**Structural and Functional Analysis of PGRP-LC Indicates Exclusive Dap-Type PGN Binding in Bumblebees**

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**Abstract:** Peptidoglycan recognition proteins (PGRPs) play an important role in the defense against invading microbes via the recognition of the immunogenic substance peptidoglycan (PGN). Bees possess fewer PGRPs than *Drosophila melanogaster* and *Anopheles gambiae* but retain two important immune pathways, the Toll pathway and the Imd pathway, which can be triggered by the recognition of Dap-type PGN by PGRP-LCx with the assistance of PGRP-LCa in *Drosophila*. There are three isoforms of PGRP-LC including PGRP-LCx, PGRP-LCa and PGRP-LCy in *Drosophila*. Our previous study showed that a single PGRP-LC exists in bumblebees. In this present study, we prove that the bumblebee *Bombus lantschouensis* PGRP-LC (Bl-PGRP-LC) can respond to an infection with Gram-negative bacterium *Escherichia coli* through binding to the Dap-type PGNs directly, and that *E. coli* infection induces the quick and strong upregulation of PGRP-LC, abaecin and defensin. Moreover, the BI-PGRP-LC exhibits a very strong affinity for the Dap-type PGN, much stronger than the affinity exhibited by the PGRP-LC from the more eusocial honeybee *Apis mellifera* (Am-PGRP-LC). In addition, mutagenesis experiments showed that the residue His³⁹⁰ is the anchor residue for the binding to the Dap-type PGN and forms a hydrogen bond with MurNAc rather than meso-Dap, which interacts with the anchor residue Arg⁴¹³ of PGRP-LCx in *Drosophila*. Therefore, bumblebee PGRP-LC possesses exclusive characteristics for the immune response among insect PGRPs.

**Keywords:** PGRP-LC; bumblebee; *Bombus lantschouensis*; innate immune; Dap-type PGNs

1. Introduction

Innate immunity is an important line of defense that protects insects from pathogen invasion. It can be activated by interactions between germline-encoded pattern recognition receptors (PRRs) and pathogen-associated molecular patterns (PAMPs) [1–4]. As an important type of PRR, peptidoglycan recognition proteins (PGRPs) were first discovered in the hemolymph of silkworms and participate in an antimicrobial host defense mechanism [5]. PGRPs have been shown to be conserved from mammals to insects, except in lower metazoans (nematodes) and plants that do not have PGRPs [6].

Nevertheless, insects possess more diverse PGRPs than mammals; for instance, *Drosophila melanogaster* has 13 PGRP genes and *Anopheles gambiae* possesses seven PGRP genes [6]. Based on their length, insect PGRPs can be divided into two classes: short PGRPs (PGRP-S) and long PGRPs (PGRP-L). Each PGRP contains a single peptidoglycan (PGN)-binding domain. These PGRPs recognize the bacterial or fungal PGN and then trigger two important immune pathways, the Toll
pathway and the Imd pathway, which activate the production and release of antimicrobial peptides against the pathogenic infection [4,7,8]. On the basis of the components, PGNs can be classified into two major categories: L-lysine type (Lys-type) PGNs which mainly exist in Gram-positive cocci, and meso-diaminopimelic acid (Dap)-type PGNs which are present in Gram-negative bacteria and several Gram-positive bacilli [9]. The Drosophila PGRP-LB structure was the first for which the PGRP-binding groove was characterized [10]. Subsequently, six other Drosophila PGRP structures were determined, including the structures of PGRP-SA [11,12], PGRP-SD [13], PGRP-LCa [14], PGRP-LE [15], PGRP-LF [16] and the PGRP-LCx/PGRP-LCa complex with tracheal cytotoxin (TCT), a naturally occurring fragment of the Gram-negative PGN [17]. Notably, the PGRP-LCx/PGRP-LCa/TCT complex structure was the first to show details of the interaction between PGRP-LC and Dap-type PGNs at the atomic level. The complex structure shows that Arg\(^{413}\) creates a positively charged patch, which is consistent with the preferential binding of PGRP-LCx to a Dap-type PGN. The residues Trp\(^{394}\), Asp\(^{395}\), His\(^{365}\), Glu\(^{480}\), Ser\(^{477}\), Ala\(^{478}\) and Thr\(^{479}\) were also shown to be essential for the ability to bind to Dap-type PGNs. Drosophila PGRP-LCx requires the cooperation of PGRP-LCa to bind to Dap-type PGNs to activate the Imd immune pathway [17]. Other studies have shown that Drosophila PGRP-LC binds to Dap-type PGNs in cooperation with PGRP-LE and triggers the Imd pathway to activate the antimicrobial peptide production and release [18].

As a group of important pollinators, bumblebees contribute greatly to the pollination of wild flowers and crops and play a key role in the maintenance of natural and agricultural ecosystems [19–24]. However, bumblebee colonies are threatened by various pathogens, which have coevolved with their immune systems for many years [25–30]. Bumblebees belong to the Hymenoptera and are members of a more ancient clade than that of Drosophila (Diptera) [31]. In addition, bumblebees possess a relatively low number of immune-related genes including PGRPs, which are involved in the Toll pathway and the Imd pathway [32]. Bumblebee and honeybee PGRP-S genes have been shown to be upregulated under challenge with bacteria or the antigenic components of bacteria [33,34]. In contrast to Drosophila PGRP-SA, the PGRP-SA proteins of the bumblebee Bombus ignitus and the honeybee Apis mellifera preferentially bind to Lys-type PGNs over Dap-type PGNs and can bind to Dap-type PGNs with high affinity [35,36]. Interestingly, unlike Drosophila PGRP-LE, which has three isoforms, namely, PGRP-LCa, PGRP-LCx and PGRP-LCy, the bumblebee B. lantschouensis PGRP-LE exists as only one isoform, which is highly conserved among different bumblebee species [37]. Nevertheless, the structural and functional details of the PGN-binding activity of PGRP-LE in bees remain unclear, impeding our understanding of the immune role of PGRP-LE in the pathogen defense of bees. In this current study, the structural and functional characteristics of B. lantschouensis PGRP-LE were investigated to fill this knowledge gap regarding PGRP-LE in bees.

2. Results

2.1. Response of Bl-PGPR-LE to Infection with Gram-Negative and Gram-Positive Bacteria

The present study showed that both the Gram-positive bacterium Staphylococcus aureus and the Gram-negative bacterium Escherichia coli can trigger the upregulation of PGRPR-LE and the antimicrobial peptides abaecin, defensin and hymenoptaecin (Figure 1). However, over time, the degree of response differs between S. aureus and E. coli. After E. coli infection, the PGRPR-LE gene showed no change within 6 h, while hymenoptaecin was significantly upregulated (Figure 1A). After 12 h, the PGRPR-LE, abaecin and defensin genes were strongly upregulated (Figure 1A). However, after 24 h, an obvious decrease in the degree of upregulation of the PGRPR-LE, abaecin and defensin genes was observed compared to that at 12 h of infection; in contrast, hymenoptaecin was upregulated, which was similar to the result at 6 h of infection (Figure 1A). In contrast to the E. coli infection, the S. aureus infection only caused the upregulation of PGRPR-LE and defensin after 12 h and none of the four genes were upregulated within 6 h (Figure 1B). After 24 h, all four genes were upregulated significantly, and the upregulation was obviously higher than that at 12 h of infection (Figure 1B).
Comparison with Drosophila PGRP-LCx

The GST protein did not bind to the Dap-type PGN or the Lys-type PGN (Figure 2D). Based on the contact residues of Drosophila and no binding to the Lys-type PGN, but the short PGRP-LCx had no ability to bind to the Dap-type PGN. Interestingly, the honeybee (A. mellifera) exhibited a weak affinity for the Dap-type PGN and no binding to the Lys-type PGN, but the short PGRP-LCx had no ability to bind to the Dap-type PGN or the Lys-type PGN (Figure 2C). To eliminate the influence of the Glutathione S-transferase (GST) tag on the binding of PGNs, the GST protein was purified and the binding test was performed on the PGNs. The GST protein did not bind to the Dap-type PGN or the Lys-type PGN (Figure 2D). Based on the contact residues of Drosophila PGRP-LCx for the binding of TCT from the PGRP-LCx/PGRP-LCa/TCT complex, we mutated the 14 corresponding residues of Bl-PGRP-LC for the recognition of the Dap-type PGN (Figure 3). In addition, the His\textsuperscript{282}, Thr\textsuperscript{283}, His\textsuperscript{305}, Arg\textsuperscript{309}, Ser\textsuperscript{312}, Tyr\textsuperscript{316}, Arg\textsuperscript{330}, His\textsuperscript{338}, Phe\textsuperscript{340}, Asn\textsuperscript{343}, His\textsuperscript{390}, Ser\textsuperscript{394}, Arg\textsuperscript{395} and Leu\textsuperscript{397} mutations did not alter the binding to Dap-type PGN (Figure 3A). Remarkably, the mutation of the His\textsuperscript{390} of bumblebee Bl-PGRP-LC completely abolished binding to the Dap-type PGN (Figure 3B). Therefore, we hypothesize that the residue His\textsuperscript{390} is the main anchor residue of Bl-PGRP-LC for the recognition of the Dap-type PGN.
wild-type Am-PGRP-LC. (Bl-PGRP-LC) mutants. Each sample includes four lanes. The first lane (M) is the marker; the second lane (+Lys) shows the corresponding protein samples after pull-down with the L-lysine type (Lys-type) PGN from *M. luteus*; and the fourth lane shows the normal corresponding protein as a control. (A) Analysis of the PGN-binding ability of the wild-type Bl-PGRP-LC. (B) The analysis of the PGN-binding ability of the wild-type Dm-PGRP-LC. (C) Analysis of the PGN-binding ability of the wild-type Am-PGRP-LC. (D) Analysis of the PGN-binding ability of the GST-tagged protein.

**Figure 3.** Analysis of the PGN-binding ability of the bumblebee *Bombus lantschouensis* PGRP-LC (Bl-PGRP-LC) mutants. Each sample includes four lanes. The first lane (M) is the marker; the second lane (+Lys) shows the corresponding protein samples after pull-down with the Dap-type PGN from *B. subtilis*; the third lane (+Dap) shows the corresponding protein samples after pull-down with the L-lysine type (Lys-type) PGN from *M. luteus*; and the fourth lane shows the normal corresponding protein as a control. (A) Analysis of the PGN-binding ability of the twelve Bl-PGRP-LC mutants, namely, H283A and T283A, H305A, R309A, S312A, Y316A, R330A, H338A, F340A, N343A, S394A, R395A and L397A. (B) Analysis of the PGN-binding ability of the wild-type Bl-PGRP-LC mutant H390A.
2.3. Structural Insight into PGRP-LC from Different Bee Species

The overall structure of Bl-PGRP-LC retains the classical PGRP structure and overlaps with Drosophila PGRP-LCx. The residues His\(^{365}\), Thr\(^{366}\), Asp\(^{388}\), Tyr\(^{399}\), Arg\(^{413}\), Gly\(^{424}\), Asn\(^{426}\), His\(^{473}\), Ser\(^{477}\) and Glu\(^{480}\) of Drosophila Dm-PGRP-LCx respond to the binding to the Dap-type PGN. In comparison, the Bl-PGRP-LC/TCT complex structure shows that the residues Arg\(^{309}\), Phe\(^{340}\), His\(^{390}\) (His\(^{473}\) in Dm-PGRP-LCx) and Ser\(^{394}\) (Ser\(^{477}\) in Dm-PGRP-LCx) interact with TCT and stabilize the binding of TCT (Figure 4A). We found that His\(^{390}\) is the anchor amino acid for Bombus Bl-PGRP-LC, which is different from the anchor amino acid Arg\(^{413}\) in Drosophila Dm-PGRP-LCx (Figure 3B). Moreover, the residue His\(^{390}\) forms a hydrogen bond with the MurNAc of TCT rather than the meso-Dap of TCT, which interacts with the corresponding anchor residue Arg\(^{413}\) in Dm-PGRP-LCx. Therefore, we hypothesize that the residues that stabilize the interaction between Bl-PGRP-LC and the Dap-type PGN are not conserved in Drosophila Dm-PGRP-LCx, especially the anchor residue, and that the binding to the Dap-type PGN might be different. Moreover, the extracellular regions of PGRP-LCs of different bee species, including Apis cerana, A. mellifera, A. dorsata, A. florea, Bombus lantschouensis, Dufourea novaeangliae, Melipona quadrifasciata, Osmia bicornis and Megachile rotundata, share a high sequence identity. The anchor amino acid His\(^{390}\) and the other contact residues Arg\(^{309}\), Phe\(^{340}\) and Ser\(^{394}\) are completely conserved among the different bee species (Figure 4B). Hence, we hypothesize that the structural characteristics of PGRP-LC for Dap-type PGN binding are extremely similar among bees.

Figure 4. Analysis of the bumblebee Bl-PGRP-LC structure and the sequence alignment of PGRP-LC from bees. (A) Comparison of the Bl-PGRP-LC/TCT complex and the Dm-PGRP-LCx/TCT complex,
colored in gray and magenta, respectively. TCT is shown as a line and the contact residues are shown as sticks. The yellow dashed line indicates the bonds between the TCT and the contact residues in the BI-PGRP-LC/TCT structure. (B) Amino acid sequence alignment of PGRP-LC from different bee species (Melipona quadrifasciata GenBank No. K0X71517.1, Apis dorsata GenBank No. XP_006618726.1, Apis florea GenBank No. XP_003693124.1, Osmia bicorsis GenBank No. XP_029039739.1, Apis mellifera X3 GenBank No. XP_006565566.1, Megachile rotundata X4 GenBank No. XP_003702406.2, Megachile rotundata X3 GenBank No. XP_012138567.1, Dufourea novuangularia GenBank No. KZC08445.1, Apis cerana GenBank No. XP_AGM19450.1, Apis mellifera X5 GenBank No. XP_006565568.1, Apis mellifera X4 GenBank No. XP_006565567.1, Apis mellifera X3 GenBank No. XP_392452.2 and Drosophila PGRP-LCx GenBank No. 2f2l_X). The residues of BI-PGRP-LC contributing to the TCT binding are boxed with red rectangles.

3. Discussion

The bumblebee has fewer genes for innate immunity, which is the unique line of defense against invading pathogens, than the dipteran insects D. melanogaster and A. gambiae. These few immune genes are involved in the recognition, signaling and effector functions, such as genes encoding PGRPs, which are important PRRs and can recognize PGNs to activate important immune pathways, the Toll and Imd pathways [2,6,31,32,34]. In contrast to the corresponding Drosophila protein, our recent study proved that bumblebee PGRP-SA preferentially binds to the Dap-type PGN rather than the Lys-PGN, and that the residue that responds to the binding to the Dap-type PGN is conserved in Apidae [35]. Our subsequent study proved that the bumblebee B. lantschouensis PGRP-LC exists as a single isoform, notably different from the three isoforms observed in Drosophila [37]. In this study, B. lantschouensis PGRP-LC was shown to be able to respond to the Gram-negative bacterium E. coli, and BI-PGRP-LC was able to bind to the DAP-type PGN directly. E. coli infection triggers a quicker and stronger upregulation of genes, including PGRP-LC and the antimicrobial peptide genes abaecin and defensin, than an infection with S. aureus. PGRP-LC expression showed no change at 6 h, and the reaction of PGRP-LC peaked at 12 h after E. coli infection. Although the upregulation level of PGRP-LC was high at 24 h after S. aureus infection, BI-PGRP-LC could not bind to the Lys-type PGN, so this upregulation might be mediated by another unknown mechanism rather than by the recognition of the Lys-type PGN by PGRP-LC. Therefore, we hypothesize that the cell wall of Gram-negative bacteria first needs to be degraded to be recognized by PGRP-LC after infection in bumblebees. These results also confirm that PGRP-LC recognizes Dap-type PGNs that constitute the bacterial cell wall [6,9]. Moreover, bumblebee PGRP-LC preferentially responded to Gram-negative bacteria through the recognition of DAP-type PGNs, where PGRP-LC expression in infected bumblebees showed a parabolic trend between 6 h and 24 h and reached a maximum at 12 h.

Moreover, bumblebee BI-PGRP-LC could recognize Dap-type PGNs with a very high affinity, and this result was consistent with the observation that bumblebee PGRP-LC exhibited an immune response to Gram-negative bacteria. However, Drosophila PGRP-LCx has been reported to recognize Dap-type PGNs with the cooperation of PGRP-LCa, and the Drosophila PGRP-LCa/PGRP-LCx/TCT complex determines the contact residues contributing to the interactions between the TCT and PGRP-LCx, especially the anchor residue Arg\(^{413}\), which interacts with meso-Dap [17]. Nevertheless, the mutation of the 12 corresponding residues did not affect the binding to the Dap-type PGN, except for the His\(^{390}\)Ala mutation, which completely abolished the binding to the Dap-type PGN in BI-PGRP-LC (Figure 3). The corresponding residue His\(^{473}\) in Drosophila PGRP-LCx only participated in the interactions; hence, we proved that BI-PGRP-LC adopts the distinct anchor residue His\(^{390}\) to form hydrogen bonds with MurNAc rather than meso-Dap to stabilize the interaction with the Dap-type PGN. The BI-PGRP-LC/TCT complex showed that the help of residues Arg\(^{397}\), Phe\(^{340}\) and Ser\(^{394}\) is required for stabilization. For now, we confirm that bumblebee PGRP-LC exhibits specific binding characteristics for the Dap-type PGN, and that these contact residues are conserved among different
bee species. The results indicate that although bumblebees possess relatively few PGRPs, these limited PGRPs have evolved a precise function and structure for the recognition of diverse pathogens.

The honeybee *A. mellifera* is a more eusocial insect than the bumblebee *B. laetschouensis* in Apidae [38]. *A. mellifera* possesses a special form of immunity, namely, “social immunity”; however, *B. laetschouensis* inhabits much smaller colonies than *A. mellifera*, with fewer social divisions, which might greatly diminish the defense provided by “social immunity” [39,40]. Therefore, the higher affinity of Bl-PGRP-LC for the Dap-type PGN than that of Am-PGRP-LC might reflect a stronger individual antibacterial immunity in bumblebees than in honeybees. Moreover, bumblebee PGRP-SA also showed a higher affinity for the Dap-type PGN than that of honeybee PGRP-SA in a previous study [35]. However, more studies on the mechanism of activation of the Toll and Imd pathways by PGRPs are needed, and would help explain the specific immune characteristics of bees and the interesting differences in the immune responses of different bee species.

In conclusion, bumblebee PGRP-LC could respond to an infection by Gram-negative bacteria and act as an effective receptor for binding to the Dap-type PGN. Moreover, the recognition of the Dap-type PGN is controlled by the specific anchor residue His\(^{390}\) in the immune response to bacterial infection. In addition, bumblebee Bl-PGRP-LC exhibits a stronger binding to the Dap-type PGN than honeybee Am-PGRP-LC.

4. Materials and Methods

4.1. Bacterial Infection and Real-Time Quantitative PCR Verification

Worker bees at 10 days of age with individual weights between 0.19 and 0.21 g were selected from colonies of the Asian bumblebee *B. laetschouensis* and then reared in separate hives in the same surroundings and fed pollen and sugars to remove unhealthy bees over the next three days. In total, 30 healthy workers (15 workers per group) were starved for 12 h in preparation for feeding with the bacteria *Staphylococcus aureus* and *Escherichia coli*. Finally, 40 µL of bacterial culture (OD value = 1.0) mixed with sugar water at 1:1 was fed to the bees, and the time of feeding for each bee was recorded. Fifteen healthy workers were used as a control, and were fed with 40 µL sugar water. After 6 h, 12 h or 24 h, workers were directly frozen in liquid nitrogen and stored at \(-80^\circ C\) until RNA extraction. There were 5 workers per timepoint in each treatment and control group. The total RNA of each worker from each treatment and timepoint was extracted by using TRIzol (Thermo Fisher Scientific, Waltham, MA, USA), and cDNA was synthesized with 5× All-In-One RT MasterMix with the AccuRT Genomic DNA Removal reactions (Applied Biological Materials, British Columbia, Canada). All the samples for relative quantitative real-time PCR (qRT-PCR) samples were run on an ABI 7500 system (Applied Biosystems, Thermo Fisher Scientific, Waltham, MA, USA) with Bestar SybrGreen qPCR Mastermix (DBI\textsuperscript® Bioscience, Ludwigshafen, Germany). Each reaction was performed in a total volume of 20 µL that contained 5 µL of first-strand cDNA as a template and processed with a preincubation program of 5 min at 95 °C, followed by 40 cycles of 15 s at 95 °C and 45 s at 60 °C and a final step for 10 min at 72 °C. Gene-specific primers were designed with Primer Premier 5 (PREMIER Biosoft, San Francisco, CA, USA) (Table 1). The bumblebee \(\alpha\)-actin gene was used as an internal reference for gene expression. Gene expression levels were calculated using the 2\(^{-\Delta\Delta Ct}\) method [41]. The credibility of 2\(^{-\Delta\Delta Ct}\) values was evaluated by the \(p\)-value from t-test statistics. The mean threshold cycle values for each gene were obtained from three independent PCRs of the different individuals.
Table 1. Primes of the genes for the real-time qPCR verification.

| Genes    | Primers Names | Primers Sequences          |
|----------|---------------|----------------------------|
| Actin    | B-actin F4    | ACGTCCTGGAGAGAGCTACG       |
|          | B-actin R4    | TACCCAGGAAGGAAGGTGGG       |
| PGRP-LC  | LC F1         | GCACTTTGACCTCTCTCTAC       |
|          | LC R1         | GAAGCACCACATCCCACCATC      |
| Abaecin  | Abaecin F1    | ATAATCCGGCAGAGGAGGGC       |
|          | Abaecin R1    | GTTTGGTAATGGGTATGGC        |
| Defensin | Defensin F2   | TCAGTCTGGCCTCTCTCTCTGATGT |
|          | Defensin R2   | TCTTCTTTGTCGTGCACCCAGCAG  |
| Hymenoptaecin | Hymenoptaecin F1 | CGGTGCTTCGTAAGTTTGTTG |  
|          | Hymenoptaecin R1 | TCACTCCGTTTCGTGTAAGAC   |

4.2. Preparation of Proteins

The wild-type *B. lantschouensis* PGRP-LC protein (Bl-PGRP-LC) was expressed and purified as described in a recent report from our laboratory [37]. *Drosophila* PGRP-LCx (Dm-PGRP-LCx) was synthesized by GENWIZ (Suzhou, China) based on the sequence deposited in GenBank (accession number: 2f2l), *A. mellifera* PGRP-LC and all Bl-PGRP-LC mutants were obtained via PCR with primers designed by Primer Premier 5, and the three segments were ligated to the PGEX-6p-1 vector (previously stored in our laboratory). Then, these plasmids were transformed into the *E. coli* strain BL21 (DE3) for expression and purification following the protocols for the wild-type PGRP-LC. Expression from these plasmids was induced with 0.5 mM IPTG. The bacteria were harvested by centrifugation at 8000 × g for 5 min and were then resuspended in cold phosphate-buffered saline (PBS). After sonication, the samples were centrifuged at 11,000 × g, and the supernatants were filtered and loaded onto a GST-tag column or a His-tag column. Finally, the collected proteins were further purified by chromatography on a Superdex 200 16/60 column (GE Healthcare, Boston, MA, USA).

4.3. Lys-Type and Dap-Type PGN-Binding Assays

This assay was performed as previously described with minor modifications: the experiment was conducted at 4 °C by incubating 30 µg of purified wild-type Bl-PGRP-LC or mutant BI-PGRP-LC with 300 µg of insoluble PGN in 300 µL of binding buffer containing 20 mM Tris-HCl (pH 8.0) and 50 mM NaCl on a shaking platform for 1.5 h with Dm-PGRP-LCx as a control [17]. Lys-type and Dap-type PGNs from Micrococcus luteus and Bacillus subtilis were provided by Sigma-Aldrich (Merck KGaA, Darmstadt, Germany). After centrifuging the incubation mixture at 16,000 × g for 5 min, the bound protein was retained in the insoluble PGN pellet and the remainder of the protein was left in the supernatant. Then, 30 µL of the supernatant was used for SDS-PAGE analysis and the PGN-free Bl-PGRP-LC was visualized by staining with Coomassie blue.

4.4. Amino Acid Sequence Alignment and Phylogenetic and Structural Analyses

Alignment of the amino acid sequences of the PGRP-LC proteins from different bee species was performed by using the Clustal Omega server (https://www.ebi.ac.uk/Tools/msa/clustalo/) (Conway Institute UCD, Dublin, Ireland) and Jalview Desktop (http://www.jalview.org/) (University of Dundee, Dundee, UK). The structure of Bl-PGRP-LC was modeled by SWISS-MODEL (https://swissmodel.expasy.org/) (Swiss Institute of Bioinformatics Biozentrum, University of Basel Klingelbergstrasse, Basel, Switzerland) and evaluated based on the GMEQ and QMEAN values [42–45] with the template (PDB ID: 2f2l), and then, the docking of the Bl-PGRP-SA/TCT complexes was completed by the AutoDock4.2 program (The Scripps Research Institute, San Diego, CA, USA) [46]. The structural analyses were completed by the PyMOL 1.8.6 (DeLano Scientific LLC) (Schrödinger, New York, NY, USA) and CCP4 programs (Research Complex at Harwell (RCAH) STFC Rutherford Appleton Laboratory Harwell Science and Innovation Campus Didcot Oxon, Oxford, UK).
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Conflicts of Interest: The authors declare no conflicts of interest.

Abbreviations

PGRPs peptidoglycan recognition proteins
PGN peptidoglycan

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