The N-terminal Domain of Drosophila Gram-negative Binding Protein 3 (GNBP3) Defines a Novel Family of Fungal Pattern Recognition Receptors

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Gram-negative binding protein 3 (GNBP3), a pattern recognition receptor that circulates in the hemolymph of Drosophila, is responsible for sensing fungal infection and triggering Toll pathway activation. Here, we report that GNBP3 N-terminal domain binds to fungi upon identifying long chains of β-1,3-glucans in the fungal cell wall as a major ligand. Interestingly, this domain fails to interact strongly with short oligosaccharides. The crystal structure of GNBP3-Nter reveals an immunoglobulin-like fold in which the glucan binding site is masked by a loop that is highly conserved among glucan-binding proteins identified in several insect orders. Structure-based mutagenesis experiments reveal an essential role for this occluding loop in discriminating between short and long polysaccharides. The displacement of the occluding loop is necessary for binding and could explain the specificity of the interaction with long chain structured polysaccharides. This represents a novel mechanism for β-glucan recognition.

The activation of the immune response is energetically costly and may be detrimental to the host, especially when inappropriately triggered. Therefore, the reliable detection of infections is a step of paramount importance in the immune response. To achieve the task of detecting potentially hazardous microorganisms, the innate immune system relies on several strategies. One of them is to sense both pathogenic and nonpathogenic microorganisms thanks to pattern recognition receptors (PRRs) that recognize intrinsic microbial molecular "signatures" (1). These immune receptors have been selected during evolution for their ability to bind to essential, conserved, structural components of the microorganisms such as flagellins, peptidoglycans of bacteria, lipopolysaccharides of Gram-negative bacteria, lipoteichoic acids of Gram-positive bacteria, and β-glucans of the fungal cell wall (2, 3). Examples of mammalian PRRs include Toll-like receptors (4), intracellular receptors of the NOD family (5), peptidoglycan recognition proteins (PGRPs) (6), and the membrane-bound Dectin-1 receptor, which detects fungal β-glucans (7).

One important arm of the innate immunity in Drosophila is a potent systemic response that relies on the synthesis in the fat body (a functional equivalent of the mammalian liver) of potent antimicrobial peptides (AMPs) that are secreted in the hemolymph where they attack invading microorganisms. Genetic analysis has delineated two major regulatory pathways of NF-κB type that control the expression of AMP genes (8). The immune deficiency (imd) pathway is mostly required in the host defense against Gram-negative bacteria (9) and is triggered by PRRs of the PGRP family, namely PGRP-SA and PGRP-LE (10). The Toll pathway is essential for fighting fungal and some Gram-positive bacterial infections (12, 13). Toll, the funding member of the Toll-like receptor family, is not itself a PRR. Rather, it is activated by a ligand of the nerve growth factor family, the Spätzle cytokine. To bind to the Toll receptor, Pro-Spätzle needs to be proteolytically processed by a protease, the Spätzle-processing enzyme (SPE) (14), which is itself activated by upstream proteolytic cascades. One such cascade is activated in response to a Gram-positive bacterial challenge by a complex of PGRP-SA, PGRP-SD, and Gram-negative binding protein 1 (GNBP1) (13, 15, 16). Flies deficient for either PGRP-SP or GNBP1 are deficient in Toll pathway activation and are susceptible to infections by several Gram-positive bacterial species but not to fungal infections. In contrast, flies mutant for GNBP3, another gene encoding a GNBP family member, fail to activate the Toll pathway in response to killed fungi and succumb rapidly to fungal but not bacterial infections (17). GNBP3 is thought to activate a proteolytic cascade, which partially overlaps that triggered by the GNBP1-PGRP-SPA complex (18). Even though they belong to the same family and activate the same

The on-line version of this article (available at http://www.jbc.org) contains supplemental Figs. 1–3.

The atomic coordinates and structure factors (code 3IE4) have been deposited in the Protein Data Bank, Research Collaboratory for Structural Bioinformatics, Rutgers University, New Brunswick, NJ (http://www.rcsb.org/).

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4 The abbreviations used are: PRR, pattern recognition receptor; GRP, glucan recognition protein; PGRP, peptidoglycan recognition protein; AMP, antimicrobial peptide; GNBP1, Gram-negative binding protein 1; AS, alkali-soluble; AI, alkali-insoluble; ITC, isothermal titration calorimetry; DP, degree of polymerization; ELISA, enzyme-linked immunosorbent assay; PBS, phosphate-buffered saline; BSA, bovine serum albumin; Nter, N-terminal.
pathway, GNBP1 and GNBP3 are required for sensing distinct classes of microorganisms.

The founding member of the GNBP family, a 50-kDa protein found in hemolymph of *Bombyx mori* and originally named p50, was characterized as a gram-negative (*Escherichia coli*) binding protein (19); hence, its name. However, it has become clear that GNBPs belong to the family of β-glucan recognition proteins (βGRP) that had first been purified on their ability to trigger the prophenol oxidase cascade (a wound response that leads to melanization at the injury site) in response to fungal infections (20). Members of the GNBP/βGRP family are extra-cellular proteins composed of a small N-terminal domain of about 100 residues and a longer C-terminal domain of about 350 residues (21, 22). In the insect *Plodia interpunctella*, both domains of βGRP bind to laminarin, a soluble β-1,3-glucan with a high affinity (*K*ₐ in the 10⁸ M⁻¹ range) (23) which is in the same range as that of the Factor G of the Japanese horseshoe crab (24). The latter factor is used as a diagnostic reagent for the detection of glucans. The C-terminal domain displays sequence similarity to bacterial glucanases, yet the catalytic residues have not been conserved, suggesting that this domain has been selected during evolution for its ability to bind to glucans (21, 22). The N-terminal domain defines a novel β-1,3-glucan binding domain that binds to curdlan, an insoluble linear β-1,3-glucan polymer, a property that the C-terminal glucanase-like domain lacks (21). Full-length recombinant GNBP/βGRPs have been reported to bind to bacteria, lipopolysaccharides, or lipoteichoic acids (19, 22, 23, 25). Although the domain(s) that mediates these interactions has not been thoroughly mapped, it appears that the N-terminal *P. interpunctella* β-1,3-glucan domain is not required for binding to these bacterial compounds (23).

Numerous three-dimensional structures of PGRPs, in some cases complexed with their ligands, have been reported (26–29). In contrast, this knowledge is currently lacking as regarding GNBPs. As a first step toward elucidating the structure/function relationships of GNBPs, we report here that a recombinant GNBP/βGRP has been selected during evolution for its ability to bind to glucans. The C-terminal domain displays β-glucan recognition for 1 h. The solution containing both recombinant protein and yeasts was either centrifuged overnight or washed with PBS solution (30 min at 70 °C and then post-quenched with 0.2 M glycine or treated with 1.5 M NaOH solution twice for 30 min at 70 °C and washed with PBS until neutrality. *p*-Formaldehyde, sodium hydroxide-treated microorganisms, and curdlan beads were used for *in vitro* binding assays of GNBP3-Nter. 1 ml of killed microbes with an *A₅₀₀* of 1 or 50 μg of curdlan beads was added to 5 μg of purified GNBP3-Nter and incubated in 200 μl of binding buffer (10 mM Tris-HCl pH 7.5, 500 mM NaCl) at room temperature with mild agitation for 1 h. The solution containing both recombinant protein and yeasts or curdlan particles was centrifuged (14,000 × *g* for 5 min), and the pellet was washed 3 times with 0.5 ml of washing buffer (10 mM Tris (pH 7.5), 500 mM NaCl, 0.02% Tween 20).

For competition assays (Western blotting coupled with immunodetection), *S. cerevisiae* A1 fraction (100 μg) or curdlan was mixed with 0.5 μg of purified GNBP3-Nter-His tagged alone or pretreated with soluble laminaritetrose/laminarin (400 μg) in a total volume of 50 μl (in 10 mM Hepes (pH 7.5) containing 30 mM NaCl) at 37 °C for 1 h with mild intermittent agitation.

**EXPERIMENTAL PROCEDURES**

**Strains**—The following strains were used in this study: the Gram-positive bacteria *Micrococcus luteus* (CIP A270), the Gram-negative bacteria *E. coli* (1106), the Ascomycetes *Candida albicans* (Ca2.1), *Candida glabrata* (BG2), *Saccharomyces cerevisiae* (BY4741, EUROSCARF, kindly provided by Unité de Génétique Moléculaire des Levures, Institut Pasteur, Paris, France), *Aspergillus fumigatus* (CBS144-89, a clinical isolate), and the Basidiomycetes *Cryptococcus neoformans* (H99).

**Materials**—Curdlan (insoluble β-1,3-glucan, a kind gift from Dr. Hidemitsu Kobayashi), laminarin (rarely branched β-1,3-glucan; Sigma), Pustulan (linear β-1,6-glucan; Calbiochem), and Schizophyllan (a highly β-1,6-branched β-1,3-glucan from Kaken, Japan, kindly provided by Dr Kazutoshi Shibuya) were the polymers used, whereas colloidal chitin was prepared using chitin (Sigma) as described by Gómez Ramírez *et al.* (30). *S. cerevisiae* and *A. fumigatus* cell wall fractions were produced following the protocol of Fontaine *et al.* (31) using actively growing yeast cells and germinating mold conidia in a medium containing 2% glucose and 1% peptone at 37 °C for 15 h. Briefly, cell walls obtained after disruption of the fungal cells was boiled with Tris (50 mM), EDTA (50 mM), SDS (2%), β-mercaptoethanol (40 mM) reagent (pH 7.5) for 15 min, twice, to release cell-wall bound proteins and subsequently treated with 1 M NaOH, 0.5 M NaBH₄ (70 °C, 1 h, twice). The supernatant (alkaline soluble (AS) fraction) obtained after centrifugation was neutralized and dialyzed against water, whereas the sediment (alkali-insoluble (AI) fraction) was washed till neutrality. Both the fractions were freeze-dried and stored at −20 °C. Laminarioligosaccharides with a degree of polymerization of 2–40 (mixture or in their pure form) were prepared according to the method described by Martín-Cuadrado *et al.* (32). Alternatively, laminaritetrose was obtained from Seikagaku. Preparation of curdlan beads was achieved following the protocol described by Ochiai and Ashida (33). All these materials were used for *in vitro* binding and pulldown assays or for direct and competition ELISA assays.

**Expression, Purification, Crystallization, and Mutagenesis**—Starting from the sequence alignment of full-length GNBP3 from the 12 known *Drosophila* genomes (supplemental Fig. 1), the N-terminal domain (that we called GNBP3-Nter) was defined from residues 1 to 128, including the signal peptide (residues 1–25). The protein was successfully expressed in *Drosophila* S2 cells. Details of expression, purification, and crystallization are described elsewhere (51). W77A and short-loop mutants of GNBP3-Nter were prepared using the QuikChange II site-directed mutagenesis kit (Stratagene). The mutation was confirmed by DNA sequencing (MWG).

**Pulldown and Competition Assays**—Overnight cultures of yeasts were collected by centrifugation, washed 3 times with PBS, and resuspended in PBS to an *A₅₀₀* = 1. Yeasts were either fixed with 4% paraformaldehyde overnight at 4 °C and then post-quenched with 0.2 M glycine or treated with 1.5 M NaOH solution twice for 30 min at 70 °C and washed with PBS until neutrality. *p*-Formaldehyde, sodium hydroxide-treated microorganisms, and curdlan beads were used for *in vitro* binding assays of GNBP3-Nter. 1 ml of killed microbes with an *A₅₀₀* of 1 or 50 μg of curdlan beads was added to 5 μg of purified GNBP3-Nter and incubated in 200 μl of binding buffer (10 mM Tris-HCl (pH 7.5), 500 mM NaCl) at room temperature with mild agitation for 1 h. The solution containing both recombinant protein and yeasts or curdlan particles was centrifuged (14,000 × *g* for 5 min), and the pellet was washed 3 times with 0.5 ml of washing buffer (10 mM Tris (pH 7.5), 500 mM NaCl, 0.02% Tween 20).

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Crystal Structure of the N-terminal Domain of GNBP3
In both types of pulldown assays or competition assays, the unbound protein was recovered from the reaction mixture by centrifugation at 3000 × g for 5 min and analyzed by SDS-PAGE (15% gel) either directly or after acetone precipitation. GNBP3-Nter bound to curdlan/Sc-Al fraction was recovered after washing (6×) the centrifugation pellet with 100 μl of 10 mM Hepes containing 150 mM NaCl followed by boiling for 10 min in SDS sample buffer (15 μl). The protein thus released into the supernatant after subsequent centrifugation was analyzed by SDS-PAGE and Western blot using a mouse peroxidase-conjugated mAb-His (Sigma) following the manufacturer’s instructions (Penta-His HRP Conjugate kit, Qiagen) or a rabbit polyclonal anti-GNBP3-Nter antibody as primary antibody.

**Antibody Production**—The purified recombinant GNBP3-Nter protein from S2 cells expression was used to produce polyclonal rabbit antisera. The anti-GNBP3-Nter antisera were screened for specific staining of GNBP3-Nter and *Drosophila* endogenous GNBP3 by Western blot analysis. The specificity of the antibody was assessed by comparing extracts of wild type flies to those of a null GNBP3 mutant strain (data not shown).

**Immunolocalization**—Recombinant His-V5-tagged GNBP3-Nter protein was incubated with paraformaldehyde-treated or NaOH-treated yeast for 1 h at room temperature in binding buffer. After coincubation, the mixture was centrifuged, and the supernatant was aspirated. The pellet was allowed to dry for 2 min. Cells were washed 3 times in washing buffer and blocked with 2% BSA in PBS for 1 h. GNBP3-Nter proteins were detected with a primary mouse anti-V5 antibody (Invitrogen). Primary antibodies were visualized with Cy3-conjugated goat anti-mouse (Zymed Laboratories Inc.). DNA was visualized with 4′,6-diamidino-2-phenylindole. Slides were mounted in Vectashield medium (Vector Laboratories) and were examined by confocal microscopy (Zeiss LSM510). Slides were kept at 4°C, and the images were processed using Adobe PhotoShop CS (Adobe Systems) and analyzed using ImageJ plugin RVB profiler.

**Direct and Competition ELISA Assays**—Fungal cell wall fractions/commercial polymers (200 μg/ml) dispersed by ultrasonication in 50 mM Na2CO3 (pH 9.6) were added (100 μl) to microtiter wells on ELISA plates and incubated overnight at room temperature. Unbound material was removed, and the wells were blocked with 1% BSA and 2% Tween 20 (in PBS) for 1 h at room temperature. His-tagged GNBP3-Nter (0.5 μg/100 μl of binding buffer containing 1% BSA in PBS) was added to each well and incubated at 37°C for 1 h followed by 3 washes with PBS containing 0.5% Tween 20. Peroxidase-conjugated mAb-His (Sigma) (1:10,000 dilution in PBS containing 1% BSA) was added, and the mixture was incubated for 1 h at 37°C. Finally, the reaction was developed in the presence of 0.1 mg/ml O-phenylenediamine (Sigma) and 0.1% H2O2.

For the competition assays, microtiter wells on ELISA plates were coated with the AI fraction (100 μg/ml, 100 μl) *A. fumigatus* or curdlan as described above. At the same time 0.5 μg of GNBP3-Nter-His tagged was incubated with different concentrations of individual laminarioligosaccharides of DP 2–16 or a laminarioligo mixture of DP 12–20 and 20–40 or laminarin in 10 mM Hepes buffer (pH 7.0) in a total volume of 50 μl for 1 h at room temperature, after which 50 μl of PBS containing 2% BSA was added to all the tubes. Then these mixtures were added to each well, and ELISA readings were performed as described above. Statistical analyses were done on GRAPHPAD PRISM using the Student-Newman-Keuls test.

**Isothermal Titration Calorimetry (ITC)**—ITC experiments were performed using an iTC200 Isothermal Titration Calorimetry system (MicroCal; Northampton, MA) at a temperature of 30°C. A typical titration profile is shown in Fig. 3. Protein and sugar samples were prepared in 20 mM Hepes and 150 mM NaCl (pH 7.5). Protein solution was taken in a syringe and loaded into the ITC sample cell (cell volume 200 μl). After the base line stabilized, 20 injections of 2 μl of the sugar ligand solution were added from the computer-controlled syringe into the protein solution, and exothermic heat changes accompanying the additions were recorded. The time period between the two consecutive injections was fixed at 340 s to allow the exothermic peak to return to the base line. The heat of mixing was measured by making identical injections into the cell containing buffer with no protein. The experimental data were fitted using software ORIGIN 7 supplied by Microcal, with ΔH (enthalpy change in kcal mol−1), K_d (association constant in M−1), and n (number of binding sites/monomer) as adjustable parameters. Other thermodynamic parameters were calculated using the standard equation, ΔG = ΔH − TΔS = −RT log K_d, where ΔG, ΔH, and ΔS are the changes in free energy, enthalpy, and entropy of binding, respectively. T is the absolute temperature in Kelvin, and R = 1.98 cal mol−1K−1.

**Structure Determination**—The structure of GNBP3-Nter was determined by the single wavelength anomalous dispersion method using a samarium derivative. Diffraction data for the samarium derivative were collected at 100 K on a 300-mm Macresearch imaging plate mounted on a Rigaku RU200 rotating anode. They were indexed, integrated, and scaled using the XDS package (34). The space group was C2 with two molecules per asymmetric unit. Samarium sites were identified, refined, and used for phase calculation with the PHENIX suite (35). An initial model was then auto-built with PHENIX in which 69% of the total amount of residues was built. At this stage, the R value was 34%, and the R_free value was 38%. This model was then refined against a high resolution diffraction data set (1.45 Å) collected on beamline ID23-1 at the European Synchrotron Radiation Facility (Grenoble, France). Refinement was performed using REFMAC5 (36), and manual rebuilding was carried out with the programs Coot (37) and Turbo-Frodo (38). The models of GNBP3-Nter lack interpretable electron density for the last residues 102–107. The final crystallographic model was refined to R and R_free values of 16.5 and 19.8%, respectively. Statistics for all the data collections and refinement are summarized in Table 1. Figs. 4 and 5 were generated with PyMOL.

**RESULTS**

The boundaries of GNBP3-Nter were delineated using an alignment of GNBP3 sequences from the genomes of 12 *Drosophila* species, as depicted in supplemental Fig. 1. The recombinant protein was overexpressed at a high level (>15 mg/liter of culture) in *Drosophila* S2 cells with the C-terminal extension V5-His6, which was used for detection and purification. The tag
was proteolytically removed to allow crystallization of the recombinant protein (51).

**GNBP3-Nter Binds to the Fungal Cell Wall**—To determine whether the recombinant N-terminal domain of GNBP3 is able to bind to fungi, we first analyzed by pulldown experiments its binding to *C. albicans*, *C. glabrata*, and *C. neoformans* yeasts. We detected a mild binding to *p*-formaldehyde-fixed *Candida* yeasts and a strong binding to NaOH-treated *Candida* yeasts using either a tagged or a cleaved tag form of the recombinant protein (Figs. 1, A–C). The NaOH treatment strips the cell wall of its proteins and alkali-soluble polysaccharides, thus making the β-1,3-glucans more accessible. We did not, however, detect any binding to *C. neoformans* or bacteria, which have no β-glucan on their surface. We confirmed by immunohistochemistry the binding of the recombinant protein to *C. albicans* (Fig. 1D) and *C. glabrata* (data not shown). We found that the recombinant protein binds to discrete patches of the yeasts. Staining appeared strong in newly formed buds and bud scars. In contrast, GNBP3-Nter bound to the entire surface of NaOH-treated *Candida* yeasts (Fig. 1E).

**GNBP3-Nter Binds Specifically to β-1,3-Glucans**—From the preceding experiments, we deduced that GNBP3-Nter binds to the fungal cell wall. However, as the latter is mainly a complex network of different polysaccharides, we performed binding assays on ELISA plates coated with different cell wall fractions or with commercially available polysaccharides. As depicted in Fig. 2A, GNBP3-Nter efficiently bound to the cell wall alkali-insoluble (AI) fraction from *S. cerevisiae* and *A. fumigatus* but not to the alkali-soluble (AS) fractions of *A. fumigatus*, which lacks β-1,3-glucans and contains mainly α-1,3-glucan and galactomannan. The structure common to the AI fraction of *S. cerevisiae* and *A. fumigatus* is a β-1,6-branched β-1,3-glucan covalently bound to chitin, suggesting that the polysaccharide recognized by GNBP3-Nter was either a β-1,3-glucan or chitin. However, we did not observe any binding with chitin, a linear polymer of N-acetylglucosamine. The binding efficacy to schizophyllan, a β-1,3-glucan with single β-(1,6)-bonded glucose at every third glucose molecule on the main chain, was less than 10% compared with the *S. cerevisiae* AI fraction. Also, there was no binding to pustulan, a linear β-1,6-glucan polymer. The highest ELISA values were obtained for curdlan, an insoluble linear β-1,3-glucan. Taken together, the binding assays indicate that GNBP3-Nter shows specific affinity toward β-1,3-glucan.

**GNBP3-Nter Binding to β-Glucans Increases with Polysaccharide Chain Length**—Competition assays for binding to GNBP3-Nter were performed between the cell wall AI fraction from *A. fumigatus* and soluble β-1,3-glucan oligosaccharides of different sizes (individually or in a mixture). After preincubation of GNBP3-Nter with laminarioligosaccharides of varying length (degree of polymerization (DP) of 2–16), there was weak or no reduction in the binding of GNBP3-Nter to the wells on the ELISA plates coated with the AI-fraction even when GNBP3-Nter was preincubated with laminarioligosaccharides

### Crystal Structure of the N-terminal Domain of GNBP3

**TABLE 1**

| Data collection and refinement statistics | GNBP3-Nter Sm | GNBP3-Nter native |
|------------------------------------------|---------------|-------------------|
| Data collection statistics              |               |                   |
| Radiation source                         | In-house      | ESRF ID23-1       |
| Wavelength (Å)                           | 1.5418        | 1.0332            |
| Space group                              | C2            | C2                |
| Cell dimensions                          |               |                   |
| a, b, c (Å)                              | 135.32, 30.80, 51.72 | 135.53, 30.68, 51.65 |
| β (°)                                    | 107.25        | 107.57            |
| Resolution range (Å)                     | 46.52-2.20 (2.32-2.20) | 46.52-1.45 (1.52-1.45) |
| Total observations                       | 132,581       | 129,875           |
| Unique reflections                       | 10,736        | 36,216            |
| Completeness (%)                         | 99.8 (99.0)   | 98.8 (95.6)       |
| Redundancy                               | 12.3 (11.9)   | 3.6 (3.5)         |
| Rmerge (°)                               | 3.3 (8.6)     | 6.7 (29.6)        |
| Average I/σ(I)                           | 53.8 (30.1)   | 12.2 (4.1)        |

**Refinement and model statistics**

| Resolution range (Å)                     | 29.85-1.45 |
| Proteins per asymmetric unit             | 2          |
| Number of reflections used               | 34,308     |
| Rwork (%) and Rmerge (%)                 | 16.5/19.8  |
| Average B values                         |            |
| All atoms (Å2)                           |            |
| Protein atoms (Å2)                       | 14.94      |
| Ethylene glycol atoms (Å2)               | 13.28      |
| Zinc atoms (Å2)                          | 35.87      |
| Water atoms (Å2)                         | 4.29       |
| Root mean square deviation from ideality |            |
| Bond lengths (Å)                         | 0.012      |
| Bond angles (°)                          | 1.433      |
| Torsion angles (°)                       | 6.756      |
| Ramachandran analysis, favored regions/allowed regions/outliers (% of residues) | 99/1/0 |
| No. of atoms                             |            |
| Protein                                 | 1,612      |
| Ethylene glycol                          | 8          |
| Zinc                                     | 6          |
| Water                                    | 192        |

& Rmerge = ΣhΣi|Fo| − |Fc|/ΣhΣi|Fo|, where |Fo| and |Fc| are the observed and calculated structure factor amplitudes, respectively, for reflection h.

b Rwork = ΣhΣi[(|Fo| − |Fc|)^2]/ΣhΣi|Fo|^2, where |Fo| and |Fc| are the observed and calculated structure factor amplitudes, respectively, for reflection h.

c Rfree is the R value for a subset of 5% of the reflection data, which were not included in the crystallographic refinement.

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Crystal Structure of the N-terminal Domain of GNBP3

The crystal structure of the N-terminal domain of GNBP3 was determined by X-ray crystallography. The structure consists of two antiparallel sheets and belongs to the immunoglobulin fold family. The first sheet is made of strands A (residues 7–10), B (residues 17–21), and E (residues 73–82), and the second sheet is made of strands C (residues 85–88), D (residues 26–35), G1 (residues 85–88), and G2 (residues 92–95). The protein is monomeric in solution up to a concentration of 0.5 mM as analyzed by gel filtration and dynamic light scattering, suggesting little functional significance for the crystallographic dimer.

The final refined model consists of residues 26–128 that contain two copies of the protein in the asymmetric unit. The two molecules were refined independently, and there are no significant differences (root mean square deviation = 0.53 Å for all Cαs). The protein is monomeric in solution up to a concentration of 0.5 mM as analyzed by gel filtration and dynamic light scattering, suggesting little functional significance for the crystallographic dimer.

The structure of GNBP3-Nter was solved at 1.45 Å of resolution by single anomalous dispersion using a samarium derivative. The crystals at a 1/800 mass ratio (Fig. 2B). In contrast, water-soluble laminariligosaccharide mixtures of higher DP (12–20, with maximum concentration of DP 14–18) and laminarin (a mixture of oligosaccharides of DP between 20 and 28 with 25-DP oligomer in the maximum concentration and having one branching point per oligosaccharide chain) competed efficiently with the AI fraction for the binding site(s) on GNBP3-Nter (Fig. 2B). However, the best inhibition was observed with a mixture of oligosaccharides of DP between 20 and 40 that was water-insoluble (Fig. 2B). These ELISA inhibition data were further confirmed by competition between laminaritetrose or laminarin and the AI fraction from _S. cerevisiae_ for GNBP3-Nter binding using pulldown assays coupled with blotting immunodetection analyses (Fig. 2C). β-1,3-Linked tetraoses did not compete with the _Sc-Ai_ fraction for GNBP3-Nter binding, even at a protein/sugar molar ratio of 1/500, whereas laminarin reduced significantly the binding of GNBP3-Nter to _Sc-Ai_ fraction when used at a protein/sugar ratio of 1/10 (Fig. 2C).

The stoichiometry for the interaction between GNBP3-Nter and laminarin was close to a ratio of 1–3 (n = 2.51), consistent with a triple helix organization of the laminarin in solution (39). The binding affinity is _K_A_ = 2.12 × 10^6 ± 0.4 × 10^6 M^-1. The fitted data also yielded the interaction with negative enthalpy (∆H = −3.34 kcal/mol) and entropy (∆S = 17.9 kcal/mole/degree). In contrast and consistent with the competition assays, no interaction was detected between GNBP3-Nter and shorter sugars such as heptaose (DP7) or hexaose (DP6) even with repeated injections (2 μl) of highly concentrated sugar (10 mM) in the cell (data not shown). Taken together, these data indicate that GNBP3-Nter binds specifically to linear β-1,3-glucans with high DP.

Overall Structure of GNBP3-Nter—

The structure of GNBP3-Nter was solved at 1.45 Å of resolution by single anomalous dispersion using a samarium derivative. The crystals...
sheets are packed in a β-sandwich conformation enclosing a highly hydrophobic core organized around a cluster of three phenylalanines (Phe-16, Phe-31, and Phe-61). The closest structural homologue found using the DALI server (40) is a fibronectin type III domain of integrin α6β4 (PDB code 1IQG3) (41) with a Z-score of 7.9. This molecule displays the same β-sheet organization, i.e. A-B-E and C-C-F-G1-G2. Despite a very low level of sequence identity (9%), superimposition of the fibronectin III domain with GNBP3-Nter shows that 66 residues of 102 are structurally conserved, giving a root mean square deviation value of 1.54 Å. The main difference is the presence in GNBP3-Nter of a large negatively charged loop between strands C and C that folds back onto the β-sheet C-C-F-G1-G2 (Fig. 4C). Interestingly, GNBP3-Nter also displays the same β-sheet organization as starch binding domains (42).

**Carbohydrate Binding Site**—Based on their binding characteristics, carbohydrate binding modules have been classified into three types named A (“surface binding,” for insoluble polysaccharides), B (“glycan chain binding,” which involves a groove), and C (“small sugar binding”) (43). The functional studies in this report show the preferential binding of GNBP3-Nter to long chain soluble or insoluble -glucans, thus classifying GNBP3-Nter either as a type A or as a type B carbohydrate binding module. Higher affinity toward curdlan/cell wall AI fractions compared with soluble short chain sugars and the absence of any groove containing aromatic residues on its surface.
The three Trp and the eight Tyr residues of GNBP3-Nter were, therefore, carefully examined (Fig. 5, A and B). Trp-47 participates in the previously described hydrophobic core located between the two β sheets. It makes van der Waals contacts with Phe-61, Phe-16, and Leu-35 and is partially buried by Thr-46. Trp-59 stands in a hydrophobic pocket and contacts Leu-26 (CG2), Ile-51 (CG2), Ala-54 (CB), Phe-29, and Phe-31. Thus, it is even more buried than Trp-47. The indole ring of Trp-77 is in stacking interaction with the His-32 imidazole group that lies beneath it. Trp-77 is only poorly accessible, as it is masked from the surface by Leu-42, which stands at the tip of the C-C loop. Five of the eight Tyr residues are distributed into two groups located at the two ends of the molecule. On one side Tyr-12 and Tyr-99 stand close to each other but are not stacked. They are accessible, with their OH groups pointing toward the surface of the molecule. On the opposite side, a stacking interaction occurs between Tyr-1 and Tyr-82. A third tyrosine, Tyr-87, positions its ring 90° from those of Tyr-1 and Tyr-82, leading to the formation of an imperfect aromatic cage. Tyr-76 is completely buried inside the molecule, whereas the Tyr-75 residue is masked from the surface by Glu-40 on the C-C loop. Finally, only Tyr-79 is fully solvent-exposed. This latter residue stands on the strand F, which is central to the β-sheet C’-C-F-G1-G2. Interestingly, strand F possesses two other aromatic residues, Tyr-75 and Trp-77, which are strictly conserved among β-glucan recognition domains. The spatial arrangement of the three aromatic side chains of Tyr-75, Trp-77, and Tyr-79 (Fig. 5C) is similar to the aromatic patch constituted by three neighboring residues described for starch binding domains, for example those of the starch recognition domain of the pullulanase PulA from Thermotoga maritima (44)(Fig. 5D). Nevertheless, the exposure of Tyr-75 and Trp-77 side chains to the surface is masked by the C-C’ loop. When this C-C’ loop is removed from GNBP3-Nter using a graphics dis-
play system, Tyr-75 and Trp-77 become accessible to solvent (Fig. 5C). Thus, we hypothesized that this loop acts as a mobile lid domain that covers the putative binding site (Fig. 5E). To verify our hypothesis, we mutated Trp-77 into Ala. Structural integrity of W77A mutant was assessed by circular dichroism spectroscopy. No structural difference between wild type and mutant proteins was detected (supplemental Fig. 3B). W77A mutant was expected to display a decreased affinity for β-1,3-glucan. Indeed, the binding to curdlan in a pulldown assay was severely decreased with the mutant protein (Fig. 5F). Moreover, in ITC experiments we failed to detect any binding of either long or short laminarioligosaccharides to the W77A mutant (data not shown), thus delineating a key role for Trp-77 in β-1,3-glucan recognition.

The C-C’ Loop of GNBP3-Nter—An unusual feature of GNBP3-Nter structure is the presence of a long loop between strands C and C’, which is composed of 10 residues and extends outwards from the compact body. Insertions between strands C and C’ have already been described for other members of the fibronectin type III-fold family and were assigned to be protein-protein interaction domains, as observed, for example, for fibronectin type III oδβ4 integrin (41). The C-C’ loop of GNBP3-Nter is quite well conserved in terms of length and sequence among the N terminus domains of GNBP/βGRPs that have been shown to bind to β-glucans (Fig. 4A). The strict conservation of seven positions often gives the following consensus motif 36NEEMGXEXG45. GNBP3-Nter carries four negatively charged amino acids (Glu-37, Glu-38, Glu-40, and Glu-43), which point outward from the surface (Fig. 5E). Two of them, Glu-37 and Glu-43, interact through side chain-side chain hydrogen bonds with the conserved Lys-34 residue. The NE2 nitrogen of Trp-77 interacts on the C-C’ loop. Finally, the side chains of the strictly conserved Met-39 and of Leu-42 contribute to the formation of a hydrophobic environment together with Tyr-75 and Trp-77 on the internal face of the C-C’ loop. As all these residues have been conserved throughout 350 million years of evolution (divergence of the Diptera and Lepidoptera lineages occurred during the early Carboniferous) (45), it is likely that these interactions have been selected to maintain the lid in a closed position in the absence of glucan ligands. A mutant protein in which the occluding loop...
was shortened and the conserved residues were mutated was cloned and produced in *Drosophila* cells (supplemental Fig. 3A).

The structural integrity of the mutated protein was assessed by circular dichroism (supplemental Fig. 3B). The binding of the short-loop mutant to killed *Candida* yeasts was not detectable by immunohistochemistry. ELISA assays showed that the binding to the A1 fraction of *S. cerevisiae* and curdlan was substantially reduced and corresponded to 30 and 10% that of the wild type, respectively (data not shown).

**DISCUSSION**

The discrimination between host and microbe-associated molecules is crucial to the function of PRRs. Short oligosaccharide chains may not constitute an ideal target for PRRs as they might also be displayed by host cells and, thus, may not represent a *bona fide* microbial signature. Therefore, it is likely that the host selected PRRs able to sense long glucan chains idiosyncratic to most fungal cell walls. In this manuscript we report that the glucan binding domain of GNBP3 binds preferentially to long β-1,3-glucan chains and shall discuss how the distinction between short and long chains of glucans is made by various PRRs.

The N-terminal domain of GNBP3 binds to the cell wall of *C. albicans* and most likely to β-glucans as indicated by the preferential binding to growing cell buds and bud scars, a pattern evocative of that of Dectin-1 (46). Indeed, the recombinant protein binds to the cell wall alkali-insoluble polysaccharide fraction of *S. cerevisiae* and *A. fumigatus*. The latter induces a GNBP3-dependent activation of the Toll pathway when injected into *Drosophila* (17). Our data indicate that the relevant biochemical moiety of these fungal cell wall AI extracts are β-1,3-glucan chains. These findings are confirmed by direct binding of GNBP3-Nter to curdlan and laminarin as assayed by ELISA, ITC, and pulldown experiments coupled to competition assays. The longer the glucan chain, the more efficient is the competition. Efficient binding to GNBP3-Nter is observed with polymeric chains that incorporate more than 16 glucan units. In keeping with this result, it has previously been shown that injection in *Drosophila* of the alkali-insoluble fraction of the *A. fumigatus* cell wall, which consists of long polysaccharides including β-1,3-glucans, induces a strong activation of the Toll pathway (17). At the same time, Gottar and colleagues in Strasbourg found that short laminarioligosaccharides with a DP ranging from 2 to 7 failed to induce Toll pathway activation when injected into flies. 5 β-1,6-Branching in the linear chain of β-1,3-glucans does not appear to be required for recognition by GNBP3-Nter as we failed to observe strong binding with schizophyllan, a highly β-1,6-branched β-1,3-glucan from *Schizophyllum commune* (Fig. 2A). Interestingly, the glucan binding properties of the mammalian fungal receptor Dectin-1 have been reported to be fairly similar, with a minimum degree of polymerization of 11 required for β-glucan binding (47).

We have solved the GNBP3-Nter crystal structure that provides structural insight into the β-1,3-glucan recognition protein (βGRP) family. The overall structure displays an immuno-globulin-like fold similar to that of the fibronectin III superfamily. Although no solvent-exposed aromatic patch is present on GNBP3-Nter (Fig. 5, A and B), Tyr-75, Trp-77, and Tyr-79 are good candidates to constitute such a binding platform (Fig. 5C) after a structural rearrangement of the loop located between strands C and C’. In keeping with this hypothesis, we found that the binding to curdlan was strongly impaired, and the binding to laminarin was completely abolished with the W77A mutant, thus underscoring the importance of this initially buried residue for glucan binding. The essential role of the C-C’ loop in terms of binding and discrimination between short and long chains of β-glucan was confirmed by mutagenesis. To free access to the binding site, the C-C’ loop should fold back toward the C-terminal domain of GNBP3 (Fig. 5, C and E). Tyr-79 may act as a primary determinant that anchors β-glucan polymers. Then the negatively charged patch formed by the four glutamic acids on the top of the loop may be expelled by the vicinity of a large sugar surface, unmasking the rest of the binding site (Tyr-75 and Trp-77).

After the lid opening, the side chain of Tyr-75 is free to re-orientate toward Trp-77. Like this, the relative spacing between the three residues would not stretch beyond a distance required to accommodate a disaccharide and, thus, would be very comparable with that of starch binding domains (Fig. 5D). The internal hydrophobic surface of the lid could hardly be fully-exposed to solvent in the open conformation and may probably interact with the ligand. This putative interaction between the lid and the ligand may explain the results obtained for the short-loop mutant. Namely, this mutant does not appear to bind efficiently to long-chain oligosaccharides, possibly because the two conserved hydrophobic residues in the lid, which are missing in the mutant, no longer stabilize the interaction.

Both the sequence of the C-C’ loop and that of the putative binding site are conserved in βGRP family members that have been reported to bind to β-glucans. Noticeably, these sequences are not conserved in *Drosophila melanogaster* GNBP1 (and GNBP1 of other *Drosophila* species), a member of the family required in the host defense against Gram-positive bacteria (supplemental Fig. 2). We infer that the GNBP1 N-terminal domain will not bind significantly to β-1,3-glucans, even though some studies have reported some binding of full-length GNBP1 to curdlan (25). Thus, the sequences of the C-C’ loop and of the glucan binding site may be useful predictors of the function of uncharacterized GNBP/βGRP family members.

Using this criterion, we predict that the function of the funding member of the GNBP family, *B. mori* p50, is not involved in defense against fungi, at least by a GNBP3/βGRP-like mechanism. In any case, GNBP full-length proteins, with their glucanase-like domains, are likely to have emergent properties not displayed by the Nter domain alone. These may include the activation of downstream proteolytic cascades (for Toll pathway and prophenol oxidase activation) and, obviously, agglutination, which requires two sugar binding domains in the protein.

An intriguing feature of the GNBP3-Nter domain is its capacity to discriminate between short and long chains of β-glucans. Many PRRs activate an immune response only

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5 M. Gottar, personal communication.
when bound to long chains of carbohydrates through the use of spatially arranged multiple subunits or multimers. Yet, in striking contrast to GNBP3, the individual domains involved in carbohydrate recognition can bind to monomeric or short carbohydrate polymers. For instance, PGRP-SA, which binds to PGN muropeptide monomers as single molecules, requires the formation of PGRP-SA clusters on longer chains to trigger downstream proteolytic cascades (48). A similar case is presented by Factor G of the Japanese horseshoe crab where a laminariheptaose is required to activate the coagulation cascade even though it binds well to laminaribiose with an affinity that is only three times lower (49). Interestingly, the recognition domain that binds to β-glucans is actually made up of two carbohydrate binding subunits arranged in a tandem repeat. Only the tandem repeat, and not each individual subunit, is able to bind to the disaccharide. Another example of the importance of the spatial arrangements of multiple carbohydrate recognition domain (CRD) is provided by the mannose-binding lectin whereby each CRD head binds to a single sugar residue (mannose or fucose). Activation only occurs when the multiple heads arranged in a bouquet-like structure of trimers bind to an array of sugar residues present on the microbial but not the host cell surface (50). Here, we propose that the lid of the array of sugar residues present on the microbial but not the host cell surface (50). Here, we propose that the lid of the carbohydrate binding site is displaced only by long chains of β-glucans. In this respect, the structures of laminarins and curdlan as triple helices in an aqueous environment may be an essential feature that triggers the opening of the carbohydrate binding site and the recognition of the fibrillar fungal structure.

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REFERENCES

1. Janeway, C. A. (1989) Cold Spring Harbor Symp. Quant. Biol. 54, 1–13
2. Akira, S., Uematsu, S., and Takeuchi, O. (2006) Cell 124, 783–801
3. Medzhitov, R., and Janeway, C. A., Jr. (2002) Science 296, 298–300
4. Takeda, K., Kaisho, T., and Akira, S. (2003) Annu. Rev. Immunol. 21, 335–376
5. Inohara, N., and Núñez, G. (2003) Nat. Rev. Immunol. 3, 371–382
6. Royet, J., and Dziarski, R. (2007) J. Immunol. 178, 264–277
7. Brown, G. D., Taylor, P. R., Reid, D. M., Willment, J. A., Williams, D. L., Geiss, J., and Willment, J. A. (1996) Proc. Natl. Acad. Sci. U.S.A. 93, 7888–7893
8. Yoshida, H., Ochiai, M., and Ashida, M. (1986) Biochem. Biophys. Res. Commun. 141, 1177–1184
9. Ochiai, M., and Ashida, M. (2000) J. Biol. Chem. 275, 4995–5002
10. Ma, C., and Kanost, M. R. (2000) J. Biol. Chem. 275, 7505–7514
11. Fabrick, J. A., Baker, J. E., and Kanost, M. R. (2004) J. Biol. Chem. 279, 26605–26611
12. Muta, T., Seki, N., Takaki, Y., Hashimoto, R., Oda, T., Iwanaga, A., Toku-naga, F., and Iwanaga, S. (1995) J. Biol. Chem. 270, 892–897
13. Lee, S. H., Carpenter, J. F., Chang, B. S., Randolph, T. W., and Kim, Y. S. (2006) Protein Sci. 15, 304–313
14. Kim, M. S., Byun, M., and Oh, B. H. (2003) Nat. Immunol. 4, 787–793
15. Chang, C. I., Pili-Floury, S., Hervé, M., Paquet, C., Chelliah, Y., Lemaître, B., Mengine-Leclercx, D., and Deisenhofer, J. (2004) PLoS Biol. 2, E277
16. Chang, C. I., Chelliah, Y., Borek, D., Mingen-Leclercx, D., and Deisenhofer, J. (2006) Science 311, 1761–1764
17. Leone, P., Bischoff, V., Kellenberger, C., Hettu, C., Royet, J., and Roussel, A. (2008) Mol. Immunol. 45, 2521–2530
18. Gómez Ramírez, M., Rojas Aveilza, L. I., Rojas Aveilza, N. G., and Cruz Cammilo, R. (2004) J. Microbiol. Methods 56, 213–229
19. Fontaine, T., Simonel, C., Dubreucq, G., Adam, O., Delepiègre, M., Lemaine, J., Vorgias, C. E., Diaquin, M., and Latgé, J. P. (2000) J. Biol. Chem. 275, 27594–27607
20. Martin-Cuadrado, A. B., Encinar del Dedo, J., de Medina-Redondo, M., Fontaine, T., del Rey, F., Latgé, J. P., and Vázquez de Aldana, C. R. (2008) Mol. Microbiol. 69, 188–200
21. Ochiai, M., and Ashida, M. (1988) J. Biol. Chem. 263, 12056–12062
22. Kabsch, W. (1993) J. Appl. Crystallogr. 26, 795–800
23. Adams, P. D., Grosse-Kunstleve, R. W., Hung, L. W., Ioerger, T. R., Mcenguy, K., and Terwilliger, T. C. (2002) Acta Crystallogr. D Biol. Crystallogr. 58, 1948–1954
24. Murshudov, G. N., Vagin, A. A., and Dodson, E. J. (1997) Acta Crystallogr. D Biol. Crystallogr. 53, 240–255
25. Emsley, P., and Cowtan, K. (2004) Acta Crystallogr. D Biol. Crystallogr. 60, 2126–2132
26. Roussel, A., and Cambillau, C. (1991) in Silicon Graphics Geometry Partners Directory, Silicon Graphics, Mountain View, CA
27. Okobita, T., Miyoshi, K., Uezu, K., Sakurai, K., and Shinkai, S. (2008) Bioinformatics 24, 783–788
28. Holm, L., Kääriäinen, S., Rosenström, P., and Snelben, A. (2008) Bioinformatics 24, 2780–2781
29. Lee, E., Shin, C., and Lee, J. (2007) Cell 129, 1277–1286
30. Palma, A. S., Feizi, T., Zhang, Y., Stoll, M. S., Lawson, A. M., Diao-
Rodriguez, E., Campanero-Rhodes, M. A., Costa, J., Gordon, S., Brown, G. D., and Chai, W. (2006) J. Biol. Chem. 281, 5771–5779

48. Park, J. W., Kim, C. H., Kim, J. H., Je, B. R., Roh, K. B., Kim, S. J., Lee, H. H., Ryu, J. H., Lim, J. H., Oh, B. H., Lee, W. J., Ha, N. C., and Lee, B. L. (2007) Proc. Natl. Acad. Sci. U.S.A. 104, 6602–6607

49. Takaki, Y., Seki, N., Kawabata Si, S., Iwanaga, S., and Muta, T. (2002) J. Biol. Chem. 277, 14281–14287

50. Hoffmann, J. A., Kafatos, F. C., Janeway, C. A., and Ezekowitz, R. A. (1999) Science 284, 1313–1318

51. Mishima, Y., Coste, F., Bobezeau, V., Hervouet, N., Kellenberger, C., and Roussel, A. (2009) Acta Crystallogr. Sect. F Struct. Biol. Cryst. Commun. 65, 870–873