Overexpression of OsHSP18.0-CI Enhances Resistance to Bacterial Leaf Streak in Rice

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Abstract

Background: The small heat shock proteins represent a large family of proteins that respond to a wide range of abiotic and biotic stresses. OsHsp18.0-CI confers tolerance to salt and cadmium and interacts with viral RNA-dependent RNA polymerase (RdRp). However, the direct function of OsHsp18.0-CI in resistance against biotic stresses remains unclear in rice.

Results: Here, we report that the expression of OsHsp18.0-CI was up-regulated upon inoculation with RS105, a strain of Xanthomonas oryzae pv. oryzicola (Xoc) that causes bacterial leaf streak in rice. In comparison with wild-type, OsHsp18.0-CI overexpression (OE) lines exhibited enhanced resistance to RS105, whereas repression lines exhibited compromised resistance to RS105. In addition, the transcriptional profiles of wild type and OE lines were compared with and without inoculation with RS105. After inoculation with RS105, most of the genes with up-regulated expression were commonly stimulated in the wild type and OE lines, with stronger induction in the OE lines than in wild type.

Conclusion: Our study reveals that OsHsp18.0-CI positively regulates resistance to Xoc by mediating an enhanced version of the basal defense response in rice.

Keywords: Bacterial leaf streak, Disease resistance, Heat shock protein, OsHsp18.0-CI, Xanthomonas oryzae pv. oryzicola

Background

Bacterial leaf streak (BLS) caused by the gram-negative bacterial pathogen Xanthomonas oryzae pv. oryzicola (Xoc) is a globally important disease affecting rice production. This pathogen often penetrates the leaf through stomata or wounds and propagates in the apoplast of mesophyll tissue, finally resulting in water-soaked lesion symptoms (NINO-Liu et al. 2006). Xoc normally causes yield losses of approximately 20% depending on the rice variety and climatic conditions (Ou 1985). Since first reported in southern China in 1957, Xoc has spread to most areas of Hainan province, southern China, southwest China and central China, becoming one of the most devastating quarantine rice diseases (NINO-Liu et al. 2006; Xu et al. 2008).

Breeding disease-resistant varieties is an ideal strategy to manage BLS disease. However, BLS resistance has been considered quantitatively inherited in rice. More than 13 quantitative trait loci (QTLs) have been mapped from the indica rice varieties Acc8558 and Dular (Tang et al. 2000; Chen et al. 2006). The major QTL is qBlsr5a, in the short arm of chromosome 5, which has the largest effect and explains approximately 14% of phenotypic variation (Tang et al. 2000; Han et al. 2008). It was identified that qBlsr5a is mainly controlled by xa5, a major gene for resistance to bacterial blight caused by Xanthomonas oryzae pv. oryzicola (XIE et al. 2014). Recently, a new dominant resistance locus named Xo1 has been identified in the qualitative resistance against the African clade of Xoc strains of the American heirloom rice variety Carolina Gold Select (TRIPLETT et al. 2016). Interestingly, this resistance can be triggered by a transcription activator-like (TAL) effector in Xoc (TRIPLETT et al. 2016). Alternatively, a non-host resistance gene, Rxol, which encodes a typical plant resistance protein containing a nucleotide-binding site-leucine rich repeat (NBS-LRR)
from maize, presents qualitative resistance to BLS in rice (Zhao et al. 2005). In addition to genetic mapping, some defense-related (DR) genes have been reported to exhibit up-regulated expression upon Xoc inoculation to positively or negatively regulate the BLS resistance (Zhou et al. 2010; Kou and Wang 2010; Xu et al. 2013). Overexpression of the NBS-LRR-type gene DEPG1 increased susceptibility to Xoc strain RS105, implying negative regulation of rice immunity (Guo et al. 2012). Consistent with negative regulation of rice immunity, suppression of the expression of DR genes also enhances the resistance to BLS, such as OsWRKY45-1 (Tao et al. 2009), OsMPK6 (Shen et al. 2010) and NRRB, which encodes a receptor-like cytoplasmic kinase (Guo et al. 2014). By contrast, positive regulation mechanisms of resistance to Xoc have been poorly explored in rice. Overexpression of OsPGIP4, which encodes a polygalacturonase-inhibiting protein, and GH3-2 which encodes an indole-3-acetic acid-amido synthetase, significantly enhance resistance to BLS in rice (Fu et al. 2011; Feng et al. 2016).

Heat shock proteins (HSPs) are found across a wide diversity of organisms. HSPs are chaperones that assist in protein folding and prevent irreversible protein aggregation (Waters 2013). They include a number of conserved protein families: HSP100s, HSP90s, HSP70s, HSP60s, and HSP20s or small HSPs (sHSPs). Among HSPs, the sHSP family is one of the most abundant and complex groups, and the monomers of these proteins range in size from 12 kDa to 42 kDa. As this alternative name of the HSP20 family suggests, most sHSPs are in the range of 15 kDa to 22 kDa, and these proteins contain a highly conserved central domain called the Alpha Crystallin Domain (ACD), which is involved in dimerization (Jaspard and Hunault 2016). There are 40 ACD-containing genes, of which 23 constitute sHSPs in rice (Sarkar et al. 2009). These sHSPs are stimulated in response to a wide range of abiotic stresses. For instance, the expression of Os03g16030, formerly named Hsp18.0-CI, OsMSR3 and OsSHSP1, was elevated in response to treatment with anoxia, heat, cold, salt, drought, L-azetidine-2-carboxylic acid (AZC) and cadmium (Guan et al. 2004; Sarkar et al. 2009; Cui et al. 2013; Ham et al. 2013). Consistent with their up-regulated expression patterns, heterogeneous overexpression of Os03g16030 in Arabidopsis enhances tolerance to salt and cadmium stresses (Cui et al. 2013; Ham et al. 2013). The expression of Os03g16030 was also activated by biotic stresses, such as Magnaporthe grisea (M. grisea) and at least seven rice viruses (Sarkar et al. 2009; Li et al. 2015). Os03g16030 has been reported to directly interact with the N-terminus of viral RNA-dependent RNA polymerase (RdRp) to change its subcellular localization and distribution pattern in Nicotiana benthamiana (Li et al. 2015). However, the function of Os3g16030 against rice pathogens remains unclear.

Here, we determined that the expression of Os03g16030 (named OsHsp18.0-CI according to Sarkar et al. 2009) was up-regulated by inoculating with Xoc strain RS105. Transgenic plants overexpressing OsHsp18.0-CI exhibited enhanced resistance to RS105, and lines with suppressed expression exhibited increased susceptibility to RS105. Using an RNA-seq strategy, we further found that the enhanced expression of defense-related genes involved in basal defense was enriched in OsHsp18.0-CI overexpressing (OE) lines compared with wild-type (WT). The expression of these genes was activated more strongly in OE than in WT plants after inoculation with RS105. Our results suggest that the sHSP OsHsp18.0-CI positively regulates resistance to Xoc in rice.

**Results**

**The Expression of OsHsp18.0-CI is Induced by Xoc**

As the expression of OsHsp18.0-CI is activated by M. grisea and rice viruses (Sarkar et al. 2009; Li et al. 2015), we determined if it is also induced by Xoc. We measured the expression of OsHsp18.0-CI post-inoculation with the Xoc strain RS105 in the susceptible rice variety Shengdao 806. The transcript level of OsHsp18.0-CI increased quickly after 6 h post-inoculation (hpi) with RS105 (Fig. 1) and reached maximum level at 24 hpi, with an increase of 16-fold compared with the control. Expression then decreased to nearly the same level as the control at 72 hpi. This result suggests that the expression of OsHsp18.0-CI is induced by Xoc, as observed for previously reported defense-related genes (Feng et al. 2016).

![Image](attachment:image-url)

**Fig. 1** OsHsp18.0-CI is induced by Xoc inoculation. The expression level of OsHsp18.0-CI in response to RS105 in Shengdao806 at 0, 6, 12, 24 and 72 h post inoculation. The bars represent the means (three replicates for gene expression) ± SD. The significance of expression compared to 0 h at P values of less than 0.05 and 0.01 are marked by “*” and “**”, respectively.
Overexpression of OsHsp18.0-CI Enhances Resistance Against Xoc in Shengdao 806

To assess whether up-regulated OsHsp18.0-CI expression could enhance resistance against Xoc, Agrobacterium-mediated transformation of the pU1301-OsHsp18.0-CI construct was used to obtain 12 independent T₀ individuals (CD39R-1 ~ CD39R-12). Nine of 12 T₀ lines were identified as positive transgenic individuals by PCR analysis and qRT-PCR, which quantified the expression level of OsHsp18.0-CI. The expression level of OsHsp18.0-CI was significantly increased in all 9 individuals (Fig. 2b). The T₀ individuals were inoculated with Xoc strain RS105 along with Shengdao806 (WT) at the seedling stage. All nine positive individuals exhibited enhanced resistance to RS105, with lesion lengths ranging from 1.58 cm to 2.02 cm, compared with an average of 2.28 cm for WT plants (Fig. 2a). The lesion length was significantly shorter in 7 of 9 OE lines (t test, P < 0.05) than in WT plants. Furthermore, the deduced lesion length in transgenic individuals was correlated (r = 0.871, α = 0.05, n = 12) with the expression level of OsHsp18.0-CI.

To evaluate the disease resistance of transgenic lines in the T₁ generation, two lines (CD39R-7 and CD39R-11) were selected for inoculation with RS105, with WT plants as a control. The progeny of the two T₁ lines showed significant reductions in lesion length and were identified as positive plants that carried the construct pU1301-OsHsp18.0-CI (Fig. 2c and d). Two lines were further identified that exhibited resistance against RS105 in the T₂ generation (Fig. 2a and b). Consistent with the enhanced resistance against RS105, the bacteria populations of CD39R-7 was dramatically less than control at 4 dpi and 8 dpi in leaves (Fig. 4c). These results suggested that OsHsp18.0-CI positively regulates resistance to Xoc in rice.

Repressing OsHsp18.0-CI Expression Enhances Susceptibility to Xoc

To further identify the role of OsHsp18.0-CI in rice-Xoc interactions, we introduced the OsHsp18.0-CI-RNAi construct (CD40R) into Shengdao 806 to silence the expression of OsHsp18.0-CI by Agrobacterium-mediated transformation. Twelve independent individuals (CD40R-1 ~ CD40R-12) were obtained. Nine of 12 individuals showed clear reductions in the expression level of OsHsp18.0-CI (Fig. 3b). At the seedling stage, all individuals and WT were inoculated with RS105. The nine positive individuals exhibited significantly increased lesion length, with lesion lengths ranging from 2.36 cm to

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**Fig. 2** Resistance of OsHsp18.0-CI-overexpressing plants to the Xoc strain RS105. a) Lesion lengths of the 12 individual T₀ transgenic plants with pU1301-OsHsp18.0-CI (CD39R) at 14 days post inoculation (dpi). b) Relative expression levels of OsHsp18.0-CI in the T₀ transgenic plants. c and d) Resistance of two OsHsp18.0-CI-overexpressing lines (CD39R-7 and CD39R-11) to Xoc in the T₁ generation. The average lesion length for each plant was measured at 10 to 15 inoculation sites at 14 dpi. The gel graph indicates the plants carrying pU1301-OsHsp18.0-CI that were validated by PCR. The bars represent the means ± SD. * and ** indicate significant (t test, P < 0.05) and extremely significant (t test, P < 0.01) differences in lesion lengths between wild-type and transgenic plants, respectively.
3.32 cm, compared with average of 2.28 cm for WT plants (Fig. 3a). Consistent with these results, three negative individuals exhibited an average lesion length of 2.20 cm, similar to that in WT plants. In addition, the lesion length in the transgenic plants was significantly correlated ($r = 0.933$, $\alpha = 0.05$, $n = 12$) with the expression level of OsHsp18.0-CI.

Two individuals (CD40R-9 and CD40R-12) were chosen for further analysis at T1 generation. All positive T1 plants exhibited significantly increased lesion lengths compared with WT or negative plants segregated from the T0 transgenic lines (Fig. 3c and d). In the T2 generation, the progenies of CD40R-9 and CD40R-12 were more susceptible to RS105 than WT, and the bacteria populations in leaves of CD40R-12 was more than WT at 4 dpi and 8 dpi (Fig. 4a, b and c).

In addition, the transgenic plants were inoculation with several Chinese Xoc isolates including JLB2-24, HNB8-47, HGA1 and HGA2 to investigate the broad spectrum resistance of OsHsp18.0-CI. Compared with WT, the OE lines (CD39R-7) significantly enhanced resistance to all four Xoc isolates, and the RNAi lines (CD40R-12) were more susceptible (Fig. 4d). All the results strongly suggest that OsHsp18.0-CI positively regulates the resistance to Xoc in rice.

**Identification of Differentially Expressed Genes in Shendao 806 (CD39R-7) and Shendao 806 by RNA-seq**

To identify genes involved in OsHsp18.0-CI-mediated resistance, RNA-seq of both infected (24 hpi) and non-infected leaves of CD39R-7 (OsHsp18.0-CI over-expression line, OE) and Shendao 806 (WT) was conducted. An average of 24 million clean reads were obtained that mapped on the rice genome at an average rate of 87%, representing an average of 28,872 genes that were expressed in each sample (Additional file 1: Table S1).

Because the introduced gene could directly trigger the plant defense response regardless of the inoculation of pathogens, we first analyzed the differentially expressed genes (DEGs) under non-infected conditions. Compared with non-infected WT (WT), a total of 403 up-regulated genes and 72 down-regulated genes were detected in non-infected CD39R-7 (OsHsp18.0-CI over-expression line, OE) and Shendao 806 (WT) was conducted. An average of 24 million clean reads were obtained that mapped on the rice genome at an average rate of 87%, representing an average of 28,872 genes that were expressed in each sample (Additional file 1: Table S1).

Because the introduced gene could directly trigger the plant defense response regardless of the inoculation of pathogens, we first analyzed the differentially expressed genes (DEGs) under non-infected conditions. Compared with non-infected WT (WT), a total of 403 up-regulated genes and 72 down-regulated genes were detected in non-infected CD39R (OE, Fig. 5a and b, Additional file 2: Table S2). GO functional annotation of the DEGs indicated that the up- and down-regulated genes could be classified into 28 categories, as shown in Additional file 3: Figure S1, such as response to endogenous stimulus, signal transduction, cellular process, response to abiotic stimulus, response to stress, and response to biotic stimulus.
To further compare the DEGs activated by inoculation of RS105 between WT and OE, the transcriptional profiles were also analyzed between infected WT (WT-24) and WT and between infected OE (OE-24) and OE. Compared with WT, a total of 898 up-regulated and 128 down-regulated genes were identified in WT-24 from the RNA-seq data, and 1844 up-regulated and 154 down-regulated genes were identified in OE-24 compared with non-infected OE. All DEGs were classified into 40 different functional categories according to Gene Ontology analysis using Plant MetGenMap (http://bioinfo.bti.cornell.edu/cgi-bin/MetGenMAP/home.cgi; Joung et al. 2009), as described in Additional file 4: Figure S2. In both OE-24 and WT-24, most of the DEGs belonged to the function categories of response to endogenous stimulus, cellular process, response to stress, signal transduction, response to abiotic stimulus and response to biotic stimulus (Additional file 4: Figure S2). The distributions of the DEGs in the different function categories were also similar ($r = 0.993$, $\alpha = 0.05$, $n = 40$). More DEGs were identified in OE-24 than in WT-24. In particular, almost all of the functional categories were enriched in more DEGs from OE-24 than WT-24. This result implies that a similar but enhanced biological process is involved in OsHsp18.0-CI-mediated resistance to Xoc as in the basal defense of WT against Xoc.

Of the 898 and 1844 up-regulated genes in WT and OE induced by RS105, a total of 738 genes were common between OE and WT (Fig. 5a). GO analysis of all 738 genes indicated that the distributions of the DEGs in 34 different function categories were similar to the results described in Additional file 5: Figure S3A. Similarly, out of 128 and 154 down-regulated genes in WT and OE induced by RS105, 29 genes in 12 function categories were commonly repressed in both OE and WT (Fig. 5b, Additional file 5: Figure S3B). Of the 738 commonly up-regulated genes, RS105 induced higher expression of 694 genes in OE-24 than in WT-24 (Fig. 5c). Among the 29 commonly down-regulated genes, 16 genes were repressed more strongly by RS105 in OE-24 than in WT-24 (Fig. 5c). In general, most of the common DEGs exhibited greater changes at the transcriptional level in the transgenic plants than in WT post inoculation with RS105. This result is also consistent with an enhanced version of resistance to Xoc in OsHsp18.0-CI-overexpressing plants.

Differential Expression of Pathogenesis-related (PR) Genes and Genes Related to the Biosynthesis of Salicylic Acid and Jasmonic Acid

Many previous reports indicated that the induced expression of pathogenesis-related (PR) genes and
increased content of salicylic acid (SA) and jasmonic acid (JA) are important for enhanced resistance to \textit{Xoc} in rice (Shen et al. 2010; Guo et al. 2012; Feng et al. 2016). Among 738 commonly up-regulated genes and 29 down-regulated genes, we identified a total of 28 genes, including 13 \textit{PRs}, and 8 and 7 genes involved in the biosynthesis of SA and JA, respectively (Fig. 6a). All 13 \textit{PR} genes exhibited higher induction of expression by RS105 in OE-24 than in WT-24 (Fig. 6a). The salicylate biosynthesis and jasmonic acid biosynthesis pathway were significantly changed with a \textit{P} value less than 0.05. All 8 SA biosynthesis-related genes belong to a gene family that encodes putative phenylalanine ammonia-lyases (PALs) involved in SA synthesis. In addition, all of these genes were more highly expressed in OE-24 than in WT-24 and encoded phospholipase A2 (LOC_Os05g51520), lipoygenase (\textit{OsLOX8} and \textit{OsLOX9}), and 12-oxophytodienoate reductase (LOC_Os06g11290, LOC_Os06g11240 and LOC_Os06g11210) (Fig. 6a). Consist with the expression of SA- and JA-biosynthesis-related genes, the content of SA and JA has increased in OE line (CD39R-7) comparing to WT (Fig. 6c and d). In addition, we also identified several DEGs involved in the ethylene or auxin signaling pathway among the 738 commonly up-regulated genes, such as ethylene-responsive transcription factors (LOC_Os04g32620, LOC_Os11g0677, LOC_Os07g47790, LOC_Os04g46400, LOC_Os02g32140, LOC_Os03g08470, LOC_Os01g64790), \textit{OsIAA2} (LOC_Os01g09450) and \textit{OsPILS1} (LOC_Os09g31478) (Additional file 6: Table S3).

In addition to the profiling analysis, the expression levels of ten DEGs were validated by quantitative reverse transcriptase-PCR. The results confirmed the RNA-seq data with regard to the expression patterns of all selected genes (Fig. 6b). In general, compared with WT, all ten genes were more highly expressed in OE post inoculation with RS105 (Fig. 6b).

**Overexpression of \textit{OsHsp18.0-CI} Results in Small Fitness Costs**

Lots of defense-related genes are resulted in enhancing resistance to particular pathogens as well as causing costly plant fitness, such as dwarfism, accelerated senescence, sterility, or reduced seed production (Chen et al. 2014; Liu et al. 2015). To estimate the resistance costs modulating by \textit{OsHSP18.0-CI}, we compared several agronomic traits with wild type Shengdao 806, CD39R-7 (OE) and CD40R-12 (RNAi). Compared to Shengdao 806, the \textit{OsHsp18.0-CI} OE lines exhibited a small decreased phenotype in plant height, 100-grain weight, grain width and grain thickness, but not significant differences in the tillers numbers per plant, grain number per panicle and grain length (Additional file 7: Figure S5). However, the RNAi lines showed no significant differences compared with Shengdao 806 for all tested traits (Additional file 7: Figure S5). Collectively, these results show that the overexpression of \textit{OsHsp18.0-CI} also has a slight negatively impact on rice fitness.

**Discussion**

\textit{OsHsp18.0-CI} Plays Important Roles in Abiotic and Biotic Stresses

HSPs are molecular chaperones that specifically prevent irreversible protein aggregation (Waters 2013). As a large family, HSPs have an important role in thermo tolerance. The expression of HSPs is regulated by heat shock transcription factors (HSFs), which bind to heat shock elements or promoters (Kumar et al. 2009; Scharf et al. 2012). Roles of HSPs in biotic stresses have also been identified, such as \textit{HSP90} and \textit{HSP70} (Park and Seo 2015). sHSPs are the most abundant of HSPs in plants and against environmental stresses such as heavy metals, drought, cold and oxidative stress (Scharf et al. 2012). Some types of biotic stresses also activate the
expression of plant sHSPs (Sarkar et al. 2009; Waters 2013; Li et al. 2015). With the exception of the increased susceptibility of *N. benthamiana* to *Ralstonia solanacearum* induced by silencing of *NtHSP17* (Maimbo et al. 2007), few reports have indicated that plant sHSPs can protect against plant pathogens.

OsHsp18.0-CI is one of 23 sHSP members in rice (Sarkar et al. 2009). OsHsp18.0-CI also belongs to class I sHSPs in the database sHSPdb (Jaspard and Hunault 2016). OsHsp18.0-CI has been reported to function in tolerance to salt and cadmium stresses and interact with the N-terminus of viral RdRp (Cui et al. 2013; Ham et al. 2013; Li et al. 2015). In this study, we found that the overexpression of *OsHsp18.0-CI* enhanced resistance against bacterial pathogenic *Xoc* in rice (Fig. 2 and Fig. 4). These results provide additional evidence that OsHsp18.0-CI plays important roles in both abiotic and biotic stresses.

**OsHsp18.0-CI Positively Regulates Resistance to Xoc in Rice**

Induction of the expression of *OsHsp18.0-CI* by the fungus *M. grisea* and at least seven rice viruses has previously been reported (Sarkar et al. 2009; Li et al. 2015). In the present study, the expression of *OsHsp18.0-CI* was activated by inoculation with a strain of *Xoc*, a bacterial pathogen (Fig. 1). This result indicates that *OsHsp18.0-CI* belongs to the DR genes, which respond to a broad spectrum of pathogens in rice. Consistent with the up-regulation of expression by *Xoc*, overexpression of *OsHsp18.0-CI* increased resistance, whereas repression of the expression of *OsHsp18.0-CI* enhanced susceptibility to *Xoc*. The expression level of *Xoc*-responsive related genes reported previously were also analyzed in the OE transgenic plants, the negatively regulators such as DEPG1, NRRB, OsWRKY45-1 and OsMPK6 have a lower expression level in both non-infected and infected OE lines compared with WT, and the positive regulated gene GH3-2 has a higher level in OE lines than WT (Additional file 8: Figure S4). These results imply that *OsHsp18.0-CI* positively regulates resistance to *Xoc* in rice. Consistent with the positive regulation mechanism, the content of SA and JA (Fig. 6c), the expression of PR genes (Fig. 6a), and the resistance costs (Additional file 7: Figure S5) were slightly higher in *OsHsp18.0-CI* overexpressing plants than in wild-type.

The chromosomal location of *OsHsp18.0-CI* is in the region of 8834823 to 8835802 on chromosome 3 (http://rice.plantbiology.msu.edu). This location coincides with the location of mapped qBlsr3a, a minor QTL for bacterial leaf streak resistance in rice Acc8558 (Tang et al. 2000). Recently, many DR genes that co-localize with QTLs have been identified that may explain the small
effect of these QTLs (Kou and Wang 2010; Feng et al. 2016). Taken together, OsHsp18.0-CI may help explain the effect of qBlsr3a.

**OsHsp18.0-CI Mediates a Basal Defense that Differs from that of Rxo1-mediated Resistance**

Plant have developed two types of pathogen perception. On the external face of the host cell, plants use extracellular pattern recognition receptors (PRRs) to recognize conserved microbial molecules known as pathogen-associated molecular patterns (PAMPs), which induce PAMP-triggered immunity (PTI) or basal defense (Zipfel 2014). The second class of perception involves the recognition of cognate effectors by resistance (R) proteins that induce effector-triggered immunity (ETI; Cui et al. 2015). Both PTI and ETI induce immune responses and consequently modulate the expression of genes in the host, referred to as differentially expressed genes (DEGs) or defense-responsive (DR) genes (Kou and Wang 2010).

To explore the possible resistance mechanism mediated by OsHsp18.0-CI, we compared the differential expression profiles between OsHsp18.0-CI in our study and Rxo1, the only cloned qualitative resistance gene to bacterial leaf streak, which was introduced into japonica rice variety 9804 (Zhao et al. 2005; Zhou et al. 2010). Zhou et al. (2010) determined differential expression profiles using microarray between 9804-Rxo1 and 9804 in response to inoculation with Xoc strain FJR5. In total, 1239 and 963 up-regulated genes were screened out in 9804-Rxo1 and 9804 induced by Xoc. Of those up-regulated genes, only 143 genes were common between the 9804-Rxo1 and 9804 (Zhou et al. 2010). In the present study, of the 1844 and 898 up-regulated genes in OsHsp18.0-CI OE and WT induced by RS105, a total of 738 genes were common between OE and WT (Fig. 5a). Because susceptible wild type is normally considered to only harbor basal defense after inoculation with Xoc, whereas the resistant variety harbors a qualitative resistance gene, the higher ratio of commonly up-regulated genes with wild-type in OsHsp18.0-CI OE plants suggests that the OsHsp18.0-CI-mediated resistance activates a basal defense similar to that found in wild-type, in contrast to the Rxo1-mediated resistance.

Consistent with this difference from to Rxo1-mediated resistance, the distribution of up-regulated genes in different function categories was similar in response to inoculation with Xoc strain RS105 in OsHsp18.0-CI OE plants and wild-type (Additional file 4: Figure S2) but were quite different between 9804-Rxo1 and 9804 (Zhou et al. 2010). As further evidence, 120 and 79 differentially regulated transcription factor (TF) genes were identified in the 9804-Rxo1 and 9804 plants infected by Xoc strain FJR5, respectively. Among them, only 7 TFs were commonly identified in both 9804-Rxo1 and 9804 (Zhou et al. 2010; Additional file 9: Table S4). In this study, 119 and 71 TF genes were differentially expressed in OsHsp18.0-CI OE and WT, and a total of 50 TFs were commonly identified in both OsHsp18.0-CI OE and WT (Additional file 9: Table S4). Among the 119 and 120 differentially expressed TFs in OsHsp18.0-CI OE and 9804-Rxo1, 104 and 105 TFs were specifically up-regulated or repressed in OsHsp18.0-CI OE and 9804-Rxo1, respectively. Fifteen TFs were common DEGs in OsHsp18.0-CI OE and 9804-Rxo1. Among them, only 7 TFs exhibited the same expression pattern, and 8 exhibited opposite expression patterns in OsHsp18.0-CI OE and 9804-Rxo1. For example, OsWRKY45, a negative regulator of BLS resistance (Tao et al. 2009), exhibited down-regulated expression in OsHsp18.0-CI OE but up-regulated expression in 9804-Rxo1.

Several DR genes have been implicated in BLS resistance by inducing the expression of PR genes and increasing the content of SA and JA (Tao et al. 2009; Shen et al. 2010; Guo et al. 2014; Feng et al. 2016). After inoculation with RS105, we detected significantly elevated expression of many genes, including SA and JA biosynthesis-related genes and 10 PR genes (Fig. 6). Interestingly, the expression of these genes was also elevated in WT after inoculation with RS105, but the expression levels were lower than in OsHsp18.0-CI OE. Together with the above evidence, these results indicate that OsHsp18.0-CI-mediated resistance occurs in a basal defense manner.

**Conclusions**

Our study revealed that OsHsp18.0-CI positively regulates resistance to bacterial leaf streak in rice. Resistance mediated by the overexpression of OsHsp18.0-CI may prime an enhanced version of basal defense after inoculation with Xoc.

**Methods**

**Plant Materials and Growth Condition**

A japonica BLS-susceptible rice variety, Shengdao 806, which is widely planted in Shandong Province of China, was used in this study. Seeds were kindly provided by Dr. Fangying Yao of Shandong Academy of Agricultural Science (Jinan, Shandong, China). All rice plants were grown in greenhouse at a temperature of 28 ± 2°C, relative humidity of 85-100%, and photoperiod of 16 h, as described previously (Feng et al. 2016).

**Vector Construction and Rice Transformation**

Genomic DNA was isolated from the leaves of Shengdao 806 with a Plant DNA Extraction Kit (CWBio, Beijing, China). Because there are no introns in Os03g16030 (OsHsp18.0-CI), a 657-bp DNA containing the coding sequence of OsHsp18.0-CI was amplified by PCR using
the specific primers OsHsp18.0-CI-1 F and OsHsp18.0-CI-1R (Additional file 10: Table S5). The PCR product was digested with BamHI and Kpn I (New England Biolabs, MA, USA) and cloned into the binary vector pU1301 cleaved with BamHI and Kpn I to prepare the construct for overexpressing OsHsp18.0-CI (pU1301-OsHsp18.0-CI) as described previously (Li et al. 2013).

To construct a RNAi vector suppressing the expression of OsHsp18.0-CI, the expected 480-bp DNA fragment containing a portion of the encoding region and the 3’ UTR was obtained by PCR using the primers OsHsp18.0-CI-2 F and OsHsp18.0-CI-2R (Additional file 10: Table S5). The appropriate restriction sites were introduced into the PCR product for subsequent cloning steps (SpeI and Kpn I at the 5’ end and SacI and BamHI at the 3’ end). The PCR product was first cloned into Kpn I/BamHI I-digested ds1301 vector and then into SpeI/SacI I-digested ds1301 to obtain ds1301-OsHsp18.0-CI as described previously (Li et al. 2013). The PCR product was inserted approximately 1.1 kb from the intron of the rice alcohol dehydrogenase (Adh) gene.

The constructs of pU1301-OsHsp18.0-CI (OE) and ds1301-OsHsp18.0-CI (RNAi) were introduced into Shendao 806 using the standard Agrobacterium-medi ed transformation system described previously (Li et al. 2013). Transgenic plants were selected with hygromycin for two constructs in our studies. OE individuals were further validated by PCR with the forward primer UbiF (Additional file 10: Table S5) derived from the sequence of the ubiquitin promoter and the reverse primer OsDRXoc8-1R (Additional file 10: Table S5). For RNAi individuals, positive plants were identified using a pair of specific primers, ds1301-F and ds1301-R, which were described previously (Feng et al. 2016).

Pathogen Inoculation and Disease Assessment

The virulent Xoc strain RS105, JLB2-24, HNB8-47, HGA1 and HGA2 were grown on polypeptone-sucrose-agar medium (10 g l⁻¹ polypeptone, 1 g l⁻¹ glutamic acid, 10 g l⁻¹ sucrose and 15 g l⁻¹ agar) at 28 °C for 2 days and then resuspended in sterile 10 mM MgCl₂ to OD₆₀₀ = 0.5. More than five newly expanded leaves were infiltrated at three positions with a non-needle syringe at the seedling stage (Liu et al. 2014). The lesion length was scored at 14 days post-inoculation (dpi). The population count experiments were progressed as previously described (Trippett et al. 2016), rice leaves of OE, RNAi and WT plants harvested at three specific time points (0, 4, 8 dpi).

RNA Extraction and Real-time RT-PCR

Total RNA was isolated from 100 mg of rice leaves using TRI Reagent (Sigma Aldrich, USA) following the procedures described in the manual. First-strand cDNA synthesis was performed using a HiFiScript gDNA Removal cDNA Synthesis Kit (CWMP, Beijing, China) according to the recommended protocols. Quantitative real-time PCR was performed on a QuantStudio™ 6 Flex Real-Time System (Applied Biosystems, USA) with UltraSYBR Mixture (CWMP, Beijing, China).

The following PCR program was used: 95 °C for 30 s, followed by 40 cycles of 95 °C for 5 s, 55 °C for 20 s, and 72 °C for 30 s. A heat dissociation curve (55-95 °C) was checked after the final PCR cycle to determine the specificity of the PCR amplification. The gene expression levels relative to the rice OsActin (LOC_Os03g50890) gene were analyzed using the 2⁻ΔΔCt analysis method. Genes expression levels were analyzed by qRT-PCR assays, which were repeated at least twice with triplicate runs. The primer sequences for each detected gene are listed in Additional file 10: Table S5. The primers qRT-1 F and qRT-1R were used to check the expression levels of OsHsp18.0-CI in the OsHsp18.0-CI-OE lines, and for the OsHsp18.0-CI RNAi plants, the primers qRT-2 F and qRT-2R were used.

RNA-seq and Analysis

RNA samples were collected from non-infected and infected leaves of Shendao 806 and Shendao 806 (CD39R-7), respectively. Library construction and sequencing were performed on a BGISEQ-500 by Beijing Genomic Institute (www.genomics.org.cn, BGI, Shenzhen, China). Clean-tags were mapped to the reference genome and genes available at the Rice Genome Annotation Project (http://rice.plantbiology.msu.edu) with a perfect match or one mismatch. The original sequence data have been submitted to the database of the NCBI Sequence Read Archive (http://trace.ncbi.nlm.nih.gov/Traces/sra) under the accession number SRP079496.

For gene expression analysis, the matched reads were calculated and then normalized to RPKM using RESM software (Li and Dewey 2011). The significance of the differential expression of genes was defined by the bioinformatics service of BGI according to the combination of the absolute value of log2-Ratio ≥1 and FDR ≤ 0.001. KOG functional classification, Gene Ontology (GO) and pathway annotation and enrichment analyses were based on the NCBI COG (https://www.ncbi.nlm.nih.gov/COG/), Gene Ontology Database (http://www.geneontology.org/) and KEGG pathway database (http://www.genome.jp/kegg/), respectively. The software Cluster and Java Treeview were used for hierarchical cluster analysis of gene expression patterns (de Hoon et al. 2004; Saldanha 2004).

Quantification of Phytohormone

The leaves of four-week-old seedlings were used for phytohormone quantification. Each of 100 ug samples were
prepared and quantified using the HPLC-MS/MS system as reported by Xu et al. (2016). Three biological replicates were analyzed.

Statistical Analysis
As described previously (Feng et al. 2016), standard deviations were checked visually by error bars, and statistical significance was determined by analysis of variance. The data were subjected to one-way analysis of variance, and the mean differences were compared by paired t test. P values <0.05 were considered significant. Correlation analysis was performed using SPSS software.

Additional files

Additional file 1: Table S1. Summary of the sequence assembly after RNA-seq. (DOCX 13 kb)

Additional file 2: Table S2. List of DEGs in OE under non-infected conditions. (XLSX 29 kb)

Additional file 3: Figure S1. GO functional annotation of DEGs in OE and WT under non-infected conditions. (TIF 1894 kb)

Additional file 4: Table S3. List of DEGs involved in the ethylene or polygalacturonase-inhibiting protein 4 (OsPGIP4) pathway. (DOCX 14 kb)

Additional file 5: Figure S2. The numbers of DEGs in different function categories for 738 commonly up-regulated genes (A) and 29 commonly down-regulated genes (B) between WT and OE induced by RS105. (TIF 2568 kb)

Additional file 6: Table S4. List of the TF genes differentially regulated under non-infected conditions. (TIF 3560 kb)

Additional file 7: Table S5. Summary of the sequence assembly after RNA-seq. (DOCX 13 kb)

Additional file 8: Figure S3. The numbers of DEGs in different function categories under infection with RS105. (TIF 2022 kb)

Additional file 9: Table S6. Primers used in this study. (DOCX 15 kb)

Additional file 10: Figure S4. Effects of OsHsp18.0-CI in transgenic plants. (A) Comparison of transgenic lines and wild type Shengdao 806 at the tillering stage, bar = 10 cm. (B-D) Grain morphology of transgenic lines and Shengdao 806, bar = 10 mm. (E) Plant height of transgenic lines and Shengdao 806 at the tillering stage, mean values were calculated from measurement on at least 10 individuals. (F) Number of tillers per plant in the wild-type and transgenic plants, Data were obtained from at least 10 individuals. (G) Number of grains per panicle in the wild-type and transgenic plants, mean values were calculated from measurement on at least 20 individuals. (H) Phenotype statistics of seeds from WT and transgenic plants, mean values were calculated from measurement on at least 20 individuals. The bars represent the means ± SD. *** indicate extremely significant differences between wild-type and transgenic plants at *P* = 0.01 by Student’s t test. (TIF 5760 kb)

Additional file 11: Table S7. The numbers of DEGs in different function categories for 738 commonly up-regulated genes (A) and 29 commonly down-regulated genes (B) between WT and OE induced by RS105. (TIF 2568 kb)

Abbreviations
BLS: Bacterial leaf streak; DEG: Differentially expressed gene; dpi: Days post inoculation; DR: Defense-related; hpi: Hours post inoculation; HSPs: Heat shock proteins; OE: Overexpression; PR: Pathogenesis related; WT: Wild-type; Xoc: Xanthomonas oryzae pv. oryzae

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Availability of Data and Materials
The data sets supporting the results of this article are included within the article and its additional files. The nucleic acid sequence of OsHsp18.0-CI is available in NCBI’s database with accession number NM_001056196 (http://www.ncbi.nlm.nih.gov/nucleotide). The RNA-seq data supporting the results of this article are available in the NCBI’s SRA with the accession numbers SRP079496 (https://trace.ncbi.nlm.nih.gov/Traces/sra_sub/sra.cgi?acc=SRP079496).

Authors’ Contribution
ZC and XD designed the experiments. YJ, HT and LZ performed the experiments. YJ and ZC analyzed data. RZ, GJ and XL help to field management. QX help to quantify the content of JA and SA. JY and ZC wrote the manuscript. All authors approved the manuscript.

Competing Interests
The authors declare that they have no competing interests.

Consent for Publication
Not applicable.

Ethics Approval and Consent to Participate
Not applicable.

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