New Insights into the Structure of (1→3,1→6)-β-D-Glucan Side Chains in the Candida glabrata Cell Wall

Douglas W. Lowman1,2, Lara J. West3, Daniel W. Bearden4, Michael F. Wempe5, Trevor D. Power6, Harry E. Ensley7, Ken Haynes3, David L. Williams1, Michael D. Kruppa8*

1 Department of Surgery, Quillen College of Medicine, East Tennessee State University, Johnson City, Tennessee, United States of America, 2 AppRidge International, LLC, Jonesborough, Tennessee, United States of America, 3 Department of Medicine, Imperial College London, London, United Kingdom, 4 Hollings Marine Laboratory, Analytical Chemistry Division, National Institutes of Standards and Technology, Charleston, South Carolina, United States of America, 5 School of Pharmacy, University of Colorado Health Sciences Center, Denver, Colorado, United States of America, 6 Department of Biochemistry and Molecular Biology, Sealy Center for Structural Biology and Molecular Biophysics, University of Texas Medical Branch, Galveston, Texas, United States of America, 7 Department of Chemistry, Tulane University, New Orleans, Louisiana, United States of America, 8 Department of Microbiology, Quillen College of Medicine, East Tennessee State University, Johnson City, Tennessee, United States of America

Abstract
β-glucan is a (1→3)-β-linked glucose polymer with (1→6)-β-linked side chains and a major component of fungal cell walls. β-glucans provide structural integrity to the fungal cell wall. The nature of the (1→6)-β-linked side chain structure of fungal (1→3,1→6)-β-D-glucans has been very difficult to elucidate. Herein, we report the first detailed structural characterization of the (1→6)-β-linked side chains of Candida glabrata using high-field NMR. The (1→6)-β-linked side chains have an average length of 4 to 5 repeat units spaced every 21 repeat units along the (1→3)-linked polymer backbone. Computer modeling suggests that the side chains have a bent curve structure that allows for a flexible interconnection with parallel (1→3)-β-D-glucon polymers, and/or as a point of attachment for proteins. Based on these observations we propose new approaches to how (1→6)-β-linked side chains interconnect with neighboring glucan polymers in a manner that maximizes fungal cell wall strength, while also allowing for flexibility, or plasticity.

Citation: Lowman DW, West LJ, Bearden DW, Wempe MF, Power TD, et al. (2011) New Insights into the Structure of (1→3,1→6)-β-D-Glucan Side Chains in the Candida glabrata Cell Wall. PLoS ONE 6(11): e27614. doi:10.1371/journal.pone.0027614

Editor: Joy Sturtevant, Louisiana State University, United States of America

Received September 29, 2010; Accepted October 20, 2011; Published November 11, 2011

This is an open-access article, free of all copyright, and may be freely reproduced, distributed, transmitted, modified, built upon, or otherwise used by anyone for any lawful purpose. The work is made available under the Creative Commons CC0 public domain dedication.

Funding: This work was supported, in part, by National Institutes of Health grants R01 GM53552 and R01 GM83016. The funders had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript. Also, AppRidge International, LLC, did have a role in the data analysis and preparation of the manuscript. DLW and MDK do not have any financial stake in AppRidge, nor do they receive any compensation from AppRidge International, LLC. The authors are in strict adherence to PLoS ONE policies on data sharing and materials. At present, there is no public repository for structural data on complex carbohydrates, however, the data and materials from this article will be provided freely upon request.

* E-mail: KRUPPA@mail.etsu.edu

Introduction
Fungal cells are surrounded by a wall that consists of four main components: glucan, chitin, mannan and mannanprotein. These carbohydrates play an important role in maintaining the shape and viability of fungal cells in response to osmotic challenge, and they are the point of contact with the environment. A major carbohydrate component of the cell wall is (1→3)-β-D-glucan [1]. Glucans are generally arrayed as triple helical polymers, which function to maintain the rigidity of the cell wall [1]. The (1→3)-β-D-glucan is thought to form a lattice support structure through the other molecules are attached. Chitin has been suggested to crosslink with both (1→3)- and (1→6)-β-D-glucan polymers [2–4] but the nature of the crosslink has not clearly been established. (1→6)-β-D-glucan side chain branches attached to a (1→3)-β-D-glucan polymer backbone, have been reported for several different fungal species [3–8]. It is thought that the (1→6)-β-D-glucan serves as attachment sites for chitin, mannan, and mannanprotein [2,3] and they may also serve to link or interconnect adjacent (1→3)-β-D-glucan polymer strands. However, the nature of the attachment to these carbohydrates has not been structurally characterized except that mannanprotein can be linked through a glycosylphosphatidylinositol anchor [9,10]. Since (1→6)-β-D-glucan appears to serve a major role for attachment of these polymers, it is important to understand the structural nature of (1→6)-β-D-glucan as it exists when attached to the (1→3)-β-D-glucan polymer.

Previously, Tada and coworkers [8] characterized the structure of Grifolan-LE, isolated from the mushroom Grifola frondosa. This is a (1→3,1→6)-β-D-glucan polymer containing a (1→3)-β-linked backbone chain and side chains composed of a single (1→6)-β-D-glucan repeat unit attached, on average, to the backbone chain every 3 repeat units. In this current study, we have isolated glucan from wild-type and several mutant C. glabrata strains. Using a modified glucan extraction procedure, we have isolated a (1→3,1→6)-β-D-glucan that contains a unique poly-(1→6)-β-D-linked side chain structure. Based on extensive 1H and 13C 1D and homonuclear (COSY, NOESY and TOCSY) and heteronuclear (HSQC, HSQC-NOESY, HSQC-TOCSY, and HMBC) 2D NMR studies, we have determined that glucan isolated from C.
*G. glabrata* contains a (1→3)-β-D-glucan polymer backbone with (1→6)-β-D-glucan-containing side chains composed of multiple (1→6)-β-linked glucosyl repeat units. Additionally, we have extended our investigation to include molecular modeling of the side chain structural conformation. The structure of the side chain appears to have a hook-like or curved structure, which may allow for interconnection with parallel (1→3)-β-D-glucan polymers, or serve as a point of attachment to chitin, mannan or a glycosylphosphatidylinositol anchor to a protein. Another possibility is that this hook-like structure plays a different role in the fungal cell wall structure, one that allows for physical grabbing of another (1→3)-β-D-glucan polymer, much in the same manner that a Velcro hook grabs a fiber strand. Based on these results, we propose that the structure of the (1→6)-β-D-glucan side chain may help to explain how the cell wall architecture allows for a rigid cell wall that also exhibits flexibility as the fungal cell grows.

### Results

**Elucidation of the (1→6)-β-linked side chain structure**

Tada and coworkers [8] recently presented the complete structural characterization of Grifolan-LE isolated from *G. fondosa*. Grifolan-LE is a purified (1→3)-β-D-glucan with single (1→6)-β-linked glucosyl side chains every third backbone repeat unit on average. They assigned specific proton NMR resonances to the anomeric proton of the glucosyl side chain repeat unit and to one of the methylene protons of the branchpoint glucosyl repeat unit in the backbone chain connected by a (1→6)-β-linkage. In the proton NMR spectrum, the resonances for the anomeric proton and methylene proton were of equal integrated area, which is consistent with a side chain structure containing only a single repeat unit. The proton NMR spectrum of the glucan isolated from *C. glabrata* acc2 strain (Fig. 1) resembles the NMR spectrum of Grifolan-LE except that the integral areas of the anomeric proton and one of the methylene protons involved in the (1→6)-β-linkages in the side chain are not of equal area. In addition, the NMR spectrum (Fig. 1) shows 5 major resonances previously assigned to glucosyl repeat units of a (1→3)-β-D-glucan polymer chain [11]. These assignments were confirmed by COSY 2D NMR (data not shown) and are summarized in Table 1.

The branchpoint glucosyl repeat unit (Br) in the (1→3)-β-linked polymer chain is unique in that it is the only repeat unit in either the backbone chain or the side chain that is substituted in three positions – the 1-, 3-, and 6-positions. Using the spectral region of the C3 carbon in the HSQC-TOCSY 2D NMR spectrum (Fig. 2), only the carbon resonating at 87.14 ppm (C3Br) exhibits long-range correlations to equivalent methylene protons at 4.00 (C3Br/H6Br) and 3.64 (C3Br/H6’Br) ppm. Other protons correlated to the C3Br resonance include those assignable to the branchpoint repeat unit (Table 1) as follows: H1 at 4.36 ppm (C3Br/H1Br), H2 at 3.24 ppm (C3Br/H2Br), H3 at 3.50 ppm (C3Br/H3Br), H4 at 3.32 ppm (C3Br/H4Br), and H5 at 3.23 ppm (C3Br/H5Br). These 1H and 13C NMR resonance assignments are based upon results from HSQC-TOCSY and HSQC 2D NMR experiments. The HSQC-TOCSY spectrum provides information about protons that are scalar coupled within the same repeat unit, or spin system, with one of the protons scalar coupled to a specific carbon. The strength of these correlations is dependent upon the distance of the individual protons from the specific carbon atom in the spin system and the relaxation times of the nuclei involved. Since the TOCSY spin lock time is short (60 ms), only 4 protons correlate to each carbon around the glucosyl residue in this study. Therefore, the correlation to H1Br observed from C3Br (C3Br/H1Br; 67.14 ppm/4.36 ppm) in the HSQC-TOCSY 2D NMR spectrum is important for establishing C1Br through C4Br in the branchpoint repeat unit while the correlation to H6Br/H6’Br (4.00 ppm/3.64 ppm) observed from C3Br is important for establishing C3Br through C6Br in the branchpoint repeat unit (Fig. 3). The correlation from the methylene carbon of the branchpoint repeat unit (C6Br) and the anomeric proton (H1) of the first side chain repeat unit (SC3) across the (1→6)-β-linked glycosidic bond is established in the HMBC 2D NMR spectrum (C6Br/H1SC1, Fig. 4: 67.86 ppm/4.25 ppm). The anomeric proton of SC1 correlates with H2 in the COSY 2D NMR spectrum (data not shown), and then H2 correlates with H3 and the remainder of the glycosyl protons H4 through H6/H6’ in the TOCSY 2D NMR spectrum (Fig. 3b). Resonances H3 through H6/H6’ of SC1 are coincident with similar resonances in internal side chain repeat units (SC INTERNAL) from the COSY (data not shown), NOESY (Fig. 5a), and TOCSY (Fig. 5b) 2D NMR spectra. Individual resonances from SC INTERNAL each exhibit the same chemical shifts (Table 1). Three types of (1→6)-β-linkages across glycosidic bonds are evident in the NOESY 2D NMR spectrum (Fig. 5a). The (1→6)-β-linked glycosidic bonds for the SC1 (H6’Br/H1SC1; 4.00, 3.64 ppm/4.25 ppm) and SC INTERNAL (H6’SC/H1SC3α+; 4.01, 3.61 ppm/4.27 ppm) have been identified. The remaining (1→6)-β-linkage exhibits H6,H6’ correlations to an anomeric proton with chemical shifts typical of a (1→6)-β-linkage. The anomeric proton in this linkage has a chemical shift characteristic of the anomeric proton in the non-reducing terminus (NRT H1) of the (1→3)-linked polymer backbone. TOCSY spectra of the NRT glycosyl repeat units (SC NRT H1 and NRT H1 in Fig. 5b) are characteristic of monosubstituted glucosyl repeat units. Therefore this third (1→6)-β-linkage is assigned to the glycosidic linkage between the methylene protons of the next to the last (second to the last) glycosyl repeat unit (SC NRT H6,H6’; 4.03, 3.66 ppm) and the anomeric proton of the non-reducing terminus of the (1→6)-β-linked side chain (SC NRT H1; 4.37 ppm). These data complete the assignment of all repeat units and glycosidic linkages in the (1→6)-β-linked side chain of this glucan isolated from *C. glabrata* and support the side chain structure containing multiple (1→6)-β-D-glucosyl repeat units (Fig. 6).

In addition, the anomeric proton for the α-conformer of the reducing terminus (α-RT) of the polymer backbone is assigned to 4.99 ppm while the chemical shift of the β-RT anomeric proton is overlapped by resonances from the anomeric protons of Br, SC NRT and NRT. Anomeric proton resonances for α- and β-glucose are observed at 4.93 and 4.30 ppm, respectively. Anomeric protons for the α- and β-conformers of SRT are observed at 4.46 and 4.49 ppm, respectively. These anomeric proton resonance assignments are in agreement with previous assignments by Kim and coworkers [12] and in general agreement with assignments by Størseth and coworkers [7].

### Calculation of the average side chain length and branching frequency

To determine the average chain length and its average branching frequency for side chains containing multiple (1→6)-β-D-glucosyl repeat units, the integrated areas of the resonances assigned to H1 of SC NRT, H1SC, and H6SC of the side chain glycosidic bond, and H1 of the (1→3)-β-linked repeat units in the polymer backbone were compared. Interestingly, the integrated areas of the resonances assigned to H1SC (4.27 ppm) and H6SC (4.01 ppm) of the glycosidic linkages in the (1→6)-β-linked side chain are not equal in the glucan isolated from *C. glabrata* (inset in Fig. 1). The integrated area of the H6SC multiplet resonance is larger than the area of the H1SC multiplet resonance. Integration
of these two resonances gives an area ratio of 0.787:1 for resonances H1SC and H6SC, respectively, not the 1:1 ratio reported for Grifolan-LE with the single (1\R6)-b-linked glucosyl repeat unit in the side chain [8]. The difference in areas of the two resonances represents the integral area assignable to the anomeric proton H1 in SC NRT. The ratio suggests that the average length of the side chain contains 4.7 (1\R6)-b-D-linked repeat units. The ratio of the areas from the resonances assigned to H1 of SC NRT and H1 of the (1\R3)-b-linked repeat units in the polymer backbone indicates that a side chain is attached to the (1\R3)-b-D-linked polymer backbone on average every 21 repeat units.

**Impact of the glucan isolation scheme on the average side chain structure**

Several strains of *C. glabrata* isolated under different conditions have been examined by proton NMR. Isolation methods 1A and 1B.

**Table 1.** Proton (top) and $^{13}$C (bottom) NMR chemical shifts (in ppm) for structural features characterized in the glucan isolated from *C. glabrata ace2* (HLS122) strain.

| Chemical Shift, ppm | Proton Assignment | (1→3)-b-Linked Backbone Chain | Br | SC1 | SC Internal (1→6)-b-Linked Side Chain | SC SNRT (1→6) | SC NRT (1→6) | NRT (1→3) |
|---------------------|------------------|--------------------------------|----|-----|--------------------------------------|---------------|---------------|----------|
| H1                  |                  | 4.53                           | 4.36| 4.25| 4.27                                 | 4.27          | 4.37          | 4.39     |
| H2                  |                  | 3.31                           | 3.24| 3.03| 3.04                                 | 3.04          | 3.10          | 3.09     |
| H3                  |                  | 3.48                           | 3.39| 3.19| 3.19                                 | 3.19          | 3.23          | 3.22     |
| H4                  |                  | 3.27                           | 3.32| 3.15| 3.15                                 | 3.16          | 3.15          | 3.15     |
| H5                  |                  | 3.27                           | 3.23| 3.32| 3.32                                 | 3.33          | 3.25          | 3.25     |
| H6                  |                  | 3.73                           | 4.00| 4.01| 4.01                                 | 4.03          | 3.72          | 3.72     |
| H6'                 |                  | 3.48                           | 3.64| 3.61| 3.61                                 | 3.66          | 3.46          | 3.46     |

| Carbon Assignment   |                  | C1                              | 102.70| 102.43| 103.10| 102.96| 102.96| 103.66| 103.48|
|---------------------|------------------|---------------------------------|-------|-------|-------|-------|-------|-------|-------|
|                     |                  | C2                              | 72.54 | 71.99| 73.60 | 73.16 | 73.16 | 76.47 | 76.57 |
|                     |                  | C3                              | 85.97 | 87.14| 76.32 | 76.31 | 76.32 | 75.89 | 75.95 |
|                     |                  | C4                              | 68.15 | 67.90| 69.74 | 69.74 | 69.74 | 69.79 | 69.79 |
|                     |                  | C5                              | 76.08 | 75.88| 75.13 | 75.23 | 75.22 | 76.48 | 76.48 |
|                     |                  | C6                              | 60.63 | 67.86| 68.24 | 68.17 | 68.23 | 60.82 | 60.84 |
1B represent duplicate glucan extractions using 1 N phosphoric acid while method 2 represents glucan extracted with 2 N phosphoric acid. All other parameters of the extraction were held constant. This was done because we have previously shown that by decreasing the concentration and/or strength of the protic acid it is possible to preserve more of the native structure of the beta-glucan [26]. At the higher acid concentration, the (1→6)-β-linked side chains are 2 repeat units in length on average while the branching frequency differs for the 4 strains examined (Table 2). The strains, other than wild type, were chosen due to their putative role in mannan synthesis as well as potential in altering the wall structure, we were curious as to whether there were any changes in the glucan of these strains, since compensatory changes could occur with other carbohydrate components in these mutants. For wild-type and anp1 strains, the branching frequency is about one-third larger than for the mnn2 and mnn11 strains. At the lower acid concentration, the average side chain length is about twice the length of that from the higher acid concentration extraction and the branching frequency is constant with side chain branches on average every twenty repeat units in the polymer backbone. Thus, the strength of the protic acid used to isolate the glucans significantly impacts the length of the side chain, i.e. lower acid concentration results in great preservation of the side chain architecture.

Molecular modeling of the glucan side chain

Based upon the structural characterization of the (1→6)-β-linked side chain, we explored molecular structure by performing computational chemistry. We first used molecular modeling – molecular mechanics (MM) – on a model compound consisting of a glucan with ten (1→3)-β-linked repeat units in the polymer backbone and five (1→6)-β-linked repeat units in the side chain attached to the third repeat unit in the polymer backbone. These results were compared to similar calculations for a linear polymer.
containing ten (1→3)-β-linked repeat units in the polymer chain (Fig. 7a). It is important to mention that other researchers have also used in silico molecular mechanics (MM) to conduct energy minimizations to probe glucans, specifically glucan:mycotoxin and glucan:zearalenone complex interactions[13,14]. It is also important to recognize that various researchers commonly use different computational programs and/or different approaches to probe theoretical structure. Our approach utilized a local minimum of the linear form and was subsequently used to conduct a semi-empirical Austin-Model (AM1) calculation. The resulting molecular structure was then used to conduct ab initio geometry optimizations at the Hartree-Fock (HF)/STO-3G basis set; an approach previously used by our group to investigate theoretical structure of a linear decasaccharide and a branched nonasaccharide[15]. Addition of the (1→6)-β-linked side chain resulted in the branchpoint repeat unit (Br) exhibiting a skewed conformation. Addition of the (1→6)-β-linked side chain, initially as a linear side chain, to a linear (1→3)-β-linked polymer backbone resulted in a calculated, lowest energy side chain structure where the repeat units near the non-reducing terminus of the (1→6)-β-linked side chain curled back onto side chain repeat units closer to the branchpoint repeat unit (Fig. 7b). While HF/STO-3G may be considered a first approximation, our main computational goal was to generate a theoretical cartoon of the branched model glucan. To improve upon the STO-3G result, we took the geometry optimization and performed ab initio geometry optimizations using Pople basis set 6-31G(d)[16–18]. This basis set was designed to obtain structures that are comparable to those obtained from experimental procedures and consistently used at the Hartree-Fock level, in the gas-phase, to obtain new geometrical parameters commonly used to conduct AMBER calculations. The data obtained at 6-31G(d) were similar to the predicted structure by STO-3G (data not shown) and thereby does not alter the general non-covalent hooking interaction hypothesis.

Discussion

Previous work by Manners and coworkers [5] proposed an often-repeated side chain structure containing a single (1→6)-β-linked glucosyl repeat unit connecting a chain of (1→3)-β-linked glucosyl repeat units to the polymer backbone. While the branchpoint repeat unit would have a 2D NMR spectral signature similar to that observed for our glucan isolate, the remainder of the side chain structure would not present a similar 2D NMR signature due to the presence of the (1→3)-β-linkages. In this current study we have presented the first, to our knowledge, detailed structural characterization of the (1→6)-β-linked side chain in (1→3,1→6)-β-D-glucan isolated from the fungal cell wall of C. glabrata. This detailed structural analysis will serve as a model for the global structural characterization of the glucan in terms of average (1→6)-β-linked side chain length and branching frequency along the (1→3)-β-linked polymer backbone. We have also observed that the methodology used to extract the glucan can impact the structure of the isolated glucan. In addition, molecular modeling based upon these NMR results provides new insight into how (1→3,1→6)-β-D-glucans may be incorporated into the carbohydrate architecture of the fungal cell wall.

We have observed that the H1SC and H6SC resonances from the (1→6)-β-linkage also exhibit several inflection points (inset in Fig. 1). Størseth and coworkers [7] suggest that these multiple inflection points reflect multiple chain lengths of the (1→6)-linked
Differences in acid concentration affect side chain length

Isolation of glucan from several strains of *C. glabrata* using two different methods illustrates the role isolation protocol plays relative to the final structure of the isolated glucan. At the lower phosphoric acid concentration, the average side chain length is about twice the length of that from the higher phosphoric acid concentration while the average distance between branches along the polymer backbone is less (Table 2). The higher acid concentration results in reduced side chain length as well as side chain content on average compared to the lower acid concentration. The resulting glucan structures from these two different isolation methods clearly demonstrate the importance of consistency in the isolation scheme. The structure of the isolated glucan can be dramatically impacted by differences in isolation protocols, which can impact ligand-receptor binding studies as well as action of the glucan in a biological system. Thus, our data indicate that isolation conditions which are less harsh, i.e. lower acid concentration, result in preservation of the glucan side chain structure. We speculate that milder isolation conditions result in side chain structures that are most closely representative of the side chains found in the fungal cell wall.

The nature of the side chains, i.e., their structure and branching frequency, and how they impact ligand-receptor interactions are important topics relative to understanding how pathogenic and non-pathogenic fungi interact with their environment, including the innate immune system. From a human health standpoint, glucans are critical to the structure and function of the fungal cell wall. They are the primary pathogen associated molecular patterns (PAMPs) recognized by the innate immune system [15,21], and provide a drug development target for glucan synthase inhibitors, the newest anti-fungal drugs.

A structural feature leads to new insights about glucan organization

Adams and coworkers [15] have discussed the importance of these side chain structures relative to interaction of (1→3,1→6)-β-D-glucan PAMPs with Dectin-1, the primary pattern recognition receptor (PRR) for glucans [21]. To better understand how these glucans with longer side chains may interact with Dectin-1 as well as neighboring polymer chains, theoretical optimized molecular geometries were investigated for the model compound containing five (1→6)-β-linked repeat units in the side chain (Fig. 7). This side chain structural proposal is interesting in that it may offer insight into how these side chains aid in not only the rigidity and structure, but also the plasticity, of the fungal cell wall [22]. At present, it is not clear whether glucan polymer strands within the fungal cell wall are interlinked, and if they are, it is not clear how this is accomplished. The side chains have non-reducing termini, which would not allow the formation of covalent linkages between glucan polymer strands. However, the curled side chain may provide a mechanism for interaction of the side chain with other glucan chains by wrapping around other polymer chains, thus not requiring covalent bonding between the side chain repeat units and other chains in the polymer. We propose that the curled side chain structures may connect neighboring glucan polymer chains in a manner that maximizes wall strength (Fig. 8a), while also glycosidic bond resulting in additional resonances in the proton NMR spectrum of the isolate [19,20]. Therefore, it is reasonable to consider that, while the multiple inflection points in the H1SC and H6SC resonances may result from different side chain lengths, they may also result from different conformations observed for the (1→6)-β-linked glycosidic bond, thus providing additional support for the presence of multiple (1→6)-β-linked glucosyl repeat units in the average side chain.

Figure 5. 2D NMR spectra of the glycosidic linkages and non-reducing termini of the (1→3,1→6)-β-D-glucan isolated from *C. glabrata* ace2 strain. (a) The three different (1→6)-β-linked glycosidic bonds from the side chain are detailed in the NOESY 2D NMR spectrum for SC1, SC Internal, and SC NRT glucosyl groups associated with H1 SC1, SC H1 and SC NRT H1. A: H6br,H6′br/H1SC1; B: H6SCn,H6′SCn/ H1SC(n+1); C: H6SC(n-1),H6′SC(n-1)/H1SCNRT. (b) Similarity of the structures of the glycosyl group associated with SC NRT H1 and NRT H1 is indicated in the TOCSY 2D NMR spectrum. doi:10.1371/journal.pone.0027614.g005

side chain. Two other studies showed that (1→6)-β-D-glucan polymers isolated from *Actinobacillus suis* and *Malassezia sympodialis* exhibits multiple conformations about the (1→6)-β-linked glycosidic bond resulting in additional resonances in the proton NMR spectrum of the isolate [19,20]. Therefore, it is reasonable to consider that, while the multiple inflection points in the H1SC and H6SC resonances may result from different side chain lengths, they may also result from different conformations observed for the (1→6)-β-linked glycosidic bond, thus providing additional support for the presence of multiple (1→6)-β-linked glucosyl repeat units in the average side chain.
allowing flexibility, or plasticity, in the fungal cell wall. In addition to a direct covalent interaction between strands of polymer, there are several possible interactions through hydrogen bonding, through which the side chains can interact. First there could be a direct hooking of two \((1\rightarrow6)\)-side chains from two individual \((1\rightarrow3)\)-linked polymers (Fig. 8b) allowing for a mesh like structure to form; this interaction with the hooking of a \((1\rightarrow6)\)-branch around a \((1\rightarrow3)\)-linked polymer (Fig. 8c) would give both strength and flexibility allowing for “slippage” of the \((1\rightarrow6)\) side chain down the length of the \((1\rightarrow3)\) polymer. Another arrangement could be the catching of a \((1\rightarrow6)\) branch that is cross-linked to a second branch with a hooked branch on a third polymer (Fig. 8d) forming a “catch” structure similar to that seen in Fig. 8b. As much of the \((1\rightarrow3)\) glucan is arranged in a triple helix (Fig. 8e) the presence of \((1\rightarrow6)\) side chains may play a lesser role in the integrity of the glucan matrix and may serve as point of attachment for other molecules such as mannan, mannoprotein, chitin or some other molecules not yet identified in the cell wall. Additional investigation will be required to fully elucidate the role of \((1\rightarrow6)\) side chains in fungal cell wall architecture. However, what we have proposed provides a potential explanation for the rigidity, flexibility and interconnections that the fungal cell wall exhibits.

Materials and Methods

Strains and growth conditions

*C. glabrata* strain ATCC2001 was obtained from the American Type Culture Collection. *C. glabrata* *ace2* (HLS122) strain has been described previously [23]. *C. glabrata* *mnn2*, *mnn11*, and *anp2* strains were constructed by gene replacement [24]. Strains were routinely cultured on YPD (1% yeast extract, 2% peptone, 2% dextrose, 2% agar) at 30°C.

Glucan isolation

Strains were cultured in 1L of YPD at 30°C for 18 hr with agitation. Cells were harvested and then lyophilized. The lyophilized cells were extracted using modifications of the methods of Williams, Mueller and coworkers [25,26]. Specifically, in method 1 the phosphoric acid concentration was 1 N, while in method 2 the phosphoric acid concentration was 2 N. Decreasing the phosphoric acid concentration reduces polymer degradation during isolation [26]. It is important to note, that this extraction method does not result in phosphorylation of the glucan. All other aspects of the method were as previously reported by our group [25,26].

Table 2. Comparison of average side chain lengths, SCn, and branching frequencies, Br Freq, for several strains of *C. glabrata* isolated by 2 different methods.

| Isolation Method | Strain | SCn  | Br Freq |
|------------------|--------|------|---------|
| 1A               | Wild-type | 4.0  | 22.2    |
| 1A               | *ace2*  | 4.2  | 20.3    |
| 1A               | *mnn11* | 4.1  | 21.8    |
| 1B               | Wild-type | 4.4  | 19.7    |
| 1B               | *ace2*  | 4.7  | 20.7    |
| 1B               | *mnn11* | 4.1  | 21.4    |
| 2                | Wild-type | 2.5  | 40.7    |
| 2                | *anp1*  | 1.8  | 43.0    |
| 2                | *mnn2*  | 2.0  | 30.8    |
| 2                | *mnn11* | 2.1  | 30.5    |

doi:10.1371/journal.pone.0027614.t002
Proton and $^{13}$C NMR spectra were collected on a Bruker Avance II 800 NMR spectrometer using a 5 mm TCI inverse cryoprobe operating at 343 K. For initial NMR spectra, ~25 mg of glucan was solvated in 950 μL DMSO-d$_6$ (Sigma-Aldrich, “100”, p/n 156914, CAS 2206-27-1) with 40 μL trifluoroacetic acid-d (Cambridge Isotope Laboratories, 99.8% deuterated, p/n DLM-46, CAS 599-00-8) to shift the exchangeable proton resonances downfield. Approximately 550 μL of the sample was placed into a 5 mm NMR tube. Trifluoroacetic acid was added just prior to NMR analysis, in order to reduce the exposure time during which depolymerization could occur. For $^{13}$C NMR spectra, a second sample was prepared using ~15 mg of glucan dissolved in 600 μL DMSO-d$_6$ with no added trifluoroacetic acid and transferred to a 5 mm NMR tube. Proton and $^{13}$C 1D and COSY [27,28], NOESY [29,30], TOCSY [31], HSQC [32–34], HSQC-TOCSY [35,36], HSQC-NOESY [35], and HMBC [37] 2D NMR spectra were obtained in this study. For 1D NMR experiments, chemical shift referencing used absolute referencing based on the $^2$H DMSO-d$_6$ lock signal and $^1$H and $^{13}$C gyromagnetic ratios. For 2D NMR experiments, chemical shift referencing was accomplished relative to the anomeric resonance of the (1→3)-β-linked backbone repeat unit at 4.53 ppm for $^1$H and 102.70 for $^{13}$C relative to DMSO-d$_6$ residual protons as referenced above. We performed all $^1$H homonuclear 2D NMR spectra on one sample and all $^{13}$C-{ $^1$H} heteronuclear 2D NMR spectra on a second solution of the same sample. 1D NMR spectra were collected and processed as follows: for $^1$H NMR, 64 90 μs scans with 8 dummy scans, acquisition of 65,536 real points zero-filled once and Fourier transformed to 65,536 complex points, 10.5 ppm sweep width, centered at 3.85 ppm with a total acquisition time of 3.9 s, exponential apodization with 0.3 Hz broadening, and 3 s pulse delay for a total interpulse period of 6.9 s; for $^{13}$C NMR, 16,384 30 μs scans with 8 dummy scans, acquisition of 65,536 real points zero-filled once and Fourier transformed to 65,536 complex points, 65 ppm sweep width, centered at 80 ppm with a total acquisition time of 2.5 s, exponential apodization with 3 Hz broadening, and 2 s pulse delay.

![Molecular modeling](doi:10.1371/journal.pone.0027614.g007)
delay for a total interpulse period of 4.5 s. 2D NMR spectra were collected and processed as follows: for homonuclear NMR experiments, 512×256 point matrix was zero-filled to 2048×1024 points, 32 scans per row with 16 dummy scans, 3 ppm sweep width centered at 3.85 ppm, cosine-squared apodization in both dimensions, and 2 s pulse delay; for heteronuclear NMR experiments, 512×512 point matrix was zero-filled to 1024×1024 points, 64 scans per row with an initial 64 dummy scans, 3 ppm sweep width centered at 3.85 ppm for the $f_2$ dimension and 80 ppm sweep width centered at 65 ppm for the $f_1$ dimension, cosine-squared apodization in both dimensions, and 1.5 s pulse delay except for HMBC which was zero-filled to 4096×1024 points and sine apodized. Adiabatic $^{13}$C decoupling was used during the $t_2$ acquisition period of the heteronuclear experiments, except for the HMBC experiment, and shaped inversion pulses were used on the $^{13}$C channel in inverse heteronuclear experiments. NMR spectra were processed using TOPSPIN 2.1 running on the Avance II 800 NMR and TOPSPIN 3.0.b.8 running on Windows XP Professional operating system under VMWare Fusion version 2.0.5 (VMWare, Inc., Palo Alto, CA) on a Macintosh MacBook Pro.

Molecular modeling

Chemical structures were drawn in a linear format using CS ChemDraw Ultra® (version 6.0.1; Cambridge Soft Corporation; Cambridge, MA). The structures were then copied into CS Chem3D Ultra®. For each polysaccharide, a molecular mechanics (MM) minimization was conducted using a root-mean-square (RMS) of 0.005. Next, molecular geometries were generated in the Gaussian Z-matrix style via the CS MOPAC application. For each compound, an Austin-Model (AM1) semi-empirical calculation (closed shell, tight convergence criteria) was conducted using the Gaussian® software package (Gaussian, Inc.; Carnegie, PA) [38]. Ab initio geometry optimizations were subsequently performed using Gaussian 03 at the Hartree-Fock level of theory using the STO-3G [16,17] basis set. Additional ab initio geometry optimizations were performed using Gaussian 03 at the Hartree-Fock level of theory using the 6–31G(d) basis set and compared (data not shown) to STO-3G results[16-18]. For STO-3G analysis geometry optimizations using implicit solvation were performed with the Onsager method [18] using the dielectric constant of water (78.39) and the solute radius; calculated with the more accurate molecular volume computation (Volume_Tight). These calculations were performed with a Microway AMD (Microway Technology, Plymouth, MA) dual 64 bit 2.0 GHz central processing unit (CPU) with 4GB RAM running Fedora Core 3 and an Aspen Systems 30 CPU cluster (Wheatridge, CO). Each blade contained two 32-bit Xeon 3.2 GHz CPUs with 4GB RAM running Red Hat Linux.

Acknowledgments

We acknowledge the support of the Hollings Marine Laboratory NMR Facility for this work.

Disclaimer

Commercial equipment or materials are identified in this paper to specify adequately the experimental procedure. Such identification does not imply recommendation or endorsement by NIST, nor does it imply that the materials or equipment identified are necessarily the best available for the purpose.

Author Contributions

Conceived and designed the experiments: DLW DWL MDK. Performed the experiments: DWL HEE DWB MFW TDP DLW MDK. Analyzed the data: DWL DWB HEE TDP MFW DLW MDK. Contributed reagents/materials/analysis tools: MFW KH LJW TDP DWB HEE DLW MDK. Wrote the paper: DLW DWL MDK.

Figure 8. Conceptual model of the role of (1→6)-β-linked glucosyl side chains in the fungal cell wall structure. Possible arrangements of (1→6)-β-D-glucan branches within the glucan matrix include (a) direct covalent cross-linking between two (1→3)-β-linked polymers, or (b) non-covalent interactions by “hooking” across (1→6)-β-linked side chains, or (c) across (1→3)-β-linked polymers, or (d) “catching” a covalent (1→6)-β-linked cross-linking branch. The triple helix arrangement (e) with associated (1→6)-β-linked side chains may serve as points for attachment to chitin, mannan, mannoprotein, GPI protein anchor or other possible molecules. doi:10.1371/journal.pone.0027614.g008
References

1. Deslandes Y, Marchessault RH, Sarko A (1980) Triple-helical structure of (1-3)-D-glucan. Macromolecules 13: 1466–71.

2. Cabib E (2009) Two novel techniques for determination of polysaccharide cross-links show that Calp B and Calp C1 attach chitin to both β(1,3)- and β(1,6)-glucan in the Schizosaccharomyces octosporus cell wall. Eukaryotic Cell 8: 1626–36.

3. Kaptyn JC, Ram AF, Groos EM, Kollar R, Montijn RC, et al. (1997) Altered extent of cross-linking of β1, 3-glucosylated mannan-proteins to chitin in Schizosaccharomyces octosporus mutant with reduced cell wall β1, 3-glucan content. Journal of Bacteriology 179: 6279–84.

4. Kollar R (1997) Architecture of the yeast cell wall, beta(1→6)-glucan interconnects mannan-protein, beta(1→3)-glucan and chitin. J Biol Chem 272: 17762–75.

5. Manners DJ, Masson AJ, Patterson JC (1978) The structure of a beta-(1→6)-glucan from yeast cell walls. Biochem J 159: 19–30.

6. Sugawara T, Takahashi S, Osumi M, Ohno N (2004) Refinement of the β(1,3)-D-glucan obtained from liquid-cultured Grifola frondosa by solution NMR spectroscopy. Carbohydr Res 339: 2253–65.

7. Størseth TR, Kirkvold S, Skjermo J, Reitan KI (2006) A branched β-D-(1,3, 1, 6)-glucan from the marine diatom Chaetoceros debilis (Bacillariophyceae) characterized by NMR. Carbohydr Res 341: 2108–14.

8. Tada R, Adachi Y, Ishibashi K-I, Ohno N (2009) An unambiguous structural elucidation of a 1,3-β-D-glucan obtained from liquid-cultured Grifola frondosa by solution NMR experiments. Carbohydr Res 344: 400–4.

9. de Nobel H, Lipke PN (1994) Is there a role for GPIs in yeast cell-wall assembly? Trends Cell Biol 4: 42–5.

10. Lu C-F, Montijn RC, Brown JL, Kias F, Kurjan J, et al. (1995) Glycosyl Phosphatidylinositol-dependent cross-linking of a-1, 2-glucan and β1, 6-glucan in the Schizosaccharomyces octosporus cell wall. The Journal of Cell Biology 128: 333–40.

11. Enyele HE, Tobias B, Perutz HA, McNamara RB, Jones EL, et al. (1994) NMR spectral analysis of a water-insoluble (1-3)-β-D-glucan isolated from Saccharomyces cerevisiae. Carbohydr Res 219: 203-13.

12. Muñoz P, Enyele H, Perutz H, McNamara R, Jones E, et al. (1997) The application of various prootic acids in the extraction of [1->3]-β-D-glucan from Saccharomyces cerevisiae. Carbohydr Res 299: 203–8.

13. Shaw AA, Sallam C, Dauphin J-F, Ancian B (1996) Artifact-Free PFG-Enhanced Double-Quantum-Filtered COSY Experiments. J Magn Reson A 120: 110–5.

14. Ancian B, Bourgeois I, Dauphin J-F, Shaw AA (1997) Artifact-Free Pure Absorption PFG-Enhanced DQF-COSY Spectra Including a Gradient Pulse in the Evolution Period. J Magn Reson 125: 548–54.

15. Jeener J, Meier BH, Bachmann P, Ernst RR (1979) Investigation of exchange processes by two-dimensional NMR spectroscopy. J Chem Phys 71: 4546–53.

16. Wagner R, Berger S (1996) Gradient-Selected NOESY–A Fourfold Reduction of the Measurement Time for the NOESY Experiment. J Magn Reson A 123: 119–21.

17. Bax A, Davis DG (1985) MLEV-17-based two-dimensional homonuclear magnetization transfer spectroscopy. J Magn Reson 65: 335–60.

18. Palmer III AG, Cavanagh J, Wright PF, Rance M (1993) Sensitivity improvement in proton-detected two-dimensional heteronuclear correlation NMR spectroscopy. J Magn Reson 93: 131–70.

19. Kay LE, Krüger P, Saarinen T (1992) Pure absorption gradient enhanced heteronuclear single quantum correlation spectroscopy with improved sensitivity. J Am Chem Soc 114: 10663–5.

20. Schleicher J, Schenkerling M, Santler M, Schmidt P, Schledtetzky O, et al. (1994) A general enhancement scheme in heteronuclear multidimensional NMR employing pulsed field gradients. J Biomol NMR 4: 301–6.

21. Norwood NJ, Boyd J, Heritage JE, Soffe N, Campbell ID (1990) Comparison of techniques for 1H-detected heteronuclear 1H-15N Spectroscopy. J Magn Reson 87: 488–501.

22. Cavanagh J, Palmer III AG, Wright PE, Rance M (1991) A general enhancement scheme in heteronuclear multidimensional NMR employing pulsed field gradients. J Am Chem Soc 113: 4899–901.

23. Kamran M, Calcagno AM, Findon H, Bignell E, Jones MD, et al. (2004) Inactivation of transcription factor gene JEC2 in the fungal pathogen Candida glabrata results in hypervirulence. Eukaryotic Cell 3: 146–52.

24. West JI. Glycosylation and virulence in Candida glabrata Imperial College London; 2009.

25. Williams DL, McNamre RB, Jones EL, Perutz HA, Enyele HE, et al. (1991) A method for the solubilization of a (1->3)-β-D-glucan isolated from Saccharomyces cerevisiae. Carbohydr Res 219: 203–13.

26. Muñoz P, Enyele H, Perutz H, McNamara R, Jones E, et al. (1997) The application of various prootic acids in the extraction of [1->3]-β-D-glucan from Saccharomyces cerevisiae. Carbohydr Res 299: 203–8.

27. Shaw AA, Sallam C, Dauphin J-F, Ancian B (1996) Artifact-Free PFG-Enhanced Double-Quantum-Filtered COSY Experiments. J Magn Reson A 120: 110–5.

28. Ancian B, Bourgeois I, Dauphin J-F, Shaw AA (1997) Artifact-Free Pure Absorption PFG-Enhanced DQF-COSY Spectra Including a Gradient Pulse in the Evolution Period. J Magn Reson 125: 548–54.

29. Jeener J, Meier BH, Bachmann P, Ernst RR (1979) Investigation of exchange processes by two-dimensional NMR spectroscopy. J Chem Phys 71: 4546–53.

30. Wagner R, Berger S (1996) Gradient-Selected NOESY–A Fourfold Reduction of the Measurement Time for the NOESY Experiment. J Magn Reson A 123: 119–21.

31. Bax A, Davis DG (1985) MLEV-17-based two-dimensional homonuclear magnetization transfer spectroscopy. J Magn Reson 65: 335–60.

32. Palmer III AG, Cavanagh J, Wright PF, Rance M (1993) Sensitivity improvement in proton-detected two-dimensional heteronuclear correlation NMR spectroscopy. J Magn Reson 93: 131–70.

33. Kay LE, Krüger P, Saarinen T (1992) Pure absorption gradient enhanced heteronuclear single quantum correlation spectroscopy with improved sensitivity. J Am Chem Soc 114: 10663–5.

34. Schleicher J, Schenkerling M, Santler M, Schmidt P, Schledtetzky O, et al. (1994) A general enhancement scheme in heteronuclear multidimensional NMR employing pulsed field gradients. J Biomol NMR 4: 301–6.

35. Norwood NJ, Boyd J, Heritage JE, Soffe N, Campbell ID (1990) Comparison of techniques for 1H-detected heteronuclear 1H-15N Spectroscopy. J Magn Reson 87: 488–501.