This is a well-documented compensatory mechanism driven by the activation of the calcineurin and the protein kinase C signaling pathways, where disruption of the synthesis of one cell wall component usually affects the levels of other components to avoid wall weakness (Bates et al., 2005, 2006; Mora-Montes et al., 2007, 2010; West et al., 2013; Lopes-Bezerra et al., 2015). Our results also point out to subtle, but significant differences in the wall organization between members of the C. parapsilosis complex. In C. orthopsilosis all the β1,3-glucan and chitin is exposed in

FIGURE 3 | Cytokine production by human PBMCs stimulated with either live, heat-killed (HK) or β-eliminated yeast cells. Human PBMCs were co-incubated 24 h with either live, HK, live and β-eliminated, or HK and β-eliminated yeast cells, supernatant collected and used to quantify the cytokine levels. Strains used are C. albicans SC5314, C. parapsilosis sensu stricto SZMC 8110, C. orthopsilosis SZMC 1545, and C. metapsilosis SZMC 1548. *P < 0.05, when compared with live cells from the same species.
live cells, which suggest its cell wall proteins have even shorter mannans than *C. parapsilosis sensu stricto* and *C. metapsilosis*, but with more glycosylation sites occupied per protein. Alternatively, it could also be possible that mannans attached to *C. orthopsilosis* wall proteins are larger than those on the surface of other members of the complex, occupying less number of sites in the proteins, making the inner wall polysaccharides highly exposed at the cell surface. Nevertheless, further experiments are required to provide a conclusive explanation to these observations.

Results presented here clearly demonstrate that the human PMBCs-*C. parapsilosis sensu lato* interaction is different to that previously characterized in *C. albicans*. While yeast cells of *C. albicans* are a poor stimulus for cytokine production, the three members of the *C. parapsilosis* complex induced a strong dectin-1-dependent cytokine production. This is in line with the higher cell wall β1,3-glucan content in live *C. parapsilosis sensu lato*, but not in *C. albicans*. It is noteworthy to mention that results obtained here with *C. albicans* cells contrast with those previously reported (Netea et al., 2006), where strong cytokine production was stimulated by yeast cells of this species. This apparent discrepancy relies in the fact that here we used live yeast cells, while in the previous work the main conclusions were based on HK organisms (Netea et al., 2006). When cells were heat killed we observed and increment in the chitin and the β1,3-glucan content, as previously reported (Gantner et al., 2005; Gow et al., 2007; Mora-Montes et al., 2011), and this correlated with a significant increment in cytokine production stimulated by *C. albicans*, *C. parapsilosis sensu stricto* and *C. metapsilosis*, stressing the importance of the dectin-1-β1,3-glucan interaction for a strong cytokine induction. In the case of *C. orthopsilosis*, we did not notice a significant increment in cytokine production upon heat killing, which support our result indicating most of the β1,3-glucan is naturally exposed on the *C. orthopsilosis* cell surface. Our results also suggest that O-linked mannans from yeast cells play a major role in the recognition of *C. parapsilosis sensu lato* than in *C. albicans*, where they are dispensable and redundant wall elements for cytokine stimulation by human PMBCs (Netea et al., 2006). It has been reported though, that O-linked mannans from *C. albicans* hypha have a major participation in cytokine stimulation by epithelial cells (Murciano et al., 2011), suggesting that the proposed redundant role during interaction with components of the immune system may be restricted to PMBCs. Here, we have evidence indicating that in the case of *C. parapsilosis sensu lato*, β-eliminated and HK cells stimulated similar cytokine levels, which suggest that upon removal of O-linked mannans, inner wall components are exposed and available to engage with PRRs, i.e., O-linked mannans have a role hiding inner wall components from immune receptors. Thus far, it has been reported that N-linked mannans mask the
recognition of inner wall components and when disrupted, a significant cytokine production can be stimulated (Wheeler and Fink, 2006; Gow et al., 2007). Similar strategies, where a pro-inflammatory wall components is masked to avoid recognition by elements of the immune system, have been reported in *Histoplasma capsulatum* and *Aspergillus fumigatus*, where α-glucans and rodlet hydrophobins hide β1,3-glucan, respectively (Rappleye et al., 2007; Aimanianda et al., 2009). Therefore, it is feasible to conceive that in the case of *C. parapsilosis sensu lato*, both N- and O-linked mannans provide a disguising strategy to avoid recognition of β1,3-glucan by dectin-1. More experiments are required to demonstrate whether these structures are more abundant on the *C. parapsilosis sensu lato* cell wall or contain more mannose residues than those reported in *C. albicans*.

In this study we assessed the relevance of dectin-1, TLR4 and MR in the recognition of *C. parapsilosis sensu lato*. We chose these receptors because they are the main PRRs involved in the *C. albicans* immune recognition (Netea et al., 2002, 2006, 2015). TLR 4 is thus far the unique receptor for *C. albicans* O-linked mannans (Netea et al., 2006), and it is likely also participating in the recognition of the same cell wall component on *C. parapsilosis sensu lato*. The blocking experiments with either antibodies against receptors or laminarin, showed that TNFα stimulation by members of the *C. parapsilosis* complex, but not with *C. albicans* cells, depends on the engagement of TLR4, dectin-1 and MR with their ligands. These results contrast with our previous observation with β-eliminated cells, where TNFα levels upon disrupting the O-linked mannan-TLR4 interaction increased. The hypothesis to explain this apparent paradoxical response is that when the TLR4 receptor is blocked with anti-TLR4 antibodies, the O-linked mannans are still present on the surface of the fungal cell wall, hiding β1,3-glucan from the recognition by dectin-1. Interestingly, IL-6 production was only affected when human PBMCs were treated with anti-TLR4 antibodies and then stimulated with *C. orthopsilosis*, suggesting the O-linked mannans from this organism are different from those present on the surface of *C. parapsilosis sensu stricto* and *C. metapsilosis*. IL-10 production was strongly depend on engagement of dectin-1, as previously reported (Reid et al., 2009). It is noteworthy to mention that the IL-1β production by *C. parapsilosis sensu lato* was partially dependent on MR recognition, but not dectin-1 nor TLR4. This observation contrast with that reported earlier (Toth et al., 2013), where IL-1β stimulation by *C. parapsilosis sensu stricto* was dectin-1 dependent. Since the *C. albicans* immune recognition via dectin-1 is fungal strain-specific (Marakalala et al., 2013), it is feasible to conceive that this discrepancy can be explained because the genetic background of the strain used here (SZMC 8110) and in the previous report (GA-1) (Toth et al., 2013). In cells stimulated...
by *C. albicans*, IL-1β production is dependent on the activation of dectin-1, TLR2 and MR (van de Veerdonk et al., 2009); thus, different PRRs or downstream signaling components after *C. parapsilosis sensu lato* recognition could be responsible of this difference.

It has been previously demonstrated that cell wall preparations or non-purified wall components, such as zymosan, are potent inductors of cytokines (Brown et al., 2003; Netea et al., 2006); while purified wall components are unable to stimulated cytokine production (Dennehy et al., 2008; van de Veerdonk et al., 2009), and can block cytokine production when immune cells are stimulated (Maneu et al., 2011; Mora-Montes et al., 2011; Huang et al., 2012; Cohen-Kedar et al., 2014). These observations have been the base to propose the co-stimulatory theory, where a strong cytokine response involves the engagement of not only one but two or more PRRs at once (Netea et al., 2015). Although it is likely mannans from *C. albicans* and *C. parapsilosis sensu lato* are different, here it was demonstrated that purified N- and O-linked mannan preparations do not stimulate cytokine production and are capable to block proper *Candida* recognition, in a similar way that anti-MR and anti-TLR4 antibodies, respectively. Thus, they can be a reliable and alternative tool to assess the contribution of these PRRs during the innate immune recognition of fungal cells.

Overall, our data clearly demonstrate that the current knowledge about *C. albicans* biology and interaction with the host cannot be extrapolated to other members of the genus. *C. parapsilosis sensu lato* displays significant differences in the cell wall that impact the recognition by human PBMCs, stimulating stronger cytokine production, via the increased exposure of β1,3-glucan at the cell surface. Finally, our results suggest different contribution of dectin-1, MR and TLR4 in the recognition of the

| Cytokine stimulation by *C. parapsilosis sensu lato* | TNFα (%) | IL-6 (%) |
|---------------------------------------------------|---------|---------|
| No mannan included                                | 100     | 100     |
| Pre-incubation with *C. albicans* N-linked mannan | 34.4 ± 4.3* | 90.4 ± 5.4 |
| Pre-incubation with *C. albicans* O-linked mannan | 43.8 ± 7.6* | 99.3 ± 3.2 |
| Pre-incubation with *C. parapsilosis sensu stricto* N-linked mannan | 39.4 ± 6.3* | 93.3 ± 4.3 |
| Pre-incubation with *C. parapsilosis* O-linked mannan | 40.3 ± 6.3* | 101.2 ± 4.3 |
| Pre-incubation with *C. orthopsilosis* N-linked mannan | 33.4 ± 9.3* | 101.4 ± 3.4 |
| Pre-incubation with *C. orthopsilosis* O-linked mannan | 42.3 ± 4.9* | 95.8 ± 4.2 |
| Pre-incubation with *C. metapsilosis* N-linked mannan | 35.6 ± 6.6* | 95.3 ± 5.5 |
| Pre-incubation with *C. metapsilosis* O-linked mannan | 43.8 ± 3.9* | 100.5 ± 6.3 |

*P < 0.05, when compared to stimulation with no mannan included.
members of the C. parapsilosis complex, which has to be taken in consideration when analyzing the C. parapsilosis sensu lato-host interaction.

**AUTHOR CONTRIBUTIONS**

ML, AG, and HM-M conceived the study. EE-M, MN-A, LP-G, EM-M, and KC performed experiments. EE-M, MN-A, LP-G, EM-M, ML, KC, AG, and HM-M analyzed data. HM-M drafted the manuscript. EE-M, MN-A, LP-G, EM-M, ML, KC, AG, and HM-M revised and approved the manuscript.

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