Eukaryotic translation initiation factor 4E family member nCBP facilitates the accumulation of TGB-encoding viruses by recognizing the viral coat protein in potato and tobacco

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Due to their limited coding capacity, plant viruses have to depend on various host factors for successful infection of the host. Loss of function of these host factors will result in recessively inherited resistance, and therefore, these host factors are also described as susceptibility genes or recessive resistance genes. Most of the identified recessive resistance genes are members of the eukaryotic translation initiation factors 4E family (eIF4E) and its isoforms. Recently, an eIF4E-type gene, novel cap-binding protein (nCBP), was reported to be associated with the infection of several viruses encoding triple gene block proteins (TGBps) in Arabidopsis. Here, we, for the first time, report that the knockdown of nCBP in potato (StnCBP) compromises the accumulation of potato virus S (PVS) but not that of potato virus M (PVM) and potato virus X (PVX), which are three potato viruses encoding TGBps. Further assays demonstrated that StnCBP interacts with the coat proteins (CPs) of PVS and PVM but not with that of PVX, and substitution of PVS CP in the PVS infectious clone by PVM CP recovered the virus infection in StnCBP-silenced transgenic plants, suggesting that the recognition of PVS CP is crucial for StnCBP-mediated recessive resistance to PVS. Moreover, the knockdown of nCBP in Nicotiana benthamiana (NbnCBP) by virus-induced gene silencing suppressed PVX accumulation but not PVM, while NbnCBP interacted with the CPs of both PVX and PVM. Our results indicate that the nCBP orthologues in potato
and tobacco have conserved function as in Arabidopsis in terms of recessive resistance against TGB-encoding viruses, and the interaction between nCBP and the CP of TGB-encoding virus is necessary but not sufficient to determine the function of nCBP as a susceptibility gene.

KEYWORDS
susceptibility genes, recessive resistance, eukaryotic translation initiation factors 4E, nCBP, TGB-encoding virus, coat protein

Introduction

Potato (Solanum tuberosum L.) is currently the third most important food crop worldwide next to rice and wheat for direct human consumption and food security (Hardigan et al., 2017; van der Waals and Krüger, 2020). However, plant viruses are one of the main limiting factors leading to a loss in quality and quantity of potato (Wang et al., 2011; Rashid et al., 2021). Due to their limited coding capacity, plant viruses recruit various host factors to help their infection (Truniger and Aranda, 2009). These host factors, whose mutation confers loss of susceptibility to related viruses, are called susceptibility genes or recessive resistance genes (Truniger and Aranda, 2009; Hashimoto et al., 2016; Garcia-Ruiz, 2018; Mäkinen, 2020).

The eukaryotic translation initiation factor 4E (eIF4E) and its isoforms [eIF(iso)4E] are mRNA 5’ cap-binding proteins which control cap-dependent translation initiation (Gingras et al., 1999). They are also the most extensively studied susceptibility genes (Lelis et al., 2002), and particular research attention has been paid to the relationship between eIF4E/eIF(iso)4E and potyvirus infection. For instance, mutation in eIF4E/eIF(iso)4E resulted in loss of susceptibility to potyviruses in multiple hosts, such as Arabidopsis, tomato, pepper, lettuce, pea, peanut, and sugarcane (Ruffel et al., 2002, 2005; Niacase et al., 2003; Gao et al., 2004; Xu et al., 2017; Yang et al., 2021). In most cases, the interaction between eIF4E/eIF(iso)4E and the viral genome-linked proteins (VPgs) of potyviruses is required for virus infection (Léonard et al., 2000; Charroin et al., 2008; Taivert-Roudet et al., 2017; Yang et al., 2021). Moreover, eIF4E/eIF(iso)4E was also found to mediate recessive resistance to other viruses. The recessive mutant conl in Arabidopsis showed reduced accumulation of a cucumovirus, and conl was found to encode an eIF4E protein (Yoshii et al., 2004). The recessive bymovirus resistance locus rym4 in barley was proved to correspond to the eIF4E gene (Kanyuka et al., 2005). In melon, an eIF4E allele confers resistance to a carmovirus, depending on the inefficient interaction between the viral 3’ cap-independent translational enhancer and eIF4E (Nieto et al., 2006; Truniger et al., 2008). In addition, the mutation of tobacco eIF(iso)4E genes was found to reduce the susceptibility to an umbravirus (Udagawa et al., 2020).

Another less studied eIF4E family member, novel cap-binding protein (nCBP), has been identified in Arabidopsis showing 41.44 and 40.50% nucleotide sequence identity with eIF4E and eIF(iso)4E, respectively, (Ruud et al., 1998). The nCBP protein also supports the translation initiation of capped mRNA in vitro and in vivo (Bush et al., 2009). In addition, nCBP has been reported to be involved in the infection of different viruses in various crops. ncbp-1/ncbp-2 mutants in cassava exhibited high resistance to cassava brown streak disease (Gomez et al., 2019), and the ncbp mutant in Arabidopsis displayed resistance to several viruses in Alphaflexiviridae and Betaflexiviridae, which encode triple gene block proteins (TGBps; Keima et al., 2017). These results indicate that nCBP, similar to eIF4E/eIF(iso)4E, has the potential to act as a recessive resistance gene, which is worthy of further investigation in other crops.

Potato virus S (PVS, Carlavirus), potato virus M (PVM, Carlavirus), and potato virus X (PVX, Potexvirus), which all encode TGBps, are important viruses posing great threats to potato production (Wang et al., 2011, 2016; Keima et al., 2017). In this study, we found that StnCBP knockdown compromised PVS accumulation but not PVM and PVX. Further analysis demonstrated that the interaction between StnCBP and PVS coat protein (CP) is crucial for the StnCBP-mediated recessive resistance to PVS, while StnCBP interacts with the CPs of both PVS and PVM. Moreover, NbnCBP knockdown significantly reduced PVX accumulation but not PVM, and NbnCBP also interacted with both their CPs in N. benthamiana. These results indicate that the interaction between nCBP and CP is necessary but not sufficient for facilitating the accumulation of TGB-encoding viruses, which improves the understanding of the underlying mechanism for nCBP to act as a susceptibility gene.

Materials and methods

Plant materials and virus isolates

Potato variety (Eshu 3) and three transgenic lines with StnCBP silencing were used in the study. The plants were maintained on MS medium (Murashige and Skoog, 1962) supplemented with 4% sucrose in vitro under a 16 h of
light/8 h of dark photoperiod with 400–1,000 μmol photons m\(^{-2}\) s\(^{-1}\) light intensity at 20°C and were transplanted into pots (12 cm) containing premixed soil in the greenhouse located in Huazhong Agricultural University (HZAU) under a 12 h of light/dark photoperiod with 90 μmol photons m\(^{-2}\) s\(^{-1}\) light intensity at 18–22°C.

Three virus isolates used in this study, PVS-HB7, PVX-HB3, and PVM-HB36, were collected from a local potato virus survey and maintained in tobacco or potato host plants under greenhouse conditions at HZAU. The virus identity and purity were determined as described previously (Nie et al., 2012; Wang et al., 2016).

BLASTn search and cloning

A BLASTn search was performed using the AtnCBP (Accession number: AF028809) nucleotide sequence as a query against the S. tuberosum genome (Spud DB) and the Solanaceae Genomics Network (SGN) to retrieve nCBP homolog sequences in potato and N. benthamiana, respectively. StnCBP and NbnCBP were amplified with specific primers (Supplementary Table 1) from the potato cultivar Eshu 3 and N. benthamiana, respectively, using Phanta Super-Fidelity DNA polymerase (Vazyme, China). Then, the PCR amplicons were cloned into the pCE-Zero vector (Vazyme, China). Five colonies were randomly selected and sent for sequencing (Sangon Biotech, China). To analyze the phylogenetic relationships of eIF4E orthologues in S. tuberosum, Arabidopsis thaliana, Solanum lycopersicum, and Nicotiana benthamiana, we performed a BLASTp search against the NCBI and SGN databases using the amino acid sequences of potato eIF4E family members as queries. The phylogenetic tree was generated using MEGA5.2 with neighbor-joining tree of potato and N. benthamiana, respectively, using Phanta Super-Fidelity DNA polymerase (Vazyme, China). Then, the PCR amplicons were cloned into the pCE-Zero vector (Vazyme, China). Five colonies were randomly selected and sent for sequencing (Sangon Biotech, China). To analyze the phylogenetic relationships of eIF4E orthologues in S. tuberosum, Arabidopsis thaliana, Solanum lycopersicum, and Nicotiana benthamiana, we performed a BLASTp search against the NCBI and SGN databases using the amino acid sequences of potato eIF4E family members as queries. The phylogenetic tree was generated using MEGA5.2 with neighbor-joining tree and 1000 bootstraps. The nCBP orthologues in Oryza sativa and Zea mays were also obtained by a BLASTp against the NCBI database. Then, an amino acid sequence alignment of nCBP orthologues from the above species was performed by DNAMAN to analyze whether the nCBP orthologues are conserved across species.

Generation of transgenic potato lines

Based on the design principles of RNAi fragments (Wesley et al., 2001; Reynolds et al., 2004), a fragment with 249 bp near the 5’ end of StnCBP gene was selected as the target fragment for RNA interference. The nucleotide sequence of the target fragment returned no other homologous genes except for StnCBP in a BLASTn search against the S. tuberosum genome. The fragment was recombined into the XhoI and XbaI sites of the pHellsGate8 vector. Agrobacterium tumefaciens (A. tumefaciens) strain GV3101 containing the recombined pHellsGate8 constructs was employed to transform the microtuber slices of Eshu 3 for transgenic lines as previously described (Zhou et al., 2019). Primers for constructing the RNAi vector are shown in Supplementary Table 1.

Virus-induced gene silencing assay

A 270-bp region in the 5’-terminus of NbnCBP was selected as the virus-induced gene silencing (VIGS) target using SGN VIGS Tool1 (Fernandez-Pozo et al., 2015). Then, it was amplified and inserted into the EcoRI and BamHI sites of pTRV2 vector (named pTRV2: NbnCBP), and the correct construction of the vector was confirmed by sequencing. The plasmids pTRV1, pTRV2, and pTRV2: NbnCBP were transformed into A. tumefaciens strain GV3101 by electroporation. The leaves of 4-week-old N. benthamiana plants were inoculated with a mixture of Agrobacterium cultures containing pTRV1 and pTRV2: NbnCBP (final OD600 = 0.5) to induce NbnCBP silencing; a mixture of Agrobacterium cultures containing pTRV1 and empty pTRV2 was used as a negative control (Control). Two weeks later, the upper leaves were collected to identify the silencing efficiency. Primers for constructing the VIGS vector are shown in Supplementary Table 1.

Virus inoculation

Potato plants at the 4–6 leaf stage were mechanically inoculated with wild-type virus inocula (leaf extract: about 1 g/10 ml leaf tissue homogenized in 10 mM phosphate buffer with 32 mM sodium sulfite, pH 7.5) as described previously (Singh et al., 2003) or agro-infiltrated with virus infectious clones as described previously (Sun et al., 2017). Foliage symptoms were monitored daily post-inoculation until harvest. For resistance assays, the upper non-inoculated leaves were sampled at 10 and 15 days post-inoculation (dpi).

Identification of virus accumulation

Total RNA of partially collected samples was extracted by Total RNAPure Kit (ZOMANBIO, Beijing, China). The first-strand cDNA synthesis was performed by All-In-One 5X RT MasterMix (Applied Biological Materials, Vancouver, Canada). Then, the accumulation level of the viruses was detected through quantitative reverse-transcription polymerase chain reaction (qRT-PCR). The qRT-PCR analyses were carried out in the LightCycler®480 Real-Time PCR system2 using 2X qPCR Real-Time Kit (Applied Biological Materials, Vancouver, Canada). The accumulation level analysis was calculated by the

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1 https://vigs.solgenomics.net/
2 https://diagnostics.roche.com
Supplementary Table 3

Fluorescent complimentary (BiFC) vectors are shown in Wetzlar, Germany. Primers for constructing bimolecular fluorescence signal of YFP was observed at 48 h post-infiltration into recombinant plasmids (final OD600 = 0.2) was infiltrated into Agrobacterium mixture of introduced into Agrobacterium strain GV3101 and agro-infiltrated the leaves of 4-week-old N. benthamiana as protocols previously described (Zhan et al., 2019). The fluorescence signal was observed at 48 h post-infiltration using a confocal laser scanning microscope (SP8, Leica, Wetzlar, Germany). Primers for constructing subcellular localization vectors are shown in Supplementary Table 3.

Yeast-two-hybrid assay

StnCBP, NbnCBP, and AtaCBP were recombined between the BamHI and EcoRI sites of the pGBK7T vector, while the proteins of PVX and PVS were recombined between the BamHI and EcoRI sites of the pGADT7 vector using SE cloning kit (Applied Biological Materials, Vancouver, Canada). Pairwise combinations for interaction analyses were co-transformed into yeast strain AH109 following BD Matchmaker Screening Kit. The interactions were identified through a medium that lacks tryptophan, leucine, adenine, and histidine and contains X-α-GAL (20 mg/L) as the interaction leads to blue plaques. Primers for constructing yeast-two-hybrid (Y2H) vectors are shown in Supplementary Table 1.

Bimolecular fluorescent complimentary

Paired combinations for interaction analysis were inserted into nYFP and cYFP plasmids, respectively, through restriction endonuclease sites BamHI and SalI, and were introduced into Agrobacterium strain GV3101. Then, a mixture of Agrobacterium cultures containing the two recombinant plasmids (final OD600 = 0.2) was infiltrated into N. benthamiana leaves at the 4–6 leaf stage, and the fluorescence signal of YFP was observed at 48 h post-infiltration using a confocal laser scanning microscope (SP8, Leica, Wetzlar, Germany). Primers for constructing bimolecular fluorescent complimentary (BiFC) vectors are shown in Supplementary Table 3.

Subcellular localization

The sequences coding for amino acids in protein (CDS) of StnCBP and PVS CP were amplified with specific primers and recombined into the BsiI407I site of pK7WGGR2 where fused behind eGFP for subcellular localization, and CDS of StnCBP was also inserted into the BsiI407I site of pK7WGR2 where fused behind RFP for co-localization analysis. The vectors were electroporated into Agrobacterium tumefaciens strain GV3101 and agro-infiltrated the leaves of 4-week-old N. benthamiana as protocols previously described (Zhan et al., 2019). The fluorescence signal was observed at 48 h post-infiltration using a confocal laser scanning microscope (SP8, Leica, Wetzlar, Germany). Primers for constructing subcellular localization vectors are shown in Supplementary Table 3.

Western blotting

Total proteins were extracted from leaves of N. benthamiana which expressed the GFP fused StnCBP and CP proteins using a protein extraction buffer [100 mM Tris–HCl (pH 8), 150 mM NaCl, 2 mM dithiothreitol, 5 mM EDTA, 10% glycerol, 2 mM phenylmethylsulfonyl fluoride, and 1% protease inhibitor tablets (A32955; Thermo Fisher Scientific)]. About 20 µl of SDS-PAGE loading buffer with 5 µM β-mercaptoethanol was added to 80 µl of the extracted protein supernatant. The supernatant was heated to 95 °C for 5 min to denature the protein, and then, these samples were loaded for Western blotting analysis. Probes with specific anti-GFP antibodies at 1:5,000 dilution (MBL, Japan) were used for Western blotting as protocols previously described (Cheng et al., 2021; Wang et al., 2021).

Results

StnCBP was an eIF4E gene belonging to the nCBP subgroup in potato

A BLASTn search against the S. tuberosum genome in the Spud DB database was performed using the nucleotide sequence of nCBP in A. thaliana (AtaCBP, Accession number: AT5G18110) as a query to identify nCBP orthologues in potato. As a result, a unique nCBP gene (named as StnCBP, Accession number: Soltu.DM.10G026730) was identified, which exhibited 59.6 and 70.0% homology to AtaCBP in nucleic acid and amino acid sequence, respectively. Potato has four members of the eIF4E family, including SteIF4E (Accession number: Soltu.DM.03G00970), SteIF4E-2 (Accession number: Soltu.DM.02G032530), SteIF(iso)4E (Accession number: 3 http://spuddb.uga.edu/
FIGURE 1

Phylogenetic analysis of eIF4E genes in potato, Arabidopsis, tobacco, and tomato (A), and an amino acid sequence alignment of nCBP proteins from various species (B). The prefix of accession numbers with "Soltu.DM" represents proteins from S. tuberosum; the prefix of accession numbers with "AT" represents proteins from A. thaliana; the prefix of accession numbers with "Solyc" represents proteins from S. lycopersicum; the prefix of accession numbers with "Niben101Scf" represents proteins from N. benthamiana. The eIF4E subgroup, eIF(iso)4E subgroup, and nCBP subgroup are distinguished by pink, blue, and yellow regions, respectively. The phylogenetic tree was generated with neighbor-joining tree and 1000 bootstraps. The red underline represents the eIF4E superfamily domain.

Soltu.DM.09G027260), and StnCBP. Then, we performed a BLASTp search against the NCBI and SGN databases using the amino acid sequences of potato eIF4E family members as queries. As a result, 12 eIF4E proteins were obtained from A. thaliana and two other Solanaceae crops (N. benthamiana and S. lycopersicum). The phylogenetic tree analysis of eIF4E proteins from the above species indicated that StnCBP belongs to the nCBP subgroup (Figure 1A). In addition, an amino acid sequence alignment of nCBP proteins from various species (including the above species, and O. sativa, and Z. mays) revealed that they contain an eIF4E superfamily domain and share a high homology of 82.32%, indicating that nCBP is highly conserved among different species and may have similar functions (Figure 1B).

**StnCBP knockdown compromised potato virus S accumulation**

To determine whether StnCBP is also involved in the infection of important potato viruses encoding TGBps, RNA interference was performed to silence StnCBP expression. Several RNAi transgenic lines were obtained (the interference efficiency is shown in Supplementary Figure 1). The three
transgenic lines with the highest interference efficiency (RiStnCBP-1, RiStnCBP-2, and RiStnCBP-3) were selected for further experiments. As shown in Figure 2A, StnCBP knockdown resulted in deformed compound leaves, indicating that StnCBP plays an important role in leaf development, but does not affect plant growth, which may be due to functional redundancy among eIF4Es. The expression of other members of the SteIF4E family was also examined in the RiStnCBP lines, and the results showed that the expression of SteIF4E, SteIF(iso)4E, and SteIF4E-2, and SteIF(iso)4E was not affected by the transgenic events (Supplementary Figure 2). Then, the RiStnCBP lines were mechanically inoculated with three TGB-encoding viruses, PVS, PVM, and PVX. No apparent symptom was observed after inoculation. The upper non-inoculated leaves were collected from control plants (WT) and transgenic lines at 10 and 15 dpi. Subsequently, the accumulation of these viruses was determined by qRT-PCR and ELISA assays. PVS accumulation was drastically reduced in the transgenic lines compared with that in WT plants (Figure 2B). In contrast, the accumulation of PVM and PVX in the RiStnCBP lines was at a similar level to that in WT plants (Figures 2C,D). These results indicated that StnCBP knockdown suppresses PVS accumulation but not PVM and PVX, which is the first report that a member of the eIF4E family is involved in PVS infection.

StnCBP interacted with potato virus S coat protein

Many susceptibility factors have been proved to function by directly interacting with viral proteins (Roudet-Tavert et al., 2007; Hwang et al., 2015; Zhang et al., 2015). To determine whether StnCBP directly interacts with any protein of PVS, we tested the interaction between StnCBP and six proteins of PVS (including RdRp, TGB1, TGB2, TGB3, CP, and NaBp proteins) in Y2H assays (Figure 3A). The results revealed that StnCBP interacted with the CP of PVS, which was further demonstrated via BiFC assays in N. benthamiana (Figure 3B). Moreover, the results of BiFC demonstrated that StnCBP and PVS CP interacted in the nucleus and cytoplasm. Therefore, we further observed their subcellular localization and colocalization. StnCBP and PVS CP tagged with a GFP at the C-terminus were co-expressed with an RFP-empty vector (RFP-EV) in the nucleus of N. benthamiana to observe their subcellular localization. As a result, the GFP fluorescence of StnCBP and PVS CP was both localized in the nucleus and cytoplasm (Figure 3C). To validate the successful expression of full-length StnCBP and PVS CP with GFP, Western blotting was performed to identify the protein size expressed by the vectors GFP-EV, StnCBP-GFP, and CP-GFP. They were predicted to encode proteins with sizes of 25.6, 52.5, and 59.4 kD, respectively, and the Western blotting results were consistent with expectation (Figure 3D), indicating the correct expression of proteins and accurate subcellular localization. Moreover, StnCBP-RFP and CP-GFP were found to be co-localized in the nucleus and cytoplasm (Figure 3C). These results indicated that StnCBP and PVS CP interact with each other and may function in the nucleus and cytoplasm.

We also tested the interaction between StnCBP and the proteins of PVX or PVM (Supplementary Figure 3). No interaction was observed between StnCBP and any PVX protein. Surprisingly, StnCBP also interacted with PVM CP, but StnCBP knockdown did not affect PVX infection. Therefore, we further determined whether PVX CP is critical for StnCBP-mediated recessive resistance to PVS.

Potato virus S coat protein played a vital role in StnCBP-mediated recessive resistance to potato virus S

Construction of plant RNA virus infectious clones has become a powerful tool to study virus molecular biology (Bao et al., 2020). Therefore, we constructed a PVS infectious clone to test whether PVS CP plays an important role in StnCBP-mediated recessive resistance to PVS. According to the previously described method (Sun et al., 2017), the PVS genome was divided into three partially overlapping cDNA fragments (PVS-A, PVS-B, and PVS-C), followed by amplification using specific primers (Supplementary Table 4) and homologous recombination assembly in the yeast with an expression vector pCB301-2µ-HDV to construct the full-length clone of PVS (named as pCB301-2µ: PVS; Figure 4A). Then, the CP in PVS infectious clone was replaced by the CP of PVM or PVX, and the infectious clones were designated as PVS\(^{\text{CP/S}}\), PVS\(^{\text{CP/M}}\), and PVX\(^{\text{CP/X}}\), respectively, (Supplementary Figure 4).

To confirm the infectivity, we inoculated the WT plants with the three infectious clones by Agrobacterium infiltration and detected the presence of the virus in the upper non-inoculated leaves by PCR assays at 14 dpi, with two pairs of primers. The first pair of primers (named as detection primers) were designed on the TGB1 region of PVS for general detection. The other pair of primers (named as CP-flanking primers) were designed based on the sequences flanking the CP region, to confirm the substitution of viral CP according to the different sizes of CPs from PVS, PVM, and PVX (Supplementary Figure 5). As a result, PVS\(^{\text{CP/S}}\) and PVX\(^{\text{CP/M}}\) infectious clones successfully infected the plants as indicated by the amplification of PCR products with expected sizes (Figure 4B). In addition, the nucleotide lengths of CP genes of PVS and PVM were 885 bp and 915 bp, respectively. As expected, the PCR product amplified by CP-flanking primers from the PVS\(^{\text{CP/M}}\) inoculated plants was slightly larger than that from PVS\(^{\text{CP/S}}\) inoculated plants (Figure 4B), indicating correct substitution of the CP. Surprisingly, PVS\(^{\text{CP/S}}\) showed no infection ability since no PCR product was amplified.
FIGURE 2
Morphology and virus resistance identification of the RiStnCBP transgenic lines. (A) Morphology of RiStnCBP transgenic lines under field conditions. (B–D) Virus resistance identification in RiStnCBP transgenic lines against PVS (B), PVM (C), and PVX (D), respectively. The virus accumulation was determined using qRT-PCR (left) and ELISA (right). Error bars indicate ± SD of three replicates (n = 3). Three independent experiments were performed with similar results. p < 0.05*, p < 0.01**, and p < 0.001*** (Student's t-test).
FIGURE 3

StnCBP interacted with PVS CP in the nucleus and cytoplasm. (A) Interaction between StnCBP and proteins of PVS in Y2H assays. -W-L represents a medium lacking tryptophan and leucine, while -W-L-H-A represents a medium lacking tryptophan, leucine, histidine, and adenine. Paired combinations PGADT7-T/PGBK7-53 and PGADT7-T/PGBK7-lam represent positive and negative controls, respectively. (B) Interaction between StnCBP and PVS CP in BiFC assays. Merge means the overlay of YFP and visible light on single confocal planes. Scale bar: 50 µm. (C) Subcellular localization and co-localization of StnCBP and PVS CP. Merge means the overlay of GFP and RFP on single confocal planes. Scale bar: 20 µm. (D) Identification of full-length proteins of StnCBP and PVS CP successfully expressed with GFP in the Western blotting. Total proteins were immunoprecipitated with anti-GFP antibody, and Rubisco was stained with Ponceau S as a loading control.

(Figure 4B). A plausible explanation is that both PVS and PVM belong to the Carlavirus genus with mutually compatible CPs, while PVX belongs to the Potexvirus genus with an incompatible CP.

Furthermore, the RiStnCBP lines were inoculated with the PVS<sup>CP/S</sup> and PVS<sup>CP/M</sup> infectious clones by Agrobacterium infiltration. The upper non-inoculated leaves were collected at 10 and 15 dpi to determine the virus accumulation using qRT-PCR and ELISA assays. The RiStnCBP lines also had significantly lower accumulation of PVS<sup>CP/S</sup> infectious clone than the control plants (Figure 4C). In contrast, the accumulation of PVS<sup>CP/M</sup> infectious clone showed no significant difference between the RiStnCBP and control plants (Figure 4D). These results indicated that PVS CP is vital for StnCBP-mediated recessive resistance to PVS. However, the interaction between StnCBP and CP may be necessary but not sufficient for the StnCBP-mediated recessive resistance to TGB-encoding viruses.
FIGURE 4

PVS CP played a vital role in the StnCBP-mediated recessive resistance to PVS. (A) Schematic illustration of the construction of PVS infectious clone. PVS-A, PVS-B, and PVS-C represent the fragments from PVS for the assembly. The gray shadow bars indicate the overlapping regions of adjacent fragments, with the number of overlapping nucleotides. (B) Electrophoretic separation of PCR products amplified with detection primers (above) and CP-flanking primers (below) from plants agroinoculated with PVS\textsuperscript{CP/S} (1–3), PVS\textsuperscript{CP/M} (4–6), and PVS\textsuperscript{CP/X} (7–9). M: Trans2K\textsuperscript{R} Plus DNA Marker. The upper non-inoculated leaves were collected 14 days after inoculation. (C,D) The accumulation of PVS\textsuperscript{CP/S} (C) and PVS\textsuperscript{CP/M} (D) infectious clones in the RiStnCBP and control plants (WT). The virus accumulation was determined using qRT-PCR (left) and ELISA (right). Error bars indicate ± SD of three replicates (n = 3). Three independent experiments were performed with similar results. *p < 0.01** and *p < 0.001*** (Student’s t-test).
Antiviral identification of NbnCBP against PVX, and interaction analyses between NbnCBP and proteins of PVX. (A) The silencing efficiency of NbnCBP in N. benthamiana by VIGS. The gene expression was determined using qRT-PCR. Error bars indicate ± SD of five replicates (n = 5). \( p < 0.001^{***} \) (Student’s t-test). (B,C) The determination of PVM and PVX accumulation in the control and NbnCBP-silenced plants by RT-qPCR (B) and ELISA (C). Error bars indicate ± SD of three replicates (n = 3). Three independent experiments were performed with similar results. \( p < 0.01^{**} \) (Student’s t-test). (D) Green fluorescence emission in PVX-GFP inoculated leaves of the control, NbnCBP-silenced, and NbnCBP-silenced: complemented plants. The photographs were taken at 8 dpi. (E,F) The determination of PVX-GFP accumulation in inoculated leaves of the control, NbnCBP-silenced, and NbnCBP-silenced: complemented plants by RT-qPCR (E) and ELISA (F). Error bars indicate ± SD of three replicates (n = 3). Three independent experiments were performed with similar results. \( p < 0.01^{**} \) (Student’s t-test). (G) Interaction analysis between NbnCBP and the proteins of PVX in Y2H. -W-L represents a medium lacking tryptophan and leucine, while -W-L-H-A represents a medium lacking tryptophan, leucine, histidine, and adenine. Paired combinations PGADT7-T/PGBKT7-53 and PGADT7-T/PGBKT7-lam represent positive and negative controls, respectively. (H) Interaction analysis between NbnCBP and PVX CP in BiFC assay. nYFP and cYFP represent their empty vectors. Merge means the overlay of YFP and visible light on single confocal planes. Scale bar: 50 µm.
The nCBP orthologue in *N. benthamiana* was involved in potato virus X infection

The function of *nCBP* gene in the infection of TGB-encoding viruses was first reported in Arabidopsis and then in potato in this study. However, it remains unknown whether the function of *nCBP* homolog is also conserved in *N. benthamiana*, which also belongs to the *Solanaceae* as potato. Therefore, the TRV-mediated VIGS assay was employed to silence the *nCBP* homolog in *N. benthamiana* (*NbnCBP*; accession number: Niben101Sc011669g04023.1). The silencing efficiency of *NbnCBP* by VIGS reached 80% (Figure 5A). The *NbnCBP*-silenced plants were then mechanically inoculated with PVS, PVM, and PVX. At 6 dpi, the upper non-inoculated leaves were collected to determine the virus accumulation by qRT-PCR and ELISA assays. Surprisingly, the accumulation of PVX was significantly reduced, while that of PVM was not affected in the *NbnCBP*-silenced plants, which was inconsistent with the results obtained for potato (Figures 5B,C). In addition, *N. benthamiana* exhibited resistance to PVS as no virus was detected in both *NbnCBP*-silenced and control plants (Figures 5B,C). It was reported that many tobacco varieties could not be infected by PVS in mechanical inoculation (De Bokx, 1970; Brattey et al., 2002; Nie and Singh, 2013), indicating that *N. benthamiana* may have some unknown resistance against PVS like these tobaccos. Therefore, it remains unclear whether *NbnCBP* is involved in the infection of PVS.

To provide more intuitive results, the *NbnCBP*-silenced plants were inoculated with a PVX infectious clone tagged with a green fluorescent protein (hereafter referred to as PVX-GFP) by agroinfiltration. The GFP fluorescence at 8 dpi revealed that PVX-GFP accumulation drastically decreased in the inoculated leaves, and the systemic infection was significantly delayed (Figure 5D). Further qRT-PCR and ELISA assays demonstrated that both the PVX RNA and protein levels decreased in *NbnCBP*-silenced plants compared with those in the control (Figures 5E,F). Obviously, these results were consistent with those of GFP fluorescence emission, indicating that *NbnCBP* knockdown compromises PVX-GFP accumulation.

A transient complementation assay was performed to confirm that the increase in the accumulation of PVX in *NbnCBP*-silenced plants was directly caused by loss of *NbnCBP*. The leaves of *NbnCBP*-silenced plants were agro-infiltrated with a mixture of *Agrobacterium* cultures carrying the PVX-GFP infectious clone and a full-length cDNA of *NbnCBP*. As a result, the GFP fluorescence of PVX-GFP was recovered in the inoculated leaves of *NbnCBP*-silenced: complemented plants. However, the systemic infection of PVX-GFP was still delayed (Figure 5D). In addition, the qRT-PCR and ELISA results also revealed the PVX-GFP accumulation in the inoculated leaves of *NbnCBP*-silenced: complemented plants was comparable to the control plants (Figures 5E,F). Thus, it could be speculated that *NbnCBP* is required for PVX infection in *N. benthamiana*.

Then, a Y2H assay showed that the CP was the only protein interacting with *NbnCBP* in the five proteins coded by PVX, which was similar to the relationship between StnCBP and PVS CP (Figure 5G), and the interaction between *NbnCBP* and CP of PVX was further confirmed by the BiFC assay (Figure 5H). Thus, PVX CP may also have a critical function in *NbnCBP*-mediated recessive resistance against PVX. In addition, we analyzed the interaction between *NbnCBP* and the proteins of PVM and PVS. Interestingly, like the case in potato, *NbnCBP* interacted with PVM CP, though *NbnCBP* knockdown did not affect PVM infection (Supplementary Figures 6A,B). *NbnCBP* also interacted with PVS CP (Supplementary Figures 6C,D), even if PVS did not infect *N. benthamiana*. This result implies that *NbnCBP* may also have the function to facilitate PVS accumulation, but this phenotype was masked by other possible anti-PVS mechanisms prevalent in tobacco.

In summary, our results indicated that the *nCBP* orthologues have conserved functions in potato and tobacco as in Arabidopsis in terms of recessive resistance against TGB-encoding viruses, and the interaction between *nCBP* and CP is necessary but not sufficient to determine the recessive resistance of *nCBP*.

**Discussion**

Recessive resistance genes have become a very important resource in plant breeding for virus resistance (Truniger and Aranda, 2009). *eIF4Es* are the most extensively studied recessive resistance genes involved in the infection of various viruses (Hashimoto et al., 2016; Xu et al., 2017; Yang et al., 2021). This study for the first time reports that *StnCBP*, a member of the *eIF4E* family in potato, is involved in PVS infection as a recessive resistance gene. PVS has become an important virus in all potato-growing areas worldwide (Song et al., 2017). However, compared with potato virus Y resistance, potato resistance to PVS has been much less studied, except for the chromosomal localization of a dominant gene (Ns) against PVS (Marczewski et al., 2002). This study identified a recessive resistance gene (*StnCBP*) to PVS, which may provide a new strategy for developing potato varieties resistant to PVS.

Numerous studies of *eIF4Es* have suggested that their interactions with VPg proteins are responsible for the susceptibility to potyviruses. For instance, the mutation of *eIF4Es* that abolishes their interaction with VPg proteins confers resistance to potyviruses, and, in turn, some isolates of potyviruses can restore the interaction to lead to a resistance-breaking phenotype (Yeam et al., 2007; Charron et al., 2008; Truniger and Aranda, 2009). In this study, *StnCBP* knockdown compromised PVS accumulation but not PVX and PVM (Figure 2), and further assays demonstrated that StnCBP...
directly interacts with PVS CP both in vitro and in vivo (Figure 3). These results indicated that PVS CP may also be crucial for the function of StnCBP just like VPgs for eIF4E/eIF(iso)4E. We further confirmed the speculation since the substitution of PVS CP by PVM CP recovered the virus accumulation in the RiStnCBP lines (Figure 4). Furthermore, our results also suggested that NbnCBP is required for PVX infection when it interacts with PVX CP (Figure 5), indicating that nCBP-CP interaction may be a common mode in nCBP-mediated recessive resistance against TGB-encoding viruses.

However, how nCBP facilitates TGB-encoding virus accumulation is still elusive. nCBP was found to be a component of the 5'cap complex in Arabidopsis and supports the translation initiation of capped mRNA, like eIF4E/eIF(iso)4E (Ruud et al., 1998; Bush et al., 2009). The genomes of PVS, PVM, and PVX consist of five or six open reading frames, a 5' cap and a 3' poly(A) tail (Poke, 2008; Petrova et al., 2013). Studies on the PVX suggested that the 5’ cap structure of RNA and PVX CP plays crucial role for viral ribonucleoprotein (vRNP) assembly (Petrova et al., 2013, 2015). Therefore, the interaction between nCBP and viral CP may promote vRNP assembly, thereby regulating viral translation and facilitating the accumulation of TGB-encoding viruses.

Notably, the knockout of StnCBP and NbnCBP did not affect PVM infection in S. tuberosum and N. benthamiana, respectively, whereas both StnCBP and NbnCBP interacted with PVM CP. One reasonable explanation is that knockdown instead of knockout was used to identify the function of nCBP homolog genes, which still produced a certain amount of transcripts. Thus, it is possible that the effect on PVM would only be observed in a knockout situation. In fact, the ncbp mutation in Arabidopsis significantly reduced the accumulation of PVM (Keima et al., 2017), and AtncBP was found to interact with PVM CP in both the Y2H and BiFC assays (Supplementary Figures 7A,B), which supports this explanation. Further knockout analysis is needed to confirm the recessive resistance of the nCBP orthologues in potato and tobacco to these TGB-encoding viruses.

Another possible explanation is that the nCBP-CP interaction may be a necessary but not sufficient condition for nCBP to act as a susceptibility gene, and some other viral proteins may also be critical in the process. The TGBps are considered as conserved gene modules involved in the viral movement, which is similar to movement proteins (MPs) of some viruses (Morozov and Solovyev, 2003; Keima et al., 2017). TGB-encoding viruses are classified into two classes: the hordei-like class in which the CP is dispensable for cell-to-cell movement of viruses, and the potex-like class in which the CP is required for cell-to-cell movement (Forster et al., 1992; Solovyev et al., 1996; Wong et al., 1998; Lin et al., 2006). The potex-like class viruses have a TGB1 ranging from 24 to 26 kDa and a TGB3 of about 15 kDa (Morozov and Solovyev, 2003; Lauber et al., 2005). In addition, the hordei-like class viruses are multicomponent viruses with rigid rod-shaped virions, while the potex-like class viruses are monopartite viruses with filamentous virions (Lauber et al., 2005). Thus, the TGBps of PVS, PVM, and PVX can be classified into the potex-like class (Morozov and Solovyev, 2003), implying that their CPs may be required for TGBs to establish the cell-to-cell movement and systemic infection of viruses. In addition, a previous study has shown that ncbp mutation in Arabidopsis conferred resistance to serval potex-like viruses (Keima et al., 2017). Among these viruses, the cell-to-cell movement of plantago asiatica mosaic virus (Potexvirus) was significantly delayed, and the accumulation of TGB2 and TGB3 was reduced, which established the association between the function of AtncBP and TGBp. In addition, we analyzed the interaction between AtncBP and the proteins of PVX by Y2H assays. As a result, AtncBP interacted with the TGB2, TGB3, and CP of PVX (Supplementary Figure 7C), indicating the association among nCBP, TGBps, and CP.

In summary, this study identified StnCBP as a susceptibility gene required for the accumulation of a TGB-encoding virus PVS by recognizing its CP in potato. Our findings revealed that the nCBP-CP interaction may also contribute to the function of nCBP orthologues for other TGB-encoding viruses. Moreover, the cassava nCBP was found to interact with the VPgs of two ipomoviruses, which do not encode TGBps (Gomez et al., 2019). It will be meaningful to further study whether nCBP is required for the infection of other types of viruses and the underlying mechanism for nCBP to facilitate the infection of TGB-encoding and non-TGB-encoding viruses.

Data availability statement

The original contributions presented in this study are included in the article/Supplementary material, further inquiries can be directed to the corresponding authors.

Author contributions

CH, BN, RC, and XH conceived and designed the experiments. RC, MY, FX, and TL performed the experiments. RC, ZT, and JC analyzed the data. RC, CH, and BN prepared the manuscript. CH, BN, and XH revised the manuscript. All authors contributed to the article and approved the submitted version.

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Conflict of interest

The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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Supplementary material

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fpls.2022.946873/full#supplementary-material
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