Genome-Wide Analysis of Soybean Lateral Organ Boundaries Domain Gene Family Reveals the Role in Phytophthora Root and Stem Rot

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The plant-specific lateral organ boundaries (LOB) domain (LBD) proteins, a family of transcription factors, play important roles in plant growth and development, as well as in responses to various stresses. However, little is known about the functions of LBD genes in soybean (Glycine max). In this study, we investigated the evolution and classification of the LBD family in soybean by a phylogenetic tree of the LBD gene family from 16 species. Phylogenetic analysis categorized these proteins into two classes (Class I and Class II) with seven subgroups. Moreover, we found that all the 18 LBD ancestors in angiosperm were kept in soybean, common bean genomes, and genome-wide duplication, suggesting the main force for the expansion of LBD from common bean to soybean. Analysis of gene expression profiling data indicated that 16 GmLBD genes were significantly induced at different time points after inoculation of soybean plants (cv. Huachun 6) with Phytophthora sojae (P. sojae). We further assessed the role of four highly upregulated genes, GmLBD9, GmLBD16, GmLBD23, and GmLBD88, in plant defense in soybean hairy roots using the transient overexpression and knockdown assays. The results showed that GmLBD9 and GmLBD23 negatively regulate plant immunity against P. sojae, whereas GmLBD16 and GmLBD88 positively manipulate plant immunity against P. sojae. Collectively, our findings expand our knowledge of the origin and evolution of the GmLBD gene family in soybean and promote the potential application of these genes in soybean genetic improvement.

Keywords: LBD gene family, phylogenetic analysis, Phytophthora root and stem rot, plant defense, soybean

INTRODUCTION

The lateral organ boundaries (LOB) domain (LBD) proteins are a family of a plant-specific transcription factor with a characteristic N-terminal LBD (Iwakawa et al., 2002). So far, LBD has only been identified in the plant genome, indicating that this unique plant gene family only regulates the plant's developmental process (Shuai et al., 2002). Following the identification of LBD in Arabidopsis, LBDs have also been found in many other plant species, such as Oryza sativa, Malus domestica, Zea mays, and Vitis vinifera. The number of LBD family members identified in different
plant genomes greatly varied ranging from $10$ to $100$ (Yang et al., 2006, 2016, 2017; Wang et al., 2013; Zhang et al., 2014; Cao et al., 2016; Luo et al., 2016; Gombos et al., 2017; Grimplet et al., 2017; Lu et al., 2018).

According to the structural characteristics of the LOB domain, the LBD family can be divided into two subclasses, namely, Class I and Class II (Shuai et al., 2002; Matsumura et al., 2009). Class I family members encode proteins containing two conserved motifs in the LOB domain, namely, a CX$_2$CXX$_2$C and an LX$_6$LX$_3$LX$_4$L leucine zipper-like motif (Shuai et al., 2002; Lee et al., 2009), while family members in Class II contain only a zinc finger-like motif, lacking a leucine zipper-like motif. Due to the incomplete LBD sequence and unstable structure in Class II LBDs, the majority of LBDs belong to Class I (Majer and Hochholdinger, 2011). In model plant Arabidopsis, among 42 LBD family numbers, 36 genes are classified into Class I and 6 genes into Class II (Iwakawa et al., 2002).

Many researches about the LBD family evolution have been performed to explore how this family was classified and originated. Chanderbali et al. (2015) found that LBD might have originated during the early evolution of charophyte algae when they constructed a comprehensive phylogenetic tree of LBD from 307 species, including angiosperms, gymnosperms, monilophytes, lycophytes, liverworts, hornworts, and charophyte algae. No LBDs were identified in Chlamydomonas reinhardtii and Volvox carteri, but several members can be found in Charales species, which suggested that the LBD family already existed before algae and evolved through extensive expansion during land plant diversification (Tang, 2013).

Identification and Phylogenetic Analysis of LBD Genes in 16 Plant Species

Soybean LBD genes (GmLBDs) have been previously identified (Yang et al., 2017). To further understand the functions of GmLBDs in their origination, classification, and even the evolutionary relationship with other species in Leguminosae, we first identified LBD family members in the Phaseolus vulgaris genome (P. vulgaris, common bean), a species with a relatively close evolutionary relationship with soybean in Leguminosae, and in the Cucumis sativus genome (C. sativus, cucumber), an eudicot species. A local BLASTP search was carried out using 42 known Arabidopsis LBD proteins as the query in common bean and cucumber genomes in the NCBI database. Subsequently, all potential LBD protein sequences were further verified by domain analysis using Pfam (LOB domain, DUF260, Pfam number: Pfam03195). As a result, a total of 42 CsLBDs in cucumber and 50 PvLBDs in common bean were finally identified. CsLBDs and PvLBDs were named according to the order of locations on the chromosomes (Supplementary Tables 1, 2).

To further improve the understanding of the phylogenetic classification and evolution of the LBD family in the soybean genome, a comprehensive phylogenetic tree was constructed using 788 amino acid sequences of LBD protein from 16 plant species, including one species each of green alga, moss, fern, and basal angiosperm; eight species in eudicots; and four species in monocots (Supplementary Table 3). The 788 amino acid sequences of LBD protein contained 696 known LBD proteins (Majer and Hochholdinger, 2011; Xu et al., 2016) and auxin signal transduction cascade (Liu et al., 2005; Lee et al., 2015). Members in Class II are involved in metabolisms, such as repressors of anthocyanin synthesis and N availability signals (Rubin et al., 2009). From expression profiles of LBD family genes in Arabidopsis, some LBD genes that belong to Class II were responsive to multiple pathogens, suggesting their functions in plant defense responses (Thatcher et al., 2012a). Further functional analysis showed that AtLBD20 showed resistance suppression against Fusarium oxysporum (F. oxysporum) infection since overexpression of AtLBD20 in roots promoted the colonization of F. oxysporum (Thatcher et al., 2012b). Expression pattern of GmLBD genes after pathogens infection indicated that several GmLBDs were induced in the root and hypocotyl after Bradyrhizobium japonicum and P. sojae mycelia infection (Yang et al., 2017). However, the detailed characterization of GmLBD functions in plant immunity remains unexplored.

In this study, we reconstructed the phylogenetic tree of the LBD gene family from 16 representative genome-available plant species and then compared the evolutionary patterns between soybean and common bean. In addition, based on the expression patterns of P. sojae infection, four GmLBDs were selected for further functional analysis to examine their roles in plant immunity.
that the homologous relationships between using full-length protein sequence. Phylogenetic analysis showed from 14 species and 92 LBD proteins that were newly identified in this study (Yang et al., 2017; Zhang et al., 2020).

Phylogenetic analysis showed that all LBD proteins were classified into two classes (Class I and Class II); Class I is further divided into five subgroups, namely, Class IA, Class IB, Class IC, Class ID, and Class IE, whereas Class II is divided into Class IIA and Class IIB (Figure 1), which is consistent with the previous results (Tang, 2013; Chanderbali et al., 2015; Zhang et al., 2020). Among the 90 GmLBDs, 74 were clustered into Class I, with 19 in Class IA, 25 in Class IB, 19 in Class IC, 4 in Class ID, and 7 in Class IE, while 16 GmLBDs were clustered into Class II.

Evolutionary Relationship of LBD Genes Between Soybean and Common Bean

Given that soybean and common bean have been demonstrated with a close genetic relationship (Vlasova et al., 2016) and both of them are important cash crops. We, therefore, constructed a phylogenetic tree between soybean and common bean to explore the evolutionary relationship of LBD genes in these two genomes using full-length protein sequence. Phylogenetic analysis showed that the homologous relationships between GmLBDs and PvLBDs were obviously observed since almost all clades were included by one PvLBD and one or two GmLBDs (Figure 2). The homologous relationships were inspected by checking GmLBDs and PvLBDs in the same clades and summarized (refer to Table 1). Intriguingly, we found that a total of 38 homologous gene groups were detected, including all PvLBDs and 91% of GmLBDs (82/90), suggesting that the gene duplication in soybean LBDs is another character. In around 70% of clades (25 in 42 clades), one PvLBD and two GmLBDs were closely clustered into one new clade. In the homologous gene group summary, 25 (65%) groups showed the gene ratio of PvLBDs:GmLBDs as 1:2, i.e., one PvLBD has two GmLBDs orthologs.

Soybean experienced one-time independent whole-genome duplication (WGD) compared with common bean, and they diverged only 19.2 million years ago, a relatively short time compared with other legume sister species. To verify the mechanism of gene duplication between soybean and common bean LBD genes, syntenic maps of LBD homologs in these two genomes were built; GmLBD89 and GmLBD90 were excluded in syntenic analysis because of the unassembled genome locations (Figure 3). As a result of 90 GmLBDs and 50 PvLBDs, 112 collinear gene pairs were detected and merged into 38 collinear groups. The collinear groups contain 93% (82/88) of GmLBDs, and 100% (50/50) of PvLBDs genes, which is perfectly consistent with the homologous gene groups summary (Table 1). Interestingly, all these collinear gene pairs contained about 1.62 pairs of conserved genes on average, which is also consistent with the summary of paralog ratios between these two species (Table 1). The syntenic analysis results again proved the close evolutionary relationship of these two species and suggested that WGD might be the main force for LBDs expansion in the soybean genome.

A total of 18 AtLBD ancestries in angiosperms have been previously retraced based on the gene collinearity investigation and phylogenetic relationships (Kong et al., 2017). To further characterize the evolutionary patterns, we also tried to trace the ancestries in GmLBDs and PvLBDs. Interestingly, our data exhibited that GmLBDs and PvLBDs can be detected in all 18 AtLBDs ancient lineages (Table 2), suggesting that no ancestor genes were lost in soybean and common bean genomes. In each AtLBD ancient lineage, 2–10 GmLBD and 1–6 PvLBD paralogs were presented. In most ancient lineage, the number of GmLBDs was much more than that of AtLBDs, such as in lineage 2, AtLOB and AtLBD25 vs 10 GmLBDs and in lineage 11, AtLBD3 vs 6 GmLBDs, which indicated the extensive expansion of GmLBDs in these ancient lineages. However, in the common bean genome, no obvious gene expansion was found except in lineages 2, 9, and 11. The decrease of gene number was also found in some ancient lineages for both GmLBDs and PvLBDs, such as in ancient lineage 8, 4 AtLBDs vs 2 GmLBDs vs 1 PvLBD and in ancient lineage 15, 6 AtLBDs vs 3 GmLBDs vs 2 PvLBD. The ancestry retracement in soybean and common bean demonstrated that LBD is reluctant to be lost and the similar expansion and decrease patterns in some ancient lineages between soybean and common bean mean that they might suffer from parallel evolution.

Expression Profiles of GmLBD Genes During P. sojae Infection

LBD proteins have been reported to play important roles in controlling plant growth and development and also in responding to stress, such as pathogen infection (Iwakawa et al., 2002; Shuai et al., 2002; Feng et al., 2012). To further examine the potential roles of GmLBD proteins in plant immunity, especially in response to P. sojae infection, some GmLBD candidate genes were first identified. Based on previous studies, there are some LBD proteins, which have previously been characterized to involve in plant immune response or upregulated by pathogen infection. AtLBD20 in Arabidopsis and GsLOB1 in Citrus sinensis were found to be involved in plant immunity response to the pathogen (Thatcher et al., 2012b; Hu et al., 2014). So a BLASTP search was performed against the soybean genome
FIGURE 2 | Phylogenetic analysis of LBD proteins in soybean and common bean. The phylogenetic tree was constructed according to the same method. Class II LBD family members were shown in red branches. GmLBDs and PvLBDs were marked with green and blue protein names, respectively. Bootstrap values of more than 60 are represented on each node.

database\(^1\) using AtLBD20 and CsLOB1 as queries and identified 6 GmLBD homologs. In total, 13 GmLBDs were reported to be highly induced in responses to biotic stresses (Yang et al., 2017). Accordingly, a total of 19 GmLBDs genes were selected as candidates for further functional characterization (Supplementary Table 4).

To determine whether these candidate genes play roles in plant defense response against P. sojae attack, the expression patterns of 19 GmLBD genes upon P. sojae infection were examined. The quantitative reverse-transcription PCR (qRT-PCR) was performed using RNA that was extracted from hairy roots of soybean susceptible species Huachun 6 and collected at different time points [0, 1.5, 3, 6, 12, 20, and 24 h after infection (hpi)] after P. sojae infection. The results showed that 16 out of 19 genes were successfully amplified by qRT-PCR, and 15 GmLBDs were found to be induced in the early infection period except for GmLBD37 when compared with the uninfected samples (Figure 4). Among them, 9 GmLBD genes (GmLBD9, GmLBD16, GmLBD23,
TABLE 1 | Summary of GmLBD and PvLBD synteny pairs derived from phylogenetic analysis.

| PvLBD genes | GmLBD genes | Ratio (Pv vs Gm) |
|-------------|-------------|-----------------|
| PvLBD17     | GmLBD30     | 1:1             |
| PvLBD18     | GmLBD86, GmLBD35 | 1:2 |
| PvLBD20     | GmLBD37     | 1:2             |
| PvLBD21     | GmLBD7, GmLBD62 | 1:2 |
| PvLBD22     | GmLBD22     | 1:1             |
| PvLBD23     | GmLBD38     | 1:2             |
| PvLBD26     | GmLBD45     | 1:2             |
| PvLBD28     | GmLBD48, GmLBD74 | 1:2 |
| PvLBD30     | GmLBD59, GmLBD30 | 1:2 |
| PvLBD32     | GmLBD49     | 1:1             |
| PvLBD33     | GmLBD65, GmLBD64, GmLBD69 | 2:3 |
| PvLBD34     | GmLBD67, GmLBD80 | 2:2 |
| PvLBD35     | GmLBD71     | 2:1             |
| PvLBD37     | GmLBD75, GmLBD84 | 1:2 |
| PvLBD38     | GmLBD27, GmLBD69, GmLBD89 | 2:4 |
| PvLBD40     | GmLBD80     | 2:2             |
| PvLBD42     | GmLBD82     | 2:2             |
| PvLBD43     | GmLBD90     | 2:1             |
| PvLBD48     | GmLBD70, GmLBD23 | 1:2 |
| PvLBD50     | GmLBD31     | 1:2             |
| PvLBD51     | GmLBD18, GmLBD69, GmLBD89 | 2:4 |
| PvLBD52     | GmLBD78, GmLBD20 | 2:2 |
| PvLBD54     | GmLBD82, GmLBD88 | 1:2 |
| PvLBD55     | GmLBD14, GmLBD82 | 2:2 |
| PvLBD56     | GmLBD88     | 1:2             |
| PvLBD57     | GmLBD80, GmLBD89 | 2:2 |
| PvLBD59     | GmLBD90     | 2:2             |
| PvLBD60     | GmLBD99     | 2:2             |
| PvLBD61     | GmLBD72, GmLBD61* | – |
| PvLBD62     | GmLBD63*    | –               |
| PvLBD63     | GmLBD90     | 2:1             |
| PvLBD64     | GmLBD20     | 1:2             |
| PvLBD65     | GmLBD18, GmLBD69, GmLBD89 | 2:4 |
| PvLBD66     | GmLBD70, GmLBD23 | 1:2 |
| PvLBD67     | GmLBD78, GmLBD20 | 2:2 |
| PvLBD68     | GmLBD80, GmLBD89 | 2:2 |
| PvLBD69     | GmLBD90     | 2:2             |
| PvLBD70     | GmLBD70, GmLBD23 | 1:2 |
| PvLBD71     | GmLBD80     | 2:2             |
| PvLBD72     | GmLBD90     | 2:2             |
| PvLBD73     | GmLBD90     | 2:2             |
| PvLBD74     | GmLBD90     | 2:2             |
| PvLBD75     | GmLBD90     | 2:2             |
| PvLBD76     | GmLBD90     | 2:2             |
| PvLBD77     | GmLBD90     | 2:2             |
| PvLBD78     | GmLBD90     | 2:2             |
| PvLBD79     | GmLBD90     | 2:2             |
| PvLBD80     | GmLBD90     | 2:2             |
| PvLBD81     | GmLBD90     | 2:2             |
| PvLBD82     | GmLBD90     | 2:2             |

*indicated GmLBDs failing to find paralogs in common bean.

GmLBD genes Closely Associated With Soybean Immunity Against P. sojae Infection

To investigate the functions of GmLBD genes in soybean immunity, four GmLBD genes (GmLBD9, GmLBD16, GmLBD23, and GmLBD88) showed remarkably upregulated expression in Huachun 6 were chosen to reveal how they regulate soybean immunity through the transient overexpression and knockdown assays. GmLBD9, GmLBD16, GmLBD23, and GmLBD88 were first transiently overexpressed in soybean hairy roots by Agrobacterium-mediated transformation, and then those transformed hairy roots were inoculated with P. sojae strain P6497-RFP. We discovered that more P. sojae oospores can be observed in hairy roots expressing GmLBD9 and GmLBD88 at 48 hpi in relative to those hairy roots expressing empty vector (EV) by microscope observation (Figure 5A). Consistently, oospores and biomass of P. sojae were much higher in soybean hairy roots inoculated with GmLBD9 and GmLBD88 than in roots inoculated with the EV control (Figures 5C,D); these data indicate that the expression of GmLBD9 and GmLBD88
FIGURE 3 | Synteny analysis of LBD genes in soybean and common bean. Circular collinearity analysis of LBD genes in soybean and common bean genomes. GmLBDs and PvLBDs were mapped to their corresponding chromosomal locations and represented in a circular diagram using Circos. Colored lines connect the pairs of orthologous LBD genes in the syntenic blocks of these two genomes. Soybean and common bean chromosomes are denoted as red and blue boxes, respectively.

could promote the colonization of *P. sojae* in soybean hairy roots. Interestingly, we found that the individual expression of GmLBD16 and GmLBD23 inhibited *P. sojae* infection in soybean hairy roots, showing fewer oospores and lower relative biomass of *P. sojae* in transiently expressing hairy roots (Figures 5A, 6C,D). Moreover, the expression of those four recombinant proteins was confirmed by western blot, respectively (Figures 5B, 6B). Overall, these results suggest that GmLBD9 and GmLBD88 may be the negative immune regulators for soybean resistance against *P. sojae*, while GmLBD16 and GmLBD23 may be the positive immune regulators.

These data prompted us to further verify their roles in soybean *Phytophthora* root and rot. Two GmLBDs (GmLBD9 and GmLBD23) were selected to perform transient silencing in soybean hairy roots by RNA interference (RNAi) technique. GmLBD9 and GmLBD23 have been proved to be negative and positive immune regulators in soybean immunity, respectively. To specifically silence GmLBD9 or GmLBD23, approximately 200–300 bp of 5′- or 3′-end UTR fragments were cloned into pK7GW1WG2D vector to generate an RNAi recombinant construct. And then, these constructs were introduced into soybean hairy roots by *Agrobacterium*-mediated transformation. Each RNAi construct successfully silenced these targets as shown by qRT-PCR analysis and revealed that the expression of GmLBD9 and GmLBD23 were obviously reduced by 70–80% in silencing hairy roots (Figure 7B). Subsequently, these silenced hairy roots were challenged with *P. sojae* strain P6497-GFP. The results displayed that GmLBD9-silenced hairy roots showed less oospores (Figures 7A, C) and lower relative biomass of *P. sojae* than those hairy roots induced by EV (Figure 7D).
Together with the above data, our results further indicated that the GmLBD9 gene negatively regulates soybean immunity against P. sojae infection. As such, the infection on GmLBD23-silenced roots exhibited that much more oospores were observed through fluorescence microscope observation (Figure 7A). Quantification of oospore number and biomass also proved that silencing of GmLBD9 promoted colonization of P. sojae in soybean hairy roots (Figures 7C,D), indicating that the GmLBD9 gene positively manipulates soybean immunity against P. sojae infection. Taken together, our data suggest that GmLBD9 and GmLBD88 are two negative immune regulators and GmLBD16 and GmLBD23 are two positive regulators of plant immunity against P. sojae infection.

**DISCUSSION**

As a plant-specific gene family, LBD family proteins have drawn many researchers’ attention to explore their phylogenetic diversification, origination, and even functional characteristics by genome-wide analysis (Wang et al., 2013; Zhang et al., 2014; Yang et al., 2016). In this study, amino acid sequences of LBD members from 16 representative species, including green alga, basal angiosperm, monocots, and dicots, were collected, and a comprehensive phylogenetic tree was constructed (Figure 1). We mainly focused on the evaluation of LBD family members in soybean and common bean. Our data have shown that all 18 ancient gene lineages for angiosperms were preserved in soybean and common bean. This is consistent with the divergence of the LBD family in early land plants, seed plants, and angiosperm. In early land plants, 7 ancient genes are deduced and kept in a stable amount. Also in angiosperm genomes, 18 major lineages can be detected in rice and Arabidopsis genomes (Kong et al., 2017). Above all the results suggested that LBDs were reluctant to be lost during evolution. Moreover, it was supposed that the additional WGD events that happened in the soybean genome were probably the major driving force behind the substantial gene content increase due to the ratio of LBD ortholog numbers and synteny analysis between soybean and common bean. Similar results were also obtained by another gene family evolutionary analysis. Wu et al. investigated WRKY transcription factors in common beans and they deduced that it was the result of genome duplication of the two WRKY transcription factors in
FIGURE 4 | Expression profiles of GmLBD genes in Huanchun 6, Williams 82, and Tianlong 1 cultivars upon P. sojae infection. (A) Soybean hairy roots were collected at 0, 1.5, 3, 6, 12, 16, 20, and 24 h after P. sojae strain P6497 infection. Total RNA was extracted and expression profiles of 9 GmLBD genes at various time points during infection were determined by qRT-PCR. The Soybean GmCYP2 gene was used as an internal control. Error bars indicate three biological replicates. Soybean hairy roots of Williams 82 (B) and Tianlong 1 (C) were collected at 0, 6, 12, and 24 h after P. sojae strain P6497 infection. Total RNA was extracted and expression profiles of 4 GmLBD genes at various time points during infection were determined by qRT-PCR. The Soybean GmCYP2 gene was used as an internal control. Error bars indicate three biological replicates.

the soybean genome rather than in the common bean genome (Wu et al., 2017).

Although the expression dataset of GmLBDs in various tissues and under biotic and abiotic stresses, including pathogen infection, is available online (Yang et al., 2017), functional characterization of GmLBDs in plant immunity has not been previously documented. In this study, four selected GmLBDs were significantly induced on P. sojae infection; however, GmLBD9 and GmLBD88 showed opposite roles with GmLBD16 and GmLBD23 in the subsequent functional analysis (Figures 5–7). It might be caused by two possible reasons: one is that the gene responses might be tissue, growth stage, or genotype-specific. In this study, three species showing different resistance to P. sojae strain P6497 were included in examining gene expression patterns. GmLBD9 was highly induced in Huachun 6 but showed no change in Williams 82 and Tianlong 1. Gene expression between four chickpea genotypes, including resistant, moderately resistant, susceptible, and wild relative genotype, was different when these four genotypes were challenged by ascochyta blight (Coram and Pang, 2006). Expression comparison of responses to volatiles in Arabidopsis revealed that genes involved in flavonoid biosynthesis were downregulated in leaves and upregulated in roots, photosynthesis genes were expressed in the seedling stage and induced at the mature stage (Hao et al., 2016). Therefore, the gene expression profile is just a useful tool providing us with the potential candidates for further functional validation since it has fast, convenient, and high throughput. The other reason is that it is common that these genes were regulated by diverse pathways. ShARP3 can be highly induced during an incompatible and compatible interaction against On-Lz infection and finally turned out to be a positive regulator of plant immunity due to its overexpression inducing rapid hypersensitive cell death and reactive oxygen generation (Sun et al., 2019). EDS1-interacting J protein 1 (EIJ1) is proved as an EDS1-dependent negative regulator of innate plant immunity with significant induction by the treatment with Pst DC3000 or SA (Liu et al., 2021).

In summary, 788 LBDs from 16 species, including 90 from soybean and 50 from common bean, were used to perform an extensive phylogenetic analysis of LBD proteins. Phylogenetic analysis categorized these proteins into two groups, namely, Class I and Class II, and Class I was further classified into five subgroups. None of the ancestor genes were lost in the soybean and common bean genomes in ancestor gene retracement. The evolutionary analysis indicated that the expansion of LBD numbers in the soybean genome was primarily driven by WGD. Based on the gene expression profiles on P. sojae infection, four GmLBDs were chosen for further functional characterization and discovered GmLBD9 and GmLBD88 function as negative immune regulators and GmLBD16 and GmLBD23 as positive immune regulators in plant immunity. So this study expands our
knowledge of the origin and evolution of the GmLBD gene family in soybean and promotes the potential application of these genes in soybean genetic improvement.

MATERIALS AND METHODS

Plant and Microbe Cultivation
Soybean cotyledons (Huachun 6, Williams 82 and Tianlong 1) were grown in a greenhouse at 25°C with a 16/8 h (light/dark) photoperiod. *P. sojae* strains, namely, P6497, P6497-RFP, and P6497-GFP, were routinely maintained on a 10% vegetable (V8) juice medium at 25°C in darkness.

Identification of LBDs in Cucumber and Common Bean
To obtain cucumber and common bean LBD protein sequences, all known 43 *Arabidopsis* LBD protein sequences were used as a query to perform BLASTP with an *e*-value of $1 \times e^{-10}$ against the protein sequences database of *C. sativus* and *P. vulgaris* (NCBI³). Redundant sequences that are partial or alternatively spliced sequences from the same locus were removed. Then conserved domain of LBDs (LOB domain, DUF260, Pfam number: Pfam03195) acquired from Pfam³ was used for a blast to identify CsLBDs and PvLBDs with DUF260 as a query (El-Gebali et al., 2019). Finally, each gene was named based on its location on the chromosome.

Phylogenetic Analyses
To construct a phylogenetic tree of LBDs in 16 species, 788 full-length LBD protein sequences were aligned using the multi-sequence alignment program ClustalW. A phylogenetic tree was constructed with ML (maximum-likelihood) method in MEGA X and 100 times of bootstrap replicates. The phylogenetic tree was further manipulated by the program Interactive Tree of Life (iTOL⁴) (Zhu et al., 2019).

Synteny Analyses
The chromosomal length and locations of each GmLBD and PvLBD were retrieved from the soybean genome database in SoyBase⁵ and common bean genome database in NCBI. Advanced Circos program in TBtools (Chen et al., 2020a) was used for collinearity analyses.

Plasmid Construction
For overexpression assay in soybean hairy roots, fragments containing full-length CDS sequences were amplified with gene-specific primers (Supplementary Table 5) and then ligated into vector PGFGC5941 by homologous recombination (Vazyme, C112-02-AB), which adds a C terminal FLAG tag. For gene silencing assay, fragments derived from the 5′ or 3′ UTR regions with 200–300 bp in length were amplified and then cloned into pK7GW1WG2D vectors with Gateway technology (Thermo Fisher Scientific, 12538120).

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1. https://www.ncbi.nlm.nih.gov/
2. http://pfam.xfam.org/
3. http://pfam.xfam.org/
4. https://itol.embl.de/
5. https://www.soybase.org/sgn/
Transformation of Soybean Cotyledons and *P. sojae* Infection Assays

Soybean seeds (Huachun 6) were sanitized with a mixture of 84 disinfectants and concentrated hydrochloric acid (96:4) and then grown on germination medium (Gm medium). After around 6 days of growth, cotyledons were removed from soybean seedlings and cut a wound (around 0.3 cm in diameter and 0.2–0.3 cm in depth) close to the petiole with a sterile knife. *Agrobacterium rhizogenes* (A. rhizogenes) K599 cell suspensions were inoculated on the wound, and then cotyledons were continued growing on Murashige and Skoog medium (MS medium). Around 3–4 weeks later, soybean hairy roots overexpressing or silencing GmLBDs were observed by fluorescence microscopy and further confirmed by western blot or qRT-PCR. Selected overexpressing hairy roots and silencing hairy roots were infected with P6497-RFP and P6497-GFP in wet and dark conditions at 25°C for around 48 h, respectively. Around 3- to 5-day-old *P. sojae* hyphae grown in 10% of V8 medium were used for soybean hairy roots infection.

RNA Extraction and Quantitative Reverse-Transcription PCR Analysis

Total RNA was isolated from hairy roots using TRIzol Reagent (TaKaRa, 9109). Isolated RNA samples were quantified using a NanoDrop spectrophotometer (Thermo Fisher Scientific, NanoDrop One) and then treated with DNase I (Thermo Fisher Scientific, AM2222) to remove any residual DNA contamination. In total, 1 μg of DNA-free RNA samples were converted to cDNA using a cDNA synthesis kit (Vazyme Biotech, R212-02-AF). First-strand cDNA was synthesized using HiScript II 1st Strand cDNA Synthesis Kit (+gDNA wiper) with Oligo-(dT) 23VN (Vazyme Biotech, R212-02-AF).

For measuring the transcript level of *GmLBD* genes from soybean during *P. sojae* infection, 1- or 2-week-old soybean (Huachun 6, Williams 82 and Tianlong 1) secondary roots was infected with R6497 and collected at different time points after infection. For measuring the transcript level of *GmLBDs* in different tissues, leaves, roots, and stems were sampled from 3- to 4-week-old soybean plants. Total RNA was extracted and used as a template for reverse transcription. To determine gene silencing efficiency, RNA was extracted from 3- to 4-week-old hairy roots induced by K599 containing PK7GWIG2D (II) and PK7GWIG2D (II)-*GmLBDs*. In all the quantitative PCR (qPCR) reactions, gene-specific primers (*Supplementary Table 5*) were designed specifically in 5'- or 3'-end UTR of genes avoiding the fragments for gene silencing. Then, qPCR was carried out using Maxima SYBR Green/ROX qPCR Master Mix (Vazyme Biotech, Q711-02-AA) (Zhang et al., 2019; Zhu et al., 2020). *GmCYP2* gene (Hu et al., 2009) was used as an
internal reference. Three independent biological replicates were conducted for each treatment with similar results.

**DNA Extraction and Biomass Assay**

Cetyltrimethylammonium bromide (CTAB) method was used for genomic DNA extraction (Li et al., 2017). Briefly, 500 µl of CTAB buffer was added into crushed soybean hairy roots to lyse plant cells in a water bath at 60°C for 1 h, and an equal volume of chloroform-isomyl alcohol (24:1) was added while vigorously shaking for 30 s. After centrifugation at 12,000 rpm at room temperature, the supernatant was transferred into a new Eppendorf tube, an equal volume of ice-cold 100% ethanol was added to precipitate DNA at −20°C for more than half an hour, and then 75% ethanol was used to wash DNA. After drying in the hood, 100 µl of sterilized H2O was added to dissolve DNA at 55°C. The extracted DNA was also quantified by Nanodrop and then diluted into the same concentration for biomass assay. Primers of *GmCYP2* (Hu et al., 2009) from soybean and *PsActin* from *P. sojae* were used for biomass assay (Shi et al., 2020; Supplementary Table 5).

**Immunoblotting Analyses**

Proteins from the sample lysate were fractionated by SDS-PAGE gel. The separated proteins were transferred from gels to PVDF blotting membrane (GE, A10203127) (pretreated with methanol for 15 s) using Transfer Buffer (BIO-RAD, Cat. #10026938). The membrane was then blocked using 5% non-fat dry milk dissolved in TBST (TBS with 0.1% Tween 20) (also called TBSTM) for 1 h at room temperature with 30–40 rpm shaking, followed by three washes with TBST, and then primary antibody anti-Flag (1:5,000; MBL, M185-3L) and secondary antibody goat anti-mouse antibody (1:5,000; MBL, Lot 366) were applied to the membranes for 1 h in order. Finally, the membrane was visualized using the Western Blotting Substrate kit (Thermo Fisher Scientific, 34580) by multifunctional fluorescent molecular imager (GE, Amersham Imager 600) at 780 and 800 nm excitation (Chen et al., 2020b).

**Statistical Analysis**

The statistical analysis in biomass assay, gene expression profile, and oospore numbers quantification was performed using an unpaired T-test in Graphpad Prism5.

**DATA AVAILABILITY STATEMENT**

The original contributions presented in the study are included in the article/Supplementary Material, further inquiries can be directed to the corresponding author/s.
AUTHOR CONTRIBUTIONS

YQ conceived and designed the experiments and revised the manuscript. SF, JS, YH, DL, and LG performed the experiments. ZZ and GL provided the suggestion for this research. SF and JS analyzed the experimental data and wrote the manuscript. All authors have read and approved the final manuscript.

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SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fpls.2022.865165/full#supplementary-material

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