A High Throughput Barley Stripe Mosaic Virus Vector for Virus Induced Gene Silencing in Monocots and Dicots

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Abstract

Barley stripe mosaic virus (BSMV) is a single-stranded RNA virus with three genome components designated alpha, beta, and gamma. BSMV vectors have previously been shown to be efficient virus induced gene silencing (VIGS) vehicles in barley and wheat and have provided important information about host genes functioning during pathogenesis as well as various aspects of genes functioning in development. To permit more effective use of BSMV VIGS for functional genomics experiments, we have developed an Agrobacterium delivery system for BSMV and have coupled this with a ligation independent cloning (LIC) strategy to mediate efficient cloning of host genes. Infiltrated Nicotiana benthamiana leaves provided excellent sources of virus for secondary BSMV infections and VIGS in cereals. The Agro/LIC BSMV VIGS vectors were able to function in high efficiency down regulation of phytoene desaturase (PDS), magnesium chelatase subunit H (ChlH), and plastid transketolase (TK) gene silencing in N. benthamiana and in the monocots, wheat, barley, and the model grass, Brachypodium distachyon. Suppression of an Arabidopsis orthologue cloned from wheat (TaPMR5) also interfered with wheat powdery mildew (Blumeria graminis f. sp. tritici) infections in a manner similar to that of the A. thaliana PMR5 loss-of-function allele. These results imply that the PMR5 gene has maintained similar functions across monocot and dicot families. Our BSMV VIGS system provides substantial advantages in expense, cloning efficiency, ease of manipulation and ability to apply VIGS for high throughput genomics studies.

Introduction

Virus-induced gene silencing (VIGS) is a particularly useful tool for plant functional genomics because it permits knockdown of genes-of-interest and observation of elicited phenotypes within 3 to 4 weeks. VIGS avoids production of knockout mutants or stable RNA interference (RNAi) and can also be performed on species that are difficult to transform [1,2,3,4]. The VIGS principle is based on antiviral responses that target RNAs for degradation and is triggered by accumulation of double-stranded RNAs (dsRNA) appearing in the infection cycle [5,6]. During the early, or “shock”, phase of infection intense symptoms develop and high levels of virus accumulate. As host defense mechanisms are activated, infections usually transit into a protracted chronic phase characterized by modulated symptoms and low virus abundance. By inserting sequence fragments derived from “genes-of-interest” into VIGS vectors, the corresponding mRNAs are selectively degraded during virus infection to result in transient silencing of the targeted gene.

The first VIGS vectors were derived from Tobamoc mosaic virus (TMV) [7], Potato virus X (PVX) [8,9] and Tobacco rattle virus (TRV) [10,11], and these vectors were initially used for Nicotiana benthamiana and tomato (Solanum lycopersicum) gene silencing. Recently, the plants in which VIGS has been employed has increased substantially, and more than 30 viruses have been shown to have potential as VIGS vectors. Examples include Apple latent spherical virus (ALSV) [12,13], Bean pod mottle virus (BPMV) [14], Brome mosaic virus (BMV) [15,16], Pea early browning virus (PEBV) [17], Rice tungro bacilliform virus [18], Tomato bushy stunt virus [19], Turnip yellow mosaic virus (TYMV) [20], and some Geminiviruses and their satellite DNAs [21,22].

Barley stripe mosaic virus (BSMV) vectors suitable for cereal VIGS have also been described [23,24,25,26,27,28,29]. BSMV is a positive-sense RNA virus with a broad experimental host range and is the type member of the Hordeivirus genus [30,31]. The tripartite genome consists of RNAs α, β, and γ, and each of the genomic [γ] RNAs has a methylated 5’ cap and a 3’ polyadenylate sequence followed by a tyrosine accepting tRNA-like structure. RNAβ of the BSMV ND18 strain encodes the methyltransferase/helicase subunit of the RNA-dependent RNA polymerase (RdRp). RNAβ specifies the coat protein (CP) and three major triple gene block (TGB) proteins (TGB1, TGB2 and TGB3) that are essential for cell-to-cell movement of the virus [30,31]. RNAγ encodes the polymerase (GDD) subunit of the RdRp and the γb protein, which...
has significant roles in viral pathogenesis, long distance movement and suppression of host RNA silencing defenses [30,31,32,33]. BSMV was first modified as a VIGS vector [23] for use in barley (Hordeum vulgare), and subsequently was used to down regulate expression of wheat (Triticum aestivum) genes [28]. Applications of BSMV-based VIGS vectors include functional genomics research in wheat [29,34,35,36], barley [25,26,27,29,37], Daucus carota (Heinola villosa) [38,39], a distant relative of wild wheat relative, Brachypodium distachyon [40,41], a model organism for cereals, and the culinary ginger (Zingiber officinale), a tropical monocot [42]. In these studies, fragments from genes-of-interest were initially inserted into RNA of monocot [42]. In these studies, fragments from genes-of-interest Zingiber officinale were inserted into RNA of monocot [42]. In these studies, fragments from genes-of-interest (CaMV) 35S promoter immediately downstream of the cDNAs. Biolistic introduction of cDNAs in Ti plasmids for initiation of BSMV infections upon infiltration of leaves resulted in replication of BSMV immediately downstream of the cDNAs. Biolistic introduction of these plasmids into barley and wheat, but none of these resulted in BSMV infections. Other inoculation methods were also tested without success, including mechanical inoculation with circular or linearized plasmids (Fig. 2A). After infiltration with equal concentrations of pCaBS-a, pCaBS-B and pCaBS-γ, the upper uninfiltrated N. benthamiana leaves developed mild mottling symptoms at 7 to 10 dpi. Western blotting revealed high levels of coat protein, and PCR analyses verified the presence of the gRNAs (data not shown), indicating systemic movement from the infiltrated leaves (Fig. 2B).

Different Agrobacterium strains (EHA105, EHA101 and GV3101, harboring pCaBS-α, pCaBS-β and pCaBS-γ) were used to infiltrate barley and wheat, but none of these resulted in BSMV infections. Other inoculation methods were also tested without success, including mechanical inoculation with circular or linearized plasmids, and vacuum infiltration of germinated seeds or seedlings with the Agrobacterium derivatives. Consequently, infiltrated N. benthamiana leaf sap was used to inoculate wheat and barley. More than 80% of the uninoculated emerging leaves of both cereals consistently developed typical mosaic symptoms at 6 to 7 dpi, and the presence of BSMV RNA and CP was confirmed by RT-PCR and Western blots (Fig. 2C). The inbred B. distachyon line Bd21-3 also consistently developed a mild systemic phenotype (Fig. 2C), similar to that shown in more limited earlier studies [41]. Moreover, in all cases BSMV symptoms elicited by the PMR5 for powdery mildew resistance 5 gene (TaPMR5) fragment whose Arabidopsis thaliana PMR5 (APMR5) loss-of-function allele elicits resistance to two powdery mildew fungi, Golovinomyces (formerly Erysiphe) cichoracearum and G. orontii, [63]. Transient down regulation of PMR5 compromises the ability of wheat powdery mildew (B. graminis f. sp. tritici) to establish infections, and illustrates the cross kingdom requirements of PMR5 for powdery mildew invasion of monocot and dicot hosts.

Results

Agrobacterium mediated BSMV infectivity of N. benthamiana, wheat, barley and B. distachyon

The three BSMV ND18-cDNAs [64] were cloned into the pCass4-Rz T-DNA vector under the control of a double 35S promoter [58] to permit precise in vivo transcription at the 5′ terminus of the gRNAs. The cDNA α was inserted into the Sυ site of pCass4-Rz, and the cDNA β and γ clones were integrated between Sυ and BamHI sites to produce pCaBS-α, pCaBS-β and pCaBS-γ (Fig. 1A). The TRSV satellite RNA ribozyme mediated cis-cleavage extending 33, 17 and 17 nts non-viral sequences beyond the 3′ ends of RNAα, β and γ respectively.

To confirm the infectivity of pCaBS-α, pCaBS-β and pCaBS-γ, the plasmids were mixed and rub-inoculated on Chenopodium amaranticolor leaves. Large necrotic lesions similar to those resulting from BSMV RNAs or purified BSMV inoculum appeared at 7 to 10 days post-inoculation (dpi) only on leaves abraded with either circular or linearized plasmids (Fig. 2A). After infiltration with equal concentrations of A. tumefaciens EHA105 mixtures harboring pCaBS-α, pCaBS-β and pCaBS-γ, the upper uninfiltrated N. benthamiana leaves developed mild mottling symptoms at 7 to 10 dpi. Western blotting revealed high levels of coat protein, and PCR analyses verified the presence of the gRNAs (data not shown), indicating systemic movement from the infiltrated leaves (Fig. 2B).

The first generation BSMV VIGS systems under the control of the T7 promoter [2] were time consuming and expensive because capped in vitro transcripts from the α, β and γ cDNA clones are required for plant inoculation. The latter problem has been circumvented in the Wise lab [26,27] by engineering plasmids containing each of the cDNAs, and incorporating a double Cauliflower mosaic virus (CaMV) 35S promoter immediately upstream of the cDNAs and a Hepatitis delta virus (HDV) ribozyme immediately downstream of the cDNAs. Biolistic introduction of these plasmids into barley leaves resulted in replication of BSMV VIGS derivatives containing candidate genes predicted to affect powdery mildew resistance, and sap from the systemically infected leaves was suitable for secondary inoculations to other cereals. However, a limitation of the Wise method is the potential instability of gene inserts that can occur during systemic invasion of the primary inoculated plants [55,56,57]. Another limitation of both strategies is that cloning multiple genes-of-interest can be time-consuming and inefficient when high throughput operations are required.

To permit more effective use of BSMV VIGS for functional genomics experiments, we have developed an approach with Agrobacterium tumefaciens strains harboring the BSMV α, β, and γ cDNAs in Ti plasmids for initiation of BSMV infections upon infiltration of N. benthamiana leaves. The Agrobacterium mediated BSMV VIGS vectors were engineered by inserting BSMV cDNAs between the double 35S promoter and a ribozyme sequence (Rz) from Tobacco ringspot virus (TRSV) satellite RNA [58]. In addition, we have inserted a ligation independent cloning (LIC) site similar to that used for TRV VIGS [39] into BSMV to facilitate efficient cloning of desired gene fragments. These two modifications permit rapid VIGS analyses of large numbers N. benthamiana plants, as we have demonstrated with phytotoxine desaturase (PDS) and plastid transketolase (TK) genes [60]. Infiltrated N. benthamiana leaves also accumulated high levels of BSMV and provided excellent sources for secondary infections to elicit VIGS in wheat, barley, and the model grass, B. distachyon [61,62]. We have also used BSMV Agro/ LIC VIGS to clone and analyze a wheat powdery mildew resistance 5 gene (TaPMR5) fragment whose Arabidopsis thaliana PMR5 (APMR5) loss-of-function allele elicits resistance to two powdery mildew fungi, Golovinomyces (formerly Erysiphe) cichoracearum and G. orontii, [63]. Transient down regulation of PMR5 compromises the ability of wheat powdery mildew (B. graminis f. sp. tritici) to establish infections, and illustrates the cross kingdom requirements of PMR5 for powdery mildew invasion of monocot and dicot hosts.

A High Throughput BSMV-Based VIGS Vector

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demonstrate that agroinfiltrated leaves of *N. benthamiana* are excellent sources of BSMV VIGS inoculum for high throughput dissection of wheat, barley and *B. distachyon* genes involved in disease development and pathogenesis.

**A LIC strategy for high-throughput VIGS**

One limitation of *in vitro* transcription and DNA based BSMV VIGS vectors [25,27] is the relatively inefficient cloning of target genes needed for efficient high throughput VIGS operations. To circumvent this limitation, a LIC site (5′-GAAGGGCCGG-TGGTGAGGTG-3′) with an *Apa*I site (underlined) was inserted into pCaBS-γ at three positions downstream of the γa (polymerase subunit) gene. The plasmid pCa-γbLIC was engineered by integrating the LIC site immediately upstream of the stop codon of the γb gene, to provide a functional γb protein with an extension of two C-terminal residues (Glu and Val) after inserting target fragments (Fig. 1B and Fig. 3). For pCa-LICγb, the LIC site was engineered to replace the AUG start codon of the γb gene and eliminate γb protein expression, and in the case of pCa-LICΔγb, the LIC site was substituted for the γb gene. The primer pairs BS-10 and BS-32 (Table S1), whose positions are indicated above pCa-γbLIC, were used for PCR amplifications to assess the stability of fragments inserted into the pCa-γbLIC vectors. (C) Western blot analysis of BSMV infections after agroinfiltration of *N. benthamiana* leaves. Analyses of infiltrated leaves are shown at 6 dpi to illustrate virus accumulation before spread to the vascular system. Upper uninfected leaves and roots were assessed at 15 dpi to compare the accumulation of BSMV pCaBS-γ and its three LIC derivatives. Protein was isolated from two plants agroinfiltrated with pCaBS-α and pCaBS-β mixtures containing pCaBS-γ (wt-γb), pCa-γbLIC (γbLIC), pCa-LICγb (LICγb) or pCa-LICΔγb (LICΔγb). BSMV coat protein was detected with BSMV antisera and healthy *N. benthamiana* leaves (H) were used as negative controls. The bottom panel shows a loading control consisting of Coomassie Blue stained Rubisco, the major protein associated with tobacco leaf extracts. doi:10.1371/journal.pone.0026468.g001

Figure 1. Construction of *Agrobacterium* mediated BSMV VIGS vectors with LIC cloning sites and comparisons of BSMV accumulation in *N. benthamiana* tissues infected with wild type BSMV and its three LIC derivatives. (A) Schematic representation of pCaBS-α, pCaBS-β and pCaBS-γ. *Agrobacterium* VIGS plasmids were engineered by inserting the respective BSMV α, β and γ cDNAs between the double CaMV 35S promoter and a ribozyme sequence (Rz) derived from TRSV satellite RNA. (B) The organization of three LIC derivatives derived from pCaBS-γ. The LIC site was inserted into pCaBS-γ at three positions downstream of the γa gene. The pCa-γbLIC plasmid was designed to express a full-length γb protein containing two C-terminal amino acid (Glu and Val) extensions preceding the VIGS target sequence. For pCa-LICγb, the LIC site was engineered to replace the AUG start codon of the γb gene and eliminate γb protein expression, and in the case of pCa-LICΔγb, the LIC site was substituted for the γb gene. The primer pairs BS-10 and BS-32 (Table S1), whose positions are indicated above pCa-γbLIC, were used for PCR amplifications to assess the stability of fragments inserted into the pCa-γbLIC vectors. (C) Western blot analysis of BSMV infections after agroinfiltration of *N. benthamiana* leaves. Analyses of infiltrated leaves are shown at 6 dpi to illustrate virus accumulation before spread to the vascular system. Upper uninfected leaves and roots were assessed at 15 dpi to compare the accumulation of BSMV pCaBS-γ and its three LIC derivatives. Protein was isolated from two plants agroinfiltrated with pCaBS-α and pCaBS-β mixtures containing pCaBS-γ (wt-γb), pCa-γbLIC (γbLIC), pCa-LICγb (LICγb) or pCa-LICΔγb (LICΔγb). BSMV coat protein was detected with BSMV antisera and healthy *N. benthamiana* leaves (H) were used as negative controls. The bottom panel shows a loading control consisting of Coomassie Blue stained Rubisco, the major protein associated with tobacco leaf extracts.
derivatives can replicate in infiltrated tissue, γb expression is important for efficient vascular movement in *N. benthamiana*, as shown previously [65,66]. The efficiency of LIC cloning into the pCa-γbLIC site was also confirmed by recovery of PDS gene fragments from different plant species. In these experiments, 43 of 44 clones had inserts corresponding in size to the targeted inserts (data not shown). Therefore, pCa-γbLIC was chosen for all subsequent BSMV VIGS experiments.

**Agrobacterium mediated BSMV VIGS in *N. benthamiana***

* N. benthamiana is the most widely used experimental dicot host for VIGS because of its susceptibility to a large number of plant viruses and the rapidity with which visible phenotypes appear [67]. To assess silencing by *Agrobacterium* mediated BSMV VIGS, a 370 bp PDS (NbPDS) fragment and a 400 bp TK (NbTK) fragment from *N. benthamiana* were cloned into pCa-γbLIC in the sense orientations to generate pCa-γb:NbPDS<sub>370</sub> and pCa-γb:NbTK<sub>400</sub> respectively. Then, the four-leaf stage of *N. benthamiana* was infiltrated with *Agrobacterium* mixtures containing pCaBS-α, pCaBS-β and pCa-γb:NbPDS<sub>370</sub> or pCa-γb:NbTK<sub>400</sub>.

Leaves infiltrated with pCaBS-α, pCaBS-β and pCa-γb:NbPDS<sub>370</sub> to elicit PDS silencing developed a mottled photobleaching phenotype in the 3th or 6th leaves at 9 to 10 dpi, and about 5 days later (15 dpi) larger areas of more uniform white PDS silencing were observed on the 6 to 8 leaf stage (data not shown). PDS silencing was most pronounced at 30 to 45 dpi when larger areas of white photobleaching were evident in most leaves and, in many cases, on stems and petioles (Fig. 4A). Leaf and stem silencing persisted until the leaves began to senesce, but silencing in younger emerging leaves began to gradually dissipate as green sectors at the tips and margins of the leaves became increasingly evident.

The TK gene, used for *N. benthamiana* VIGS, encodes an enzyme required for photosynthesis and phenylpropanoid metabolism [60]. When pCa-γb:NbTK<sub>100</sub> was co-infiltrated with pCaBS-α, pCaBS-β, an obvious chlorosis became evident at 9 to 10 dpi on the 5th or 6th emerging leaves (data not shown). The chlorosis gradually transitioned into a more generalized silencing phenotype and by 23 dpi all upper uninfiltrated leaves displayed yellowing chlorotic sectors with a distinctive carotenoid coloration interspersed with pale green regions (Fig. 4B) that appear to be similar to the TK knockdown phenotype noted in previous studies [68].

Overall, silencing was efficient with the NbPDS and NbTK genes and, in most experiments, distinctive symptoms appeared in >80% of the infiltrated plants grown under our growth chamber conditions. The lower transcript abundances of the NbPDS and NbTK genes in *N. benthamiana* infected with BSMV:NbPDS<sub>370</sub> or BSMV:NbTK<sub>400</sub> respectively were confirmed by semi-quantitative RT-PCR and by use of a tobacco 18S rRNA as an internal control (Fig. 4C). To the best of our knowledge, this is the first report of BSMV-based VIGS in *N. benthamiana*, and it is notable that the high efficiency and intensity of the silencing phenotypes are comparable to those triggered by other VIGS vectors [4]. Hence, our results demonstrate that BSMV VIGS is a viable choice for use in conjunction with *N. benthamiana* genomics studies.

**BSMV accumulation and insert stability in infiltrated *N. benthamiana* leaves***

Because a high percentage of plants consistently developed gene silencing phenotypes after agroinfiltration of *N. benthamiana*, we wished to determine the accumulation of BSMV to provide a basis for efficient transfer of BSMV vectors suitable for VIGS elicitation in secondary plants. When the first four to eight leaves above the *N. benthamiana* cotyledons were infiltrated, the levels of BSMV coat protein increased for the first 6 days and then leveled off and

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**Figure 2.** *Agrobacterium* mediated BSMV infection of *N. benthamiana*, wheat, barley and *B. distachyon*. (A) Appearance of large necrotic local lesions on leaves of *C. amaranticolor* at 10 days after mechanical inoculation with either circular (right) or linearized (left) plasmids. CP shows western blots of BSMV CP isolated from two separate leaves. (B) *N. benthamiana* leaves infiltrated with equal concentrations of *Agrobacterium* mixtures harboring pCaBS-α, pCaBS-β, and pCaBS-γ. Infiltrated leaves at 10 dpi (left) and upper uninfiltrated leaves developing mild mottling symptoms at 15 dpi (right). CP is as designated panel A. (C) Typical BSMV symptoms on systemically infected leaves of wheat, barley and *B. distachyon* emerging at 14 dpi after inoculation of plants with infected *N. benthamiana* leaf sap. The presence of BSMV RNA and CP was confirmed by RT-PCR and Western blots. doi:10.1371/journal.pone.0026468.g002

46.2% of pCaBS-γ). Moreover, at 15 dpi only pCa-γbLIC accumulated to the same levels as wild-type pCaBS-γ in the upper uninfiltrated leaves, and only wt pCaBS-γ and pCa-γbLIC were detected in the upper uninfiltrated leaves and the roots (Fig. 1C). These results indicated that although the three BSMV
remained relatively constant for the next six days (Fig. 5A). Moreover, several grams of tissue were recovered from the infiltrated leaves and extracts from this tissue were able to provide >80% infections of cereals used for VIGS tests (data not shown). These results demonstrate that *N. benthamiana* amplification of BSMV from primary infiltrated leaves provides an inexpensive inoculum source for VIGS trials of a large number of plants.

To compare the stability of inserts in infiltrated and systemically infected leaves at different times after infection, *N. benthamiana* leaves were infiltrated with pCa-γb:TaPDS<sub>400</sub>. Then at 12 dpi, the next four fully expanded leaves above the infiltrated leaf, and the fifth emerging leaf, which was still expanding, were harvested. The primer pair BS-10/BS-32 flanking the LIC site (Fig. 1B; All primers used in this work are listed in Table S1) was

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**Figure 3. Schematic representation of use of the pCa-γbLIC vector for target gene fragment cloning.** Individual steps in LIC cloning and insertion of target fragments to be used for VIGS assessment are illustrated. doi:10.1371/journal.pone.0026468.g003
RT-PCR products of the empty pCa-ybLIC vector appeared in the 5th emerging smaller leaves. Moreover, degradation products were more pronounced in upper systemically infected leaves at 20 dpi (Fig. 5C). At this time, the infiltrated leaves were becoming necrotic, so we were unable to isolate intact RNA at later stages. Taken together, these results suggest that infiltrated leaves harvested within 12 days after BSMV infection contain a high abundance of intact inserts with negligible amounts of degradation products, and should be the most suitable source for secondary inoculations for VIGS tests. This is particularly important for VIGS analyses with larger inserts because infiltrated leaves maximizes foreign sequence stability and minimizes loss of inserts occurring during long distance BSMV movement [55,56,57].

**Agrobacterium mediated BSMV VIGS in cereals**

Although we were unable to agroinfect cereals directly, *N. benthamiana* infiltrated leaves provided an excellent source for secondary inoculations to wheat, barley, and the model grass, *B. distachyon* (Fig. 3; Table S2). Using LIC cloning techniques, a series of *PDS* or magnesium chelatase subunit H (*ChlH*) target gene fragments of different lengths from each species to be tested were cloned into the LIC site of pCa-ybLIC (Fig. 3). When the first two emerging leaves of wheat, barley, and *B. distachyon* were inoculated with infected *N. benthamiana* sap, silencing phenotypes appeared 10 to 14 dpi on upper leaves emerging after infection that corresponded to the cognate *PDS* or *ChlH* sequences were used for silencing (Fig. 6; Fig. S1; Table S2). Leaves of all three cereal species targeted for down regulation of *PDS* developed a longitudinal white streaks that appeared to be devoid of carotenoid pigment. These photo-bleached segments occupied various portions of the parallel venation and were interspersed within green regions of the leaf blades. In the case of the *ChlH* phenotype, a more intense yellowing sometimes covered large areas of the leaf, but the yellowing was often interspersed with green sections. In contrast, BSMV or BSMV:00 infected leaves developed typical mosaic symptoms consisting of mild light yellow chlorotic streaks (Fig. 6 and Fig. S1). In all three species, the virus symptoms could be easily distinguished visually from the *PDS* or *ChlH* phenotypic effects. In most cases, a high proportion of the plants (>80%) developed the silencing phenotype as illustrated in Fig. 6B, which shows several pots of wheat at 14 dpi, in which >90% of the plants exhibit the chlorotic phenotype associated with suppression of *TaChlH* transcripts.

To determine the effects of fragment lengths on silencing efficiency, we evaluated phenotypes induced by 200 and 400 bp *TaPDS* fragments, and 250, 300 and 547 bp *TaChlH* fragments in wheat, a 300 bp *HvChlH* fragment, 300 and 400 bp *HvPDS* fragments in barley, and 102, 303 and 402 bp *HvPDS* fragments in *B. distachyon* (Fig. 6). The 300 and 400 bp *HvPDS* fragments appeared to elicit similar levels of photobleaching in barley leaves (Fig. 6A). The extent of silencing in wheat appeared to be substantially more efficient with the 250 bp and 300 bp *TaChlH* inserts than the 547 bp insert. In these cases, the two smaller inserts consistently induced more uniform bright yellowing phenotypes than the 547 bp insert, which often elicited a mottled phenotype on emerging leaves. The results of semi-quantitative RT-PCR also show that the *TaChlH* transcript abundance was higher in BSMV:*TaChlH*47 infected wheat than BSMV:*TaChlH*300 or BSMV:*TaChlH*547 infections, and that *TaPDS* RNA levels were reduced to similar levels in BSMV:*TaPDS*250 or BSMV:*TaPDS*300 infected leaves (Fig. 6B). In another comparison, the 102 bp *BdPDS* fragment was less effective than the 303 bp or 402 bp

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**Figure 4. Silencing of endogenous phytoene desaturase (*PDS*) and plastid transketolase (*TK*) genes in *N. benthamiana* by *Agrobacterium* mediated BSMV VIGS.** The four-leaf stage of *N. benthamiana* was infiltrated with an *Agrobacterium* mixture containing pCaBS-α, pCaBS-β and pCa-yb:NbPDS370 (BSMV:NbPDS370) or pCa-yb:NbTK400 (BSMV:NbTK400). (A) Large areas of photobleaching occurring in leaves infected with BSMV:NbPDS370 40 dpi, and an intense white photobleaching often appearing in stems and petioles (see magnified inset panel). (B) Chlorotic sectors with a distinctive carotenoid coloration interspersed with pale green regions in upper uninfected leaves of plants infected at 23 dpi with BSMV:NbTK400. A leaf of a plant infiltrated with the empty vector (BSMV:00) is shown in the inset panel. (C) Relative transcript levels of *NbPDS* and *NbTK* in leaves infected with the BSMV:00, BSMV:NbPDS370 or BSMV:NbTK400. RNA extracted from the leaves was subjected to semi-quantitative RT-PCR amplification (27 cycles) with the gene-specific oligonucleotide primers shown in Table S1. Amplified tobacco 18S rRNA served as an internal control.

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used for RT-PCR of RNA extracted from each leaf. Although agarose gels revealed that the inserted sequence is relatively stable in the infiltrated leaves, a distinct shadow below the major 1350 bp band became progressively more intense in the upper systemically infected leaves 1 to 4 (Fig. 3B). In addition, an obvious smaller band of ~590 bp, whose size is similar to the
fragments in induction of photobleaching in *B. distachyon*, and these photobleaching phenotypes were consistent with the *BdPDS* transcript abundance assayed by semi-quantitative RT-PCR (Fig. 6C). These results indicate that sequences ranging from 200 to 400 bp provide effective silencing with BSMV VIGS, and that the gene silencing is specific for the cognate gene fragments used to induce silencing and for the anticipated silencing phenotype [2,3,24].
The potential of Agrobacterium mediated BSMV VIGS for analyses of genes involved in disease development was assessed by investigating a orthologue of the Arabidopsis PMR5 gene, which is known to interfere with powdery mildew (Golovinomyces cichoracearum and G. cichoracearum) colonization of A. thaliana [63].

To determine the possible presence of a wheat orthologue of PMR5, we carried out a BLAST search with the A. thaliana PMR5 gene sequence and found a 337 bp region in common wheat with a 60.5% nucleotide sequence identity. Because this 337 bp region has 100% identity in the wheat Chinese Spring and Xuezao lines, and shares 98.8% identity with a wheat EST sequence (GenBank accession: CV759357), we predicted that the fragment might be a wheat orthologue of the A. thaliana PMR5 gene and provisionally designated the putative gene TaPMR5.

To determine the effects of TaPMR5 on wheat powdery mildew (B. graminis f. sp. triticum) infections, a 170 bp fragment was amplified from wheat DNA with the primer pair PMR5-8/PMR5-10 (Table S1) and inserted into pCa-cbLIC in the sense orientation to generate pCa-cb:TaPMR5170. This derivative was infiltrated into N. benthamiana, and sap from the infected leaves was inoculated to the first two emerging leaves of the highly susceptible wheat line Xuezao to assess the effects on mildew pathogenesis (Fig. 7). BSMV:00 (pCa-cbLIC) was tested as a vector alone control, and BSMV:TaPDS200 was used to confirm viral infection and spread.

The potential effects of TaPMR5 silencing on powdery mildew infections were assessed at 3 weeks after BSMV:00 or BSMV:TaPMR5170 inoculation of Xuezao plants. Colonies emerged on all plants by 4 days after dusting with conidia, but the number of colonies on BSMV:TaPMR5170 inoculated leaves were obviously lower than those on the BSMV:00 treated leaves at 4 and 5 days after mildew infection (Fig. 8A). By 11 days after conidia applications, most BSMV:00 treated plants had wilted and displayed extensive mildew colonization and growth, whereas the BSMV:TaPMR5170 treated plants were more robust and had much lower levels of mildew development (data not shown).

TaPMR5 mRNA suppression in plants used for the mildew trials were evaluated by using the primer pair TaPMR5-7/TaPMR5-14 in RT-PCR reactions to amplify a 337 bp sequence from RNA of plants infected for 3 weeks with the three BSMV VIGS derivatives.

**Figure 6. PDS and magnesium chelatase subunit H (ChlH) gene silencing phenotypes in barley, wheat and B. distachyon. (A) Barley showing phenotypes typical of suppression of ChlH and PDS by BSMV:HvChlH or BSMV:HvPDS inserts of the indicated lengths. (B) Upper panel: Wheat inoculated with BSMV:TaChlH250 showing more than 90% of inoculated plants developing the chlorotic phenotype associated with suppression of the ChlH gene. Middle panel: PDS and ChlH gene silencing phenotypes on wheat leaves infected with BSMV:TaChlH and BSMV:PDS derivatives with different length inserts. Bottom panel: PCR amplification of transcripts from leaves shown in the middle panel. (C) B. distachyon leaves showing effects of PDS suppression after infection with BSMV:8dPDS inserts. In these experiments, plants were inoculated at the two-leaf stage with infected N. benthamiana sap harboring BSMV derivatives targeting PDS and ChlH cognate genes from each species. Leaf photographs were taken at 14 dpi, and fragment lengths from each source are indicated as subscripts above each leaf. Relative transcript levels of PDS and ChlH genes in the leaves infected with the different BSMV derivatives are shown under the leaf photographs. RNA extracted from the leaves was subjected to semi-quantitative RT-PCR amplification (28 cycles for wheat and barley, 31 cycles for B. distachyon) with the gene-specific oligonucleotide primers shown in Table S1. Amplified species-specific 18S rRNA served as internal controls for each species.

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**Functional analysis of a gene required for powdery mildew invasion**

The potential of Agrobacterium mediated BSMV VIGS for analyses of genes involved in disease development was assessed by investigating a orthologue of the Arabidopsis PMR5 gene, which is known to interfere with powdery mildew (Golovinomyces cichoracearum and G. cichoracearum) colonization of A. thaliana [63]. To determine the possible presence of a wheat orthologue of PMR5, we carried out a BLAST search with the A. thaliana PMR5 gene sequence and found a 337 bp region in common wheat with a 60.5% nucleotide sequence identity. Because this 337 bp region has 100% identity in the wheat Chinese Spring and Xuezao lines, and shares 98.8% identity with a wheat EST sequence (GenBank accession: CV759357), we predicted that the fragment might be a wheat orthologue of the A. thaliana PMR5 gene and provisionally designated the putative gene TaPMR5.

To determine the effects of TaPMR5 on wheat powdery mildew (B. graminis f. sp. triticum) infections, a 170 bp fragment was amplified from wheat DNA with the primer pair PMR5-8/PMR5-10 (Table S1) and inserted into pCa-cbLIC in the sense orientation to generate pCa-cb:TaPMR5170. This derivative was infiltrated into N. benthamiana, and sap from the infected leaves was inoculated to the first two emerging leaves of the highly susceptible wheat line Xuezao to assess the effects on mildew pathogenesis (Fig. 7). BSMV:00 (pCa-cbLIC) was tested as a vector alone control, and BSMV:TaPDS200 was used to confirm viral infection and spread.

The potential effects of TaPMR5 silencing on powdery mildew infections were assessed at 3 weeks after BSMV:00 or BSMV:TaPMR5170 inoculation of Xuezao plants. Colonies emerged on all plants by 4 days after dusting with conidia, but the number of colonies on BSMV:TaPMR5170 inoculated leaves were obviously lower than those on the BSMV:00 treated leaves at 4 and 5 days after mildew infection (Fig. 8A). By 11 days after conidia applications, most BSMV:00 treated plants had wilted and displayed extensive mildew colonization and growth, whereas the BSMV:TaPMR5170 treated plants were more robust and had much lower levels of mildew development (data not shown).

TaPMR5 mRNA suppression in plants used for the mildew trials were evaluated by using the primer pair TaPMR5-7/TaPMR5-14 in RT-PCR reactions to amplify a 337 bp sequence from RNA of plants infected for 3 weeks with the three BSMV VIGS derivatives.
As an internal control, the primer pair SQ18S-1/SQ18S-2 was used to amplify wheat 18S rRNA. The results show that leaves infected with BSMV:00, BSMV:TaPDS$^{200}$ or BSMV:TaPMR$^{170}$ had similar levels of 18S rRNA transcripts (Fig. 8B). However, the TaPMR5 transcript abundance was much lower in BSMV:TaPMR$^{170}$ infected plants than in the BSMV:00 or BSMV:TaPDS$^{200}$ infections, and TaPDS RNA levels were specifically reduced in BSMV:TaPDS$^{200}$ infected leaves (Fig. 8B). These results demonstrate that suppression of TaPMR5 transcripts occurs only during infections with BSMV:TaPMR$^{170}$ sequences and provides an additional basis for the specificity of TaPDS VIGS.

For statistical evaluations of the mildew infections, the 4th emerging wheat leaves were cut into 4 cm long segments, placed in a Petri dish and dusted with powdery mildew conidia (Fig. 8C). The numbers of infection foci and hyphal growth between 3 and 4 days after conidial dusting revealed that leaves inoculated three weeks previously with BSMV:TaPMR$^{170}$ were significantly less susceptible to mildew infections than those infected with BSMV:00. In three sets of independent experiments in which more than 500 colonies were counted and measured for each treatment, the average numbers of colonies on leaf segments that had been infected with BSMV:00 and BSMV:TaPMR$^{170}$ respectively were 294.37±82.16 μm and 267.44±79.92 μm respectively at 3 days after dusting with significant differences (P<0.01). The average colony sizes increased to 604.82±122.51 μm and 306.22±126.66 μm at 4 days after dusting, with significant (P<0.0001 differences) between the two groups (Fig. 8D). In sum, significant colony growth occurred between 3 and 4 days after application of conidia to leaf sections infected with BSMV:00, whereas foci in BSMV:TaPMR$^{170}$ infected leaves exhibited no significant differences in growth (Fig. 8C). Taken together, the data presented above indicate that down regulation of the TaPMR5 gene has a substantial negative effect on rates of wheat powdery mildew infection, as was also the case with the previous A. thaliana powdery mildew infections [63].

**Discussion**

BSMV VIGS vectors provide a reliable and efficient VIGS system and are now contributing to substantial advances in functional genomics research in a diverse range of monocot hosts [3]. The most frequent studies have focused on the natural host, barley, and on wheat, which is occasionally infected in the field. However, short communications have shown that BSMV VIGS can be used for down regulation of genes in B. distachyon [40,41].
and *Daucus vulgaris* (H. villosa) [38,39], as well as the more distantly related monocot, culinary ginger (Z. officinale) [42]. In addition to these plants, selective BSMV strains can be mechanically transferred to a wide range of experimental hosts, including maize (Z. mays), rice (Oryza sativa), hexaploid oat (Avena sativa) and diploid oat (A. strigosa) [41,69]. Hence, it is likely that BSMV VIGS can be applied more widely to other crop species. In this regard, we (Li Lab members; unpublished) have shown that BSMV ND18 infected barley sap can be used to establish systemic infections of millet (Setaria italica), several inbred maize lines and other *Triticum* species. Among these, wild emmer (*Triticum turgidum* var. dicoccoides, AABB, 2n = 4X = 28), the progenitor of cultivated tetraploid wheat, has particular promise as a donor of powdery mildew resistance genes [70]. Crosses of wild emmer with cultivated tetraploid durum wheat developed obvious photo-bleaching phenotypes on emerging leaves at 14 dpi when wheat *PDS* sequences (*TaPDS* or *TaPDSb*) were used for BSMV VIGS (data not shown) and this finding permits the powdery mildew resistance in these hybrids to be evaluated in more detail. In addition, use of the Agro/LIC silencing strategy should permit more effective and widespread applications of BSMV VIGS for high-throughput functional genomics of sorghum (*Sorghum bicolor*), *B. distachyon*, and other cereals, and to assist in evaluating gene functions before cross species introgression or transformation.

Among dicots, *N. benthamiana* is being developed rapidly for molecular genetic and genomics analyses, and our results show that BSMV VIGS provides another tool for functional genomics of this host. In addition to *N. benthamiana*, BSMV has been reported to elicit systemic infections in spinach (*Spinacia oleracea*) and a limited number of other dicots [69]. Primarily inoculated leaves of *Chenopodium* species, tobacco (*N. tabacum*) and beet (*Beta vulgaris*) also develop local infections [69] that might serve to initiate VIGS. These results suggest that the BSMV Agro/LIC strategy has potential for high-throughput functional genomics of susceptible dicots, as well as untested members of the Zingiberales and possibly other monocot families. Hence, BSMV VIGS appears to be well positioned to contribute to a more refined understanding of plant biology in the future [31].

Infections of barley and wheat with *in vitro* transcripts [25,28] and bombardment with BSMV plasmids [26,27] has been used with considerable success with VIGS studies of host genes involved in disease responses. In order to improve on these methods for large-scale functional genomics, we have developed agroinfiltration to permit secondary BSMV transfers from infiltrated *N. benthamiana* leaves and have introduced a LIC insertion site to facilitate efficient gene cloning. The silencing efficiencies of the resulting BSMV Agro/LIC VIGS vectors is comparable to those of the first generation systems [3], and the inserts appear to be very stable in infiltrated leaves. In addition to being a useful amplification host for BSMV, *N. benthamiana* can be grown under a wide range of temperature, humidity and light intensity conditions, and the 4 to 8 leaves that can be infiltrated on a single plant accumulate high levels of BSMV from 6 to 12 dpi under our growth conditions. In addition, infiltrated leaves or sap can be frozen or used to purify recombinant BSMV derivatives for subsequent experiments. Moreover, the inexpensive leaf sap inoculation procedure is very convenient and can be used to facilitate reliable and stable VIGS analyses of large numbers of plants. Taken together, our data indicate that the BSMV Agro/LIC VIGS strategy provides substantial advantages over previous BSMV VIGS systems in expense, cloning efficiency, ease of manipulation and ability to rapidly scale up preliminary experiments for high throughput genomics trials. To facilitate the use of the BSMV VIGS vectors, we will honor all requests for plasmids and other resources, and we encourage interested researchers to contact DL (Dawei.Li@cau.edu.cn) for their distribution. We also plan to deposit the vectors into a resource comparable to that of the Arabidopsis Biological Resource Center (ABRC) as soon as a comparable resource becomes available for cereal genomics.

Host susceptibility genes (S-genes) have been intensively studied in recent years to better understand the pathogenesis of powdery mildew fungi, *G. cichoracearum* and *G. cichoracearum* [63]. PMR5 also affects rice blast (Magnaporthe oryzae) infections in *A. thaliana* [72], but to date has not been reported to be involved in disease development of other fungal or bacterial pathogens. Other PMR genes identified in the *Arabidopsis* screen, such as *PMR4* and *PMR6* [63,73,74], also contribute to microbe-plant interactions in a resistance gene (R-gene) independent manner, and have been proposed to be S-genes [71,75,76]. The recessive *PMR5* gene encodes a novel protein that localizes in the endoplasmic reticulum and is predicted to participate in pectin biosynthesis, modification or deposition. This has lead to suggestions that *PMR5* (and *PMR6*, which is a putative wall degrading enzyme) may alter wall composition sufficiently to interfere with critical host defense pathways or signaling and recognition requirements for hyphal penetration [77].

To identify potential PMR5 cereal orthologous, we conducted a BLAST search with the 337 bp *TaPMR5* sequence, which revealed the presence of related cDNA or EST sequences in wheat, barley, rice, maize, sorghum, *B. distachyon*, fescue species (*Festuca arundinacea*), and a wild rye interspecific hybrid (*Leymus cinererus* x *Leymus triticoides*). Multiple alignments of the 337 bp sequence revealed >90% sequence identity in these cereals when the *A. thaliana* *PMR5* gene and the Chinese cabbage orthologue provided a control outgroup (Fig. 9). These results support previous suggestions that *PMR5*, like other PMR genes involved in powdery mildew invasion [78], represents an ancient lineage whose functions have been conserved across the plant kingdom over ~200 million years since divergence of the monocots and dicots [79].
Our results clearly show that down regulation of TaPMR5 in a susceptible wheat line has a pronounced interference with the early stages of powdery mildew infection. Moreover, the resulting disease phenotype appears to mimic that previously noted with the 
\textit{Apmr5} mutant [63]. Both the \textit{Apmr5} allele and VIGS down regulation of \textit{TaPMR5} result in reduced hyphal growth during the initial stages of infection and in reduced sporulation at later stages of powdery mildew infection, rather than blocking specific infection steps. Although additional work needs to be carried out to more thoroughly define \textit{TaPMR5} regulatory interactions in cereal disease pathogenesis and roles in development, \textit{Arabidopsis} studies have revealed that \textit{Apmr5} mutant cells are significantly smaller than wild type cells and chemical analyses showed elevated pectin and possibly aberrant pectin cross-linking [63]. Cell wall composition has become increasingly implicated in pathogen infection and defense responses, and the available evidence suggests that \textit{PMR5} along with other \textit{PMR} genes have diverse roles in formation of wall complexity elements needed for powdery mildew infections [77]. Although, the pectin composition of Type 1 cell walls of dicots differs substantially from Type 2 cell walls of the grasses [80,81], our current results suggest that certain of these features have been conserved during evolution, and that BSMV Agro/LIC VIGS can provide a valuable tool for more refined understanding of the roles of these elements and the functions of “loss-of-susceptibility” orthologous during pathogenesis.

**Materials and Methods**

**Plant material and growth conditions**

\textit{N. benthamiana} plants were grown in a controlled environmental climate chamber at 20–25°C with a 14/10 h light (~75 μmol/m².s)/dark photoperiod. Wheat (Yangmai 11), barley (Black Hulless) and \textit{B. distachyon} (Bd21-3) plants used for VIGS experiments were grown in a greenhouse until the two leaf-stage, then inoculated with BSMV and transferred to a climate chamber at 23–25°C until evaluated. For powdery mildew experiments, the highly susceptible common wheat line Xuezao was inoculated with infected \textit{N. benthamiana} sap and grown at 23–25°C under a 16/8 h light (~160 μmol/m².s)/dark regimen.

**Construction of BSMV VIGS vectors with LIC cloning sites**

The \textit{Cass4-Rz} [58] \textit{Agrobacterium} plasmid was used for construction of VIGS vectors. Full-length \(\alpha\), \(\beta\) and \(\gamma\) cDNAs were amplified from BSMV ND18 clones [64] with high-fidelity \(Pfu\) DNA polymerase (Tiangen, Beijing, China) with the primer pairs BS-22/BS-32, BS-23/BS-26, or BS-24/BS-26, respectively (Table S1). The cDNA \(\alpha\) clone was inserted into the \textit{StuI} site, and the cDNA \(\beta\) and \(\gamma\) clones were integrated between the \textit{StuI} and \textit{BamHI} sites of \textit{pCass4-Rz}, and the resulting clones were confirmed by sequencing, and designated pCaBS-\(\alpha\), pCaBS-\(\beta\) and pCaBS-\(\gamma\) (Fig. 1A).

A LIC site (5'-TAAGGAAGGGCCGGTGGTGTGGTT-A-3') was generated by inserting a 21 nt sequence (5'-GAA-GGGCCGGTGGTGTGGTT-3') containing an Apol restriction enzyme site (underlined sequence) into pCaBS-\(\gamma\) at the 3' terminus of the \(\gamma\) gene to facilitate efficient cloning. The LIC vector was constructed by amplifying pCaBS-\(\gamma\) DNA with the primer pair BS-24/LIC-1 to produce fragment 1 containing cDNA residues 1–2536 fused to the 3' LIC sequence upstream of the \textit{Apol} site. A second amplification with the primer pair LIC-2/BS-32 generated fragment 2, whose sequence consisted of the \textit{Apol} site, the downstream 12 nt LIC sequence, and cDNA residues 2537–2790. Fragments 1 and 2 were then digested with \textit{Apol} and \textit{BamH}I, mixed in equal amounts with \textit{StuI} and \textit{BamH}I-digested \textit{pCass4-Rz}, and ligated to produce pCa-\(\gamma\)-LIC\(b\) (Fig. 1B). Similar strategies with different primer pairs (Table S1) were used to construct pCa-LIC\(b\)\(\gamma\) and pCa-LIC\(b\)\(\alpha\).

The LIC strategy [59], was used to clone a series of \(PDS, \textit{Ta}\textit{ChlH}, \textit{Ta}\textit{PDS}\) and \(\textit{magnesiun chelatase subunit H} \textit{(ChlH)} [82]\) fragments. The selected sequences were amplified with primer pairs, in which the forward primers consisted of 5'-AAGGAGGTTAA-3' fusions to a 5' plant sequence, and the reverse primers contained 5'-AACCACCACCCGGT-3' fusions to each 3' gene sequence. PCR products were purified, treated at room temperature with T4 DNA polymerase (New England Biolabs, MA, USA) in 1X reaction buffer containing 5 mM dATP to generate sticky ends, and heat-treated at 75°C for 10 min to inactivate the polymerase. The \textit{Apol}-linearized pCa-\(\gamma\)-LIC\(b\)\(\gamma\) vector was also treated with T4 DNA polymerase in the presence of 5 mM dTTP to generate sticky ends complementary to those of the PCR products. The PCR products (~200 ng) and pCa-\(\gamma\)-LIC\(b\)\(\gamma\) vector (20 ng) were mixed, incubated at 66°C for 2 min and slowly cooled to room temperature to anneal their complementary termini. Then, 10 μl aliquots were used for cloning into \textit{Escherichia coli DH10B}, and transformants identified by colony PCR and endonuclease digestions.

**BSMV gene silencing derivatives for VIGS assessment**

The \textit{N. benthamiana} 370 bp \(PDS\) (\(Nb\)\(PDS\); GenBank accession: AJ571700) and 400 bp plastid transketolase (\(\textit{PDS}\); GenBank accession: HQ200305) fragments were obtained by RT-PCR using the \(Nb\)\(PDS\)-1/\(Nb\)\(PDS\)-2 and \(Nb\)\(TK\)-1/\(Nb\)\(TK\)-2 primer pairs respectively (Table S1). The resulting fragments were integrated into pCa-\(\gamma\)-LIC\(b\) in the sense orientations to generate pCa-\(\gamma\)-\(Nb\)\(PDS\)\(_{270}\) and pCa-\(\gamma\)-\(Nb\)\(TK\)\(_{400}\). In a similar fashion, 200 and 400 bp wheat \(PDS\) (\(Ta\)\(PDS\); GenBank accession: FJ517553), 250, 300 and 547 bp wheat \(\textit{ChlH}\) (\(Ta\)\(ChlH\); TIGR accession: TC169257), 300 bp barley \(\textit{ChlH}\) (\(Hv\)\(ChlH\); GenBank accession: U26545), 300 and 400 bp barley \(PDS\) (\(Hv\)\(PDS\); GenBank accession: AY062039), and 102, 303 and 402 bp \(B. \textit{distachyon}\) \(PDS\) (\(Bd\)\(PDS\); GenBank accession: HQ317869) fragments were amplified by RT-PCR with appropriate primer pairs (Table S1) and inserted into pCa-\(\gamma\)-LIC\(b\).

**Agroinfiltration of \textit{N. benthamiana} and viral inoculation of cereals**

The BSMV pCa-\(\gamma\)-LIC\(b\)\(\gamma\) derivatives were transformed into \textit{A. tumefaciens} strain EHA105. For agroinfiltration, single colonies
were grown overnight at 28°C with constant shaking in 3 ml of LB containing rifampicin (25 μg/ml) and kanamycin (100 μg/ml). Then, 0.5 ml of the cultures were used to inoculate 30 ml LB containing the same antibiotics and grown at 28°C for 10 to 12 h. Bacterial cells were pelleted at 2200 g for 10 min, resuspended in infiltration buffer [10 mM MgCl₂, 10 mM 2-(N-morpholino)ethanesulfonic acid (MES), pH 5.2, and 0.1 mM acetylsyringone] to ~0.7 OD₆₀₀ and incubated at room temperature for at least 3 h. For agroinfiltration, equal amounts of bacteria harboring pCaBS-w, pCaBS-b, and pCa-ybLIC (or its derivatives) were mixed, and infiltrated into four to eight N. benthamiana leaves immediately above the cotyledons with a 1-ml needleless syringe. After maintenance in a growth chamber for 5 to 12 days post infiltration (dpi), the infiltrated leaves were harvested, ground in 20 mM Na-phosphate buffer (pH7.2) containing 1% celite, and the sap was mechanically inoculated onto the two-leaf stages of wheat, barley and B. distachyon.

RNA extraction and semi-quantitative RT-PCR analysis
Total RNA was extracted with TRIzol reagent as described by the manufacturer (Invitrogen, Carlsbad, CA, USA), treated with DNase, and first-strand cDNA synthesis carried out with M-MLV reverse transcriptase (Promega, Madison, WI, USA). Target gene expression levels were monitored by semi-quantitative RT-PCR using gene-specific primers that anneal outside the region targeted for silencing (Table S1). The wheat 18S rRNA gene (GenBank accession: AF490940), barley 18S rRNA gene (GenBank accession: AK251731), B. distachyon 18S rRNA (GenBank accession: DV483201), and tobacco 18S rRNA (GenBank accession: HQ384692) were used as individual internal RNA controls for appropriate assays.

Powdery mildew inoculations
At 3 weeks after inoculation with N. benthamiana leaf sap, wheat developing visible BSMV symptoms (stripe mosaic on the 3th leaf with a mild mosaic on the 4th leaf) was challenged with B. graminis f. sp. tritici isolate E09, a prevalent pathotype in the Beijing area, by dusting test plants with conidia from infected wheat. Plants were maintained as described above until mildew observations were recorded. For detached leaf observations of VIGS effects, the 4th leaf exhibiting BSMV symptoms was cut into ~4 cm segments, dusted with mildew conidia and incubated in a growth chamber for 5 days in Petri dishes containing 0.7% (w/v) agar and 40 ppm benzimidazole in distilled water.

Staining and Microscopy of Powdery Mildew Infected Wheat
Leaves used for mildew observations were stained according to the Wise lab protocol [26,27] with minor modifications. Leaves were fixed in 75% ethanol/25% glacial acetic acid overnight, and subsequently rinsed twice with deionized water. Then, tissue was immersed in Coomassie blue stain (0.05% Coomassie blue R250, 10% acetic acid, 50% methanol and 40% H₂O) for 10–15 min, and examined immediately or stored in buffer (20% glycerol and 5% acetic acid) for up to 48 h before observation with an Olympus IX71 microscope.

References

VIGS efficacy and data collection
To score susceptibility of VIGS treated wheat leaves, visible colonies were counted and their sizes determined by microscopy at 3 and 4 days after mildew inoculation. Colony numbers per visual field were calculated, the longest axis of the colonies was measured and the data were subjected to statistical analyses. Differences between the groups were evaluated by using the Student’s t-test, where P<0.01 was considered statistically significant.

Supporting Information
Figure S1 Comparison of BSMV symptoms with PDS and ChlH silencing phenotypes at 10 to 14 days after inoculation. (A) Barley leaves to illustrate uninoculated plants (Mock), with BSMV systemic mosaic on the upper emerging leaves of plants inoculated with BSMV:00, and the PDS and ChlH silencing phenotypes on leaves of plants inoculated with BSMV:He:PDS₃₀₀ and BSMV:He:ChlH₃₀₀, respectively. Compare the faint yellow BSMV mosaic with the more expansive PDS white photobleaching phenotype and the intense yellowing phenotype elicited by ChlH silencing. (B) Comparison of systematically infected Brachypodium distachyon plants showing BSMV symptoms and the PDS silencing phenotype. Note that the upper leaves of the BSMV-infected plant have a mild chlorotic mosaic whereas the PDS silenced plant inoculated with BSMV:BdPDS₃₀₃ exhibits intense white chlorotic streaks on the second, third and fourth leaves emerging above the inoculated leaves. (C) B. distachyon leaves from uninoculated plants, mild systemic mosaic symptoms and PDS phenotype on leaves of plants inoculated with BSMV:00 and BSMV:BdPDS₃₀₃.

Table S1 Primers used in vector construction and molecular analyses.

Table S2 Infection characteristics and VIGS phenotype details in cereals inoculated with N. benthamiana sap from infected leaves infiltrated with BSMV Agro/LIC VIGS derivatives.

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Author Contributions
Conceived and designed the experiments: DL, AOJ. Performed the experiments: CY, CL, LY. Analyzed the data: ZL, CH, JY. Wrote the paper: DL, AOJ, CY, CL.


