Heterologous expression of the cysteine proteinase gene VaCP17 from the cold-adapted grapevine *Vitis amurensis* increases cold tolerance in Arabidopsis

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Abstract

Cysteine proteinases (thiol) carry out diverse and critical functions in plants through their ability to hydrolyze peptide bonds in target proteins. Here, we cloned a cysteine proteinase gene designated *VaCP17* from a highly cold-resistant wild *Vitis amurensis* accession ‘Shuangyou’, and then its potential function in cold resistance was investigated. The results showed that the CDS of *VaCP17* is 1404 bp, encoding 467 amino acids, the VaCP17 protein localized to the cell membrane. Expression of *CP17* was highly distinctive among different structures of ‘Shuangyou’ and the cold-sensitive *Vitis vinifera* cultivar ‘Red Globe’, with the highest expression in the stem of ‘Shuangyou’ and the flower of ‘Red Globe’. Arabidopsis plants constitutively expressing a *VaCP17*-GREEN FLUORESCENT PROTEIN fusion (35S::*VaCP17*-GFP) showed increased survival after transient exposure to freezing (-6 °C), and showed lower electrolyte leakage and MDA content, higher soluble sugar content and SOD, POD and CAT activities, as compared with non-transgenic Arabidopsis controls. The expression of nine cold-resistance related genes (*CBF1, CBF2, CBF3, RD29A, COR15A, KIN1, NCED3, AOC1* and *JAZ10*) in 35S::*VaCP17*-GFP plants was increased under cold treatment at 4 °C, relative to control plants. Using a yeast two-hybrid system, we identified *VaNAC72, VaCAM7* and *VaDi19* as potential interactors of *VaCP17*, and their interactions were demonstrated by a bimolecular fluorescence complementation assay. In conclusion, we revealed that *VaCP17* can enhance cold resistance by influencing physiology and biochemistry and the expression of cold resistance related genes under cold stress.

Key message

*VaCP17* gene was cloned from the high cold-resistant wild *Vitis amurensis* acc. ‘Shuangyou’. The heterologous expression of *VaCP17* can enhance the cold tolerance of *Arabidopsis thaliana*.

Keywords *Vitis amurensis* · *VaCP17* · Transgenic Arabidopsis · Cold resistance · Interacting proteins
Introduction

Cold is one of the most important abiotic stresses limiting plant geographical distribution and crop production (Ye et al. 2017). The damage caused by low temperature to plants can be classified as cold injury (low temperatures above 0 °C) or freezing injury (temperatures below 0 °C). Cold stress can destabilize the plant cell membrane, increase the accumulation of reactive oxygen species and disrupt protein structure, all of which negatively impact photosynthesis, metabolism, and growth and development (Chinnusamy et al. 2007; Liu et al. 2018; Yin et al. 2021).

Plants have evolved complex regulatory networks that mediate responses to cold stress to increase survival. These include the rapid deployment of the CBF/DREB (C-repeat binding/dehydration element binding) family of transcription factors, which recognize the C-repeat and DRE cis elements in downstream COLD RESPONSIVE (COR) genes to modulate their expression (Shi et al. 2012). The expression of COR genes, such as COR15A, RD29A, KIN1 and GOLS3, can improve plant tolerance to low temperature (Knight et al. 2004; Thomashow 2010; Park et al. 2015; Guo et al. 2019). In addition, cold stress can induce the expression of abscisic acid (ABA) and jasmonic acid (JA) biosynthetic genes, resulting in the accumulation of ABA and JA, which also regulate genes required for cold response and survival (Knight et al. 2004; Hu et al. 2013; Feng et al. 2016). Allene oxide cyclase (AOC) is one of the key enzymes in jasmonic acid (JA) biosynthesis, which can affect the stereoisomerization and biological activity of jasmonic acid molecules, and plays an important role in plant stress resistance (Sun et al. 2020). Jasmonic acid induces the degradation of JAZ protein and activates the expression of cold resistance-related genes, thus improving plant cold resistance (An et al. 2021). Abiotic stresses such as cold, salt and drought usually increase the content of ABA, which in turn induces stomatal closure and reduces transpiration rate; ABA can also regulate the expression of stress-related genes (Mulholland et al. 2003; Jiang et al. 2014; Li et al. 2019). The rate-limiting enzyme of ABA biosynthesis is 9-cis-epoxycarotenoid dioxygenases (NCEDs). Together, these mechanisms act to enhance the stability of cellular membranes by promoting accumulation of osmolytes such as proline, soluble sugar and soluble protein (Ashraf and Foolad 2007).

Cysteine proteinases, also known as thiol or sulfhydryl proteinases, play various roles in plant growth and development, including senescence and programmed cell death (PCD) (Xu et al. 2003; Paireder et al. 2016; Burke et al. 2020). Senescence can be conditioned by several factors, including water deficit, phytohormones, and pathogen infection (Yang et al. 2003; Munné-Bosch and Alegre 2004; Mercedes et al. 2016). Likewise, PCD can be triggered by a variety of biotic and abiotic stresses (Paireder et al. 2016; Burke et al. 2020). Both abiotic and biotic stress can lead to damage of intracellular proteins, and proteolysis of damaged proteins by cysteine proteinases can minimize the potentially negative effects of aberrant protein activity (García-Lorenzo et al. 2006; López-Barón et al. 2017). Abiotic stresses will aggravate the production of reactive oxygen species (ROS), and the excessive accumulation of ROS will lead to oxidative stress, leading to various metabolic disorders and affecting plant growth and development (Bose et al. 2014). Cysteine proteinases are up-regulated under oxidative stress and play an important role in maintaining cell metabolism (Usui et al. 2007). Cysteine proteinases can also regulate the level of reactive oxygen species in cells, thus reducing oxidative damage to membranes and proteins (Hoorn and Renier 2008).

Several studies have revealed that cysteine proteinase genes can be induced by abiotic stress (Jones and Mullet. 1995; Zang et al. 2010; Zheng et al. 2018). For example, in Arabidopsis thaliana (Arabidopsis), Pisum sativum, Nicotiana tabacum and Ipomoea batatas, cysteine proteinase genes are upregulated in response to drought, salt, cold or exogenous hormones such as gibberellin, jasmonic acid and abscisic acid (Koizumi et al. 1993; Jones and Mullet 1995; Beyene 2006; Chen et al. 2010). Zhang et al. exposed wheat seedlings to low temperature, high temperature, salt and drought stresses, and found that wheat TaCP3 was induced by all these conditions, especially drought (Zhang et al. 2019).

Ectopic expression of the cysteine protease SpCP3 from sweet potato in transgenic Arabidopsis enhanced the sensitivity of the plants to drought stress (Chen et al. 2013). Heterologous expression of Salix matsudana SmpCP in Arabidopsis improved salt tolerance, and this was associated with increased stability of cell membrane and activity of the antioxidant enzyme system (Zheng et al. 2018). Therefore, plant cysteine proteases play a key role in the defense to abiotic stress.

Vitis amurensis (Amur grape) is native to present-day northeastern China and far-eastern Russia, and is adapted to extremely low winter temperatures. Currently there is wide interest in exploiting genes/alleles conferring cold tolerance in V. amurensis for cultivar improvement of domestic grapes, including the commerically valuable Vitis vinifera. In this study, we identified a cysteine proteinase gene from V. amurensis that is induced in response to cold. We generated transgenic Arabidopsis expressing VaCP17, and analyzed phenotypic, physiological and biochemical indices related to cold stress response, and evaluated expression of cold-resistance related genes. We further identified proteins that could interact with VaCP17 by yeast two-hybrid technology. Together, the results of
these experiments lay the foundation for further studies of the molecular mechanisms of VaCP17 in cold resistance of V. amurensis.

Materials and methods

Plant materials and treatments

Vitis amurensis acc. ‘Shuangyou’ and Vitis vinifera cv. ‘Red Globe’ were maintained in the grapevine germplasm repository at Northwest A & F University, Yangling, Shaanxi, China. Seeds of A. thaliana and Nicotiana benthamiana were obtained from the State Key Laboratory of Crop Stress Biology in Arid Areas, Northwest A & F University.

To assess freezing tolerance of Arabidopsis, plants were maintained in a controlled environment chamber at 10 °C for 24 h, and then the temperature was changed to -6 °C. To assess physiological and biochemical indices, another set of plants were acclimated at 10 °C for 24 h and then subjected to 4 °C for 48 h. Samples were taken respectively at 0, 3, 6, 12, 24 and 48 h. Three biological replicates were included at each time point. Samples were immediately stored at −80 °C.

Cloning and sequence analysis of VaCP17 gene

The VaCP17 coding sequence (CDS) was cloned by PCR from cDNA derived from leaves of V. amurensis acc. ‘Shuangyou’, using oligonucleotide primers VaCP17-F and VaCP17-R as designed by Primer Premier 5.0 software (Supplementary Table 1). The PCR reaction contained PrimeSTAR® (Takara) Buffer, 200 μM dNTPs, 0.4 μM primer VaCP17-F, 0.4 μM primer VaCP17-R, 2 μL cDNA, and 1.25 units PrimeSTAR®HS, in a total volume of 50 μL. The reaction profile was 32 cycles of 98 °C for 10 s; 57 °C for 15 s, and 72 °C for 1 min 30 s, with a final incubation at 72 °C for 10 min. Amplified DNA was purified and cloned using the pMD19-T vector to generate pMD19-T-VaCP17. The VaCP17 CDS in pMD19-T-VaCP17 was subjected to Sanger sequencing on both strands to ensure that no mutations had resulted from the PCR amplification.

A draft genome annotation of V. amurensis was used to determine the chromosomal location of VaCP17 (Wang et al. 2021). Homologous sequences were identified from the NCBI nr (non-redundant protein) database using BLASTP and the predicted VaCP17 amino acid sequence as a query. DNAMAN was employed to construct a multiple amino-acid alignment and MEGA7.0 was employed to construct a phylogenetic tree by the neighbor-joining method (Tamura et al. 2004).

VaCP17 subcellular localization assay

The cloned VaCP17 CDS was inserted into the pCAMBIA2300 vector downstream of the 35S promoter to generate the 35S::VaCP17-GFP expression vector. 35S::VaCP17-GFP and pCAMBIA2300 plasmid DNAs were introduced into Arabidopsis protoplasts by a polyethylene glycol (PEG)-mediated method (Yu et al., 2013). The protoplasts were cultured at 22 °C under weak light for about 18 h. The localization of VaCP17-GFP fusion protein was observed by confocal microscopy (LEICA TCS SP8, Leica, Germany).

Analysis of CP17 expression in structures of ‘Shuangyou’ and ‘Red Globe’

Young roots, stems, leaves, flowers and tendrils of ‘Shuangyou’ and ‘Red Globe’ were collected from plants that had grown in the field natural conditions for more than 10 years. RNA was extracted and first-strand cDNA generated using the E.Z.N.A. Plant RNA Kit from Omega Bio-tek and FastQuant RT with gDNAase from TIAN-GEN, respectively. qRT-PCR analysis was performed using SYBR®Premix Ex TaqTMII (ThermoFisher) and the QS6 real-time quantitative PCR detection system (Bio-Rad). RT-CP17-F and RT-CP17-R were used as primers (Supplementary Table 1), and VvActin (XM_002265440) was employed as the internal reference gene. The 2−ΔΔCT method was used to calculate gene relative expression levels.

Production and identification of transgenic Arabidopsis expressing VaCP17

Production of VaCP17 transgenic Arabidopsis

Agrobacterium tumefaciens strain GV3101 harboring 35S::VaCP17-GFP was used to transform Arabidopsis plants by the floral dip method. T1 seedlings resistant to Kan were identified by germination and growth on ½x MS medium containing 50 mg/L Kan. Homozygous lines were identified by progeny testing, and initiated from T2 plants.

Identification of VaCP17 transgenic Arabidopsis

Genomic DNA was extracted from rosette leaves of T3 homozygous plants, and the presence of pCAMBIA2300 sequence was determined using PCR and primers specific for pCAMBIA2300 (Supplementary Table 1). PCR-positive plants were evaluated for VaCP17 expression using real-time fluorescence quantitative RT-PCR and gene-specific primers as shown in Supplementary Table 1.

To detect VaCP17 protein, total proteins were extracted from Arabidopsis leaves using the method described by Wang et al. (2006), separated on a 10% SDS-PAGE gel and
transferred to PVDF film using a semi-dry transfer membrane method. Anti-GFP-tagged mouse monoclonal antibody (TransGen Biotech) was used as the primary antibody and a HRP-labeled goat anti-mouse IgG (H + L) (TransGen Biotech) was used as the secondary antibody.

**Evaluation of tolerance of 35S:VaCP17 transgenic Arabidopsis to cold stress**

**Phenotypic changes of transgenic plants under cold stress**

Seeds of 35S::VaCP17-GFP lines or non-transgenic control plants were sown in an artificial potting medium, and seedlings/plants were cultivated for four weeks at 22 °C under 16 h-light/8 h-dark photoperiods. Subsequently plants were transferred to 10 °C for 24 h, and then −6 °C for 8 h. After the freezing treatment, the temperature was raised at 3 °C/h to 22 °C, and then plants were returned to previous conditions for 10 d prior to phenotypic observations.

**Physiological and biochemical indices of 35S::VaCP17-GFP plants under cold stress**

Leaves of 35S::VaCP17-GFP and non-transgenic control plants lines were removed after cold treatment and used to evaluate electrolyte leakage, contents of MDA, proline and soluble sugar, and activities of SOD, POD, and CAT, as described by Gao (2006).

**Analysis of expression of VaCP17 and cold-resistance-related genes in 35S::VaCP17-GFP and control plants under cold stress**

Four-week-old 35S::VaCP17-GFP and non-transgenic control plants were placed at 10 °C for 24 h, then at 4 °C for 48 h, and samples were taken at 0, 3, 6, 12, 24 and 48 h, with three biological replicates at each time point. RNA was extracted from the samples and subjected to reverse transcription into cDNA. AtACTIN (AT3G18780) was used as an internal reference gene. Primers used for quantitative analysis of expression of VaCP17 and cold resistance-related genes are shown in Supplementary Table 1.

**Screening and verification of VaCP17-interacting proteins**

**Verification of VaCP17 self-activation**

The VaCP17 CDS was cloned by PCR using primers described in Supplementary table 1 and inserted into pGBKKT7 (Supplementary Table 1). The pGBKKT7/VaCP17 plasmid was extracted and transferred into yeast Y2HGold cells by the LiAc method, and tranformed cultures were spread onto SD/-Trp agar medium. After incubating at 30 °C for 3 d, P53/pGBKKT7 and Lam/pGBKKT7 were co-transformed with pGADT7-T, and cultures were spread on SD/-Trp-Leu medium as positive and negative controls, respectively. Three isolates each for pGBKKT7/VaCP17, negative and positive controls were selected and suspended in 150 μl sterile water, then spotted onto SD/-Trp medium containing 200 ng/ml AbA and 20 mg/ml X-α-gal.

**Identification of VaCP17-interacting proteins by yeast two-hybrid screening**

A cDNA library was constructed RNA derived from leaves of a V. amurensis acc. ‘Shuangyou’ individual that had been subjected to 0 °C, and the cDNA library plasmid was constructed by Takara Biotech Inc. The library was constructed in the plasmid pGADT7, and the VaCP17 CDS was cloned into the bait vector plasmid pGBKKT7/VaCP17. Prey and bait vectors were co-transformed into Y2HGold yeast cells, spread on SD/-Trp/-Leu/-Ade/-His solid medium, and positive clones were characterized by DNA sequencing. In order to confirm initially observed protein interactions, selected candidate genes were cloned independently into pGADT7 and co-transformed with the pGBKKT7/VaCAP17 bait vector into Y2HGold cells, and then spread onto SD/-Trp/-Leu solid medium. After 3 d of growth, clones were selected and verified on SD/-Trp/-Leu/-Ade/-His/AbA/X-α-gal solid medium.

**Bimolecular fluorescence complementation assay for VaCP17-interacting proteins**

The CDS sequence of VaCP17 was incorporated into the PBI221NE vector by homologous recombination to generate PBI221NE/VaCP17, and the CDS sequence of interacting proteins were incorporated into PBI221CE to generate PBI221CE/X, using oligonucleotide primers as listed in Supplementary Table 1. The PBI221NE/VaCP17 and PBI221CE/X plasmids were extracted and co-transformed into Arabidopsis protoplasts by a polyethylene glycol (PEG)-mediated method (Yu et al. 2013). The protoplast suspension was then cultured at 22 °C under weak light for about 18 h. YFP fluorescence was observed with a confocal microscope (LEICATCS SP8, Leica, Germany), using an excitation wavelength of 514 nm and emission wavelength of 527 nm.

**Statistical analysis**

Data are presented here as the mean ± standard deviation (SD) of three biological replicates. All data were analyzed by LSD and Duncan methods of SPSS one-way ANOVA (*P < 0.05; **P < 0.01; ***P < 0.001).
Results and analysis

Cloning of VaCP17 and phylogenetic analyses

As a first step to assess a potential role for VaCP17 in cold tolerance, we cloned a 1404-bp, full-length VaCP17 coding sequence (CDS) using RT-PCR and RNA from ‘Shuangyou’ leaf as template (Fig. 1a).

According to the genome of V. amurensis, the VaCP17 gene is located on Chromosome 18 (Fig. 1b). The presumed VaCP17 protein contains three conserved domains: Inhibitor_I29, Peptidase_C1A and GRAN (Fig. 1b). Homologous sequences to VaCP17 in A. thaliana, Carica papaya, Citrus sinensis, Durio zibethinus, Hevea brasiliensis, Juglans regia, Malus domestica, N. tabacum, Prunus avium, Populus trichocarpa, Pistacia vera, Ziziphus jujuba were identified in the NCBI nr database, and the amino acid sequences of VaCP17 and CPs with high homology were compared. It was found that the amino acid sequence similarity between VaCP17 and CsCP was the highest, at 80.6% (Fig. 1c).

Results of a phylogenetic analysis including the most closely homologous sequences showed that VaCP17 was most closely related to CPs from Nicotiana tabacum and Carica papaya (Fig. 1d).

VaCP17 localises to the cell membrane

To determine the subcellular localization of VaCP17, the 35S::VaCP17-GFP construction in pCAMBIA2300, or the non-modified pCAMBIA2300 plasmid, were transformed into Arabidopsis protoplasts. The results showed that green fluorescence was observed on cell membrane, while in the control green fluorescence was visible in the whole cell. This suggests that VaCP17 localizes to the cell membrane (Fig. 2).

Developmental expression of CP17 in ‘Shuangyou’ and ‘Red Globe’

We evaluated the developmental expression of CP17 in five structures (root, stem, leaf, flower, and tendril) of ‘Shuangyou’ and ‘Red Globe’ plants growing under field conditions. CP17 expression was detectable in all of the structures in both genotypes. In ‘Shuangyou’, expression was highest in the stem, while in ‘Red Globe’ expression was relatively low in the stem and highest in the flower (Fig. 3). This finding suggests that the CP17 gene is regulated distinctly between the two genotypes during development.

Fig. 1 Cloning and sequence analysis of VaCP17 isolated from Vitis amurensis accession ‘Shuangyou’. a Cloning of VaCP17 from V. amurensis acc. ‘Shuangyou’. M. DNA marker III; 1–2. Independent VaCP17 amplification products. b Chromosomal location of VaCP17 and conserved domains within the protein. c Multiple sequence alignment of VaCP17 with strongly homologous CPs from other plant species. d Phylogenetic analysis of VaCP17 and other plant CPs. The accession numbers are as follows: AtCP, Arabidop-
Production and molecular identification of transgenic plants

To assess the potential effects of heterologous expression of VaCP17 in Arabidopsis, we engineered a VaCP17 fusion with the GFP sequence under control of the strong, constitutive 35S promoter and introduced this into Arabidopsis via Agrobacterium-mediated transformation. Ten independent T1 lines were initially analyzed by qRT-PCR, using oligonucleotide primers specific for the VaCP17 CDS (Fig. 4a, b), and immunoblotting using anti-GFP antibody (Fig. 4c). Western blot analysis revealed an immunoreactive species of the expected VaCP17-GFP molecular mass (~76 kDa) in lines OE-1, OE-3 and OE-5, and these three lines were further evaluated (Fig. 4c). Based on these analyses, we selected three strongly expressing T1 individuals (OE-1, OE-3 and OE-5) and established homozygous lines for further study.

Fig. 2 Subcellular localization of VaCP17. 35S::VaCP17-GFP was transformed into Arabidopsis protoplasts by a PEG mediated method. Cells were observed using fluorescent microscopy. Bars, 20 μm

Fig. 3 Developmental expression patterns of CP17 in ‘Shuangyou’ and ‘Red Globe’. SY. ‘Shuangyou’; RG. ‘Red Globe’

Fig. 4 Identification and analysis of 35S::VaCP17-GFP Arabidopsis. a qRT-PCR analysis of VaCP17 expression in ten independent 35S::VaCP17-GFP Arabidopsis lines. M. DNA Marker III; 1–10, 35S::VaCP17-GFP lines; 11–12, positive plasmid control; 13, Negative control. b Quantification of qRT-PCR analysis for lines OE-1, OE-3, and OE-5. Non-transgenic Arabidopsis was used as a negative control. c Western blot analysis of VaCP17-GFP protein expression in lines OE-1, OE-3, and OE-5. WT, non-transgenic control. The predicted molecular mass for VaCP17 is 51.42 kDa, while that for GFP protein is 25 kDa, so the total expected molecular mass of VaCP17-GFP protein is 76.42 kDa
Overexpression of VaCP17 confers cold resistance in Arabidopsis

We noticed no striking difference in growth or development between 35S::VaCP17-GFP transgenic plants and non-transgenic controls when maintained at 22 °C (Fig. 5a). However, when plants were subjected to a freezing treatment of −6 °C for 8 h, non-transgenic controls showed severe wilting, whereas 35S::VaCP17-GFP plants showed only mild wilting. After a 10-day recovery period at 22 °C, almost all control plants died, while 35S::VaCP17-GFP lines quickly resumed normal growth (Fig. 5a). This result suggested that expression of VaCP17 significantly increased the resistance of Arabidopsis to transient freezing. Electrolyte leakage (EL) and malondialdehyde content (MDA) are two indicators of cell damage caused by freezing. After the freezing treatment, the contents of EL and MDA in both control and 35S::VaCP17-GFP plants increased, but the increases in control plants were much greater than in the 35S::VaCP17-GFP lines (Fig. 5b, c). These results indicated that the increased freezing tolerance presumably conferred by VaCP17 was associated with reduced cellular damage.

Plants under cold stress can increase synthesis of proline and soluble sugar as osmolytes, enhancing their water retention ability and cold resistance (Ashraf and Foolad 2007). As shown in Fig. 5d, e, both control and 35S::VaCP17-GFP plants accumulated proline and soluble sugar during the cold treatment. However, proline and soluble sugars both accumulated to higher levels in the 35S::VaCP17-GFP plants relative to the controls; this difference was most striking, and highly significant, for soluble sugars. In addition, we also evaluated the activities of three antioxidase enzymes that are known to be upregulated in plants under stress-superoxide dismutase (SOD), peroxidase (POD), and catalase (CAT). All three showed increased activity in the cold-treated plants; however, the increase in activities was significantly greater for the 35S::VaCP17-GFP lines (Fig. 5f–h). These results suggest that the active oxygen scavenging system in the 35S::VaCP17-GFP plants was more active under cold stress. In summary, the results of these physiological and biochemical experiments demonstrate that VaCP17 could confer increased cold tolerance to Arabidopsis.

Fig. 5 Heterologous expression of VaCP17 in Arabidopsis enhances cold resistance. a Plants of non-transgenic control (WT) and 35S::VaCP17-GFP lines (OE-1, OE-3 and OE-5) before and after a chilling treatment, and after recovery at 25 °C. b Electrolyte leakage and c malondialdehyde (MDA) content. d Proline content and e soluble sugar content. f–h Activities of SOD, POD and CAT, respectively. Data are the mean (± SD) of three biological replicates. Significance was analyzed with SPSS using the LSD and Duncan methods of one-way ANOVA (*P < 0.05; **P < 0.01, ***P < 0.001)
**VaCP17 promotes cold resistance-related genes expression in Arabidopsis**

To further explore the mechanism by which VaCP17 promotes cold tolerance, we analyzed the relative expression of VaCP17 and nine cold-resistance related genes (CBF1, CBF2, CBF3, RD29A, COR15A, KIN1, NCED3, AOC1 and JAZ10) in the 35S::VaCP17-GFP and control plants at various times after plants were moved to the cold (4 °C). Among them, CBF1, CBF2 and CBF3 are early cold-response genes, while RD29A, COR15A and KIN1 respond later to cold. AOC1 and JAZ10 are related to JA synthesis, and NCED3 is related to ABA synthesis (Jiang et al. 2014; Sun et al. 2020; An et al. 2021). Interestingly, although VaCP17 was expressed under control of the constitutive 35S promoter, expression of VaCP17 increased rapidly during the cold treatment, reaching a maximum at 6 h and then declining to near-initial levels by 24 h (Fig. 6). All of the nine cold-responsive genes were also induced during the cold treatment in both 35S::VaCP17-GFP and non-transgenic control plants (Fig. 7), as anticipated. However, each gene was expressed to significantly higher levels in the representative 35S::VaCP17-GFP line tested, at least one time point. These observations suggest that VaCP17 increased the cold resistance of Arabidopsis by promoting cold-responsive gene expression and participating in the JA and ABA pathways.

**Identification of VaCP17-interacting proteins**

To further our knowledge of potential mechanisms by which VaCP17 may promote cold tolerance in grapevine, we used the yeast two-hybrid approach to identify proteins that interact with VaCP17. Yeast colonies containing VaCP17/pGBKT7 did not grow on the SD/-Trp/AbA/X-α-gal medium and did not develop a blue color (Fig. 8a), indicating that VaCP17 had no independent transcriptional activation activity and could be used for subsequent cDNA library screening. Sixteen cells were plated after co-transformation of bait vector plasmid VaCP17/pGBKTK7 and library plasmid constructed from the leaves of cold-treated ‘Shuangyou’, and three candidate genes associated with low temperature were detected as VaNAC72, VaDi19, and VaCAM7.

All three candidate genes were inserted into the pGADT7 vector and co-transformed into Y2HGold yeast cells with VaCP17/pGBKTK7 for verification (Fig. 8b). These results confirmed that VaCP17 could interact with VaNAC72, VaDi19 and VaCAM7 in yeast cells.

**VaCP17 interacts with VaNAC72, VaDi19 and VaCAM7 in vivo**

We tested whether these three proteins could interact with VaCP17 in vivo using bimolecular fluorescence complementation (BiFC). The CDS sequence of VaCP17 was inserted into the PBI221NE vector to generate PBI221NE/VaCP17. At the same time, the CDS sequences of VaNAC72, VaDi19 and VaCAM7 were inserted into PBI221CE to generate PBI221CE/VaNAC72, PBI221CE/VaDi19 and PBI221CE/VaCAM7. As shown in Fig. 8c, YFP signal could be observed upon co-transformation of PBI221NE/VaCP with PBI221CE/VaNAC72, PBI221CE/VaDi19 or PBI221CE/VaCAM7, while no YFP signal was observed after co-transformation of PBI221NE/VaCP with the non-modified PBI221CE plasmid. These results supported a physical interaction of VaCP17 with VaNAC72, VaDi19 and VaCAM7 in vivo.

**Discussion**

Cysteine proteinases play important roles in plant growth and development, senescence and programmed cell death, and it has been reported that they may participate in response to various stresses (Xu et al. 2003; Zang et al. 2010; Zheng et al. 2018). However, to date, there have been no reported studies of cysteine proteinases in grapevines. In this study, we explored the potential position of VaCP17 in mechanisms of cold and freezing tolerance of *V. amurensis*. Similar to cysteine proteases identified in other species, the VaCP17 protein contains three conserved domains, Inhibitor_I29, Peptidase_C1A and GRAN, indicating that it belongs to the PapainC1A family (Drenth et al. 1968; Rawlings and Barrett 2010). The phylogenetic tree shown in Fig. 1 indicates that VaCP17 is closely related to AtCP from Arabidopsis. In Arabidopsis, the *AtRD19A* and *AtRD21* genes encode
cysteine proteinases included in the papain family, and can be upregulated in response to drought or salt stress (Masa-hiro et al. 1993). So we speculated that VaCP17 may be involved in response to cold stress in grape.

Interestingly, we found that VaCP17 expression in 35S::VaCP17-GFP plants increased in response to cold, even though the gene was under control of the constitutive 35S promoter (Fig. 6). This suggests that VaCP17 expression is promoted through post-transcriptional mechanisms in Arabidopsis. Compared with non-transgenic control plants, the three independent Arabidopsis 35S::VaCP17-GFP lines showed better survival after a short freezing treatment, with higher proline and soluble sugar content, higher antioxidant activities, and lower MDA and EL (Fig. 5). In this study, the activities of three antioxidant enzymes (SOD, POD and CAT) in three transgenic lines were significantly higher than those in the non-transgenic control plants, indicating that VaCP17 is able to recruit the antioxidant enzyme system to scavenge ROS. This finding is similar to the previous demonstration that heterologous expression of a cysteine proteinase from S. matsudana in Arabidopsis was sufficient to enhance the antioxidant enzyme system (Zheng et al. 2018).

Plants respond to adverse environmental conditions by modulating the expression of many stress-responsive genes. To further explore the molecular mechanism by which VaCP17 promotes cold tolerance, we analyzed the expression of nine cold-resistance related genes under low temperature treatment at 4 °C. In this study, the expression of CBF genes, as well as RD29A, COR15A and KIN1, were induced to higher levels in 35S::VaCP17-GFP transgenic plants relative to non-transgenic control plants. In addition, JA and ABA synthesis genes and signal pathway genes were significantly induced in 35S::VaCP17-GFP transgenic plants relative to non-transgenic control plants. In addition, JA and ABA synthesis genes and signal pathway genes were significantly induced in 35S::VaCP17-GFP. Therefore, we speculate that VaCP17 may be involved in responding to JA and ABA signaling pathways and respond to cold stress by...
cooperating with these cold resistance related genes, thereby significantly improving the cold resistance of Arabidopsis.

To further reveal the mechanism by which \textit{VaCP17} promotes cold resistance, three proteins (\textit{VaNAC72}, \textit{VaDi19} and \textit{VaCAM7}) interacting with \textit{VaCP17} were identified by Y2H and BiFC (Fig. 8). NAC is one of the largest transcription factor families in plants (Puranik et al. 2012), and individual NAC proteins are involved in responses to various biotic and abiotic stresses (Takada et al. 2001; Mallory et al. 2004; Liu et al. 2014). For example, transgenic rice overexpressing \textit{ONAC022} are more tolerant to drought and salt stress (Hong et al. 2016). The strawberry \textit{FaNAC2} gene could be significantly induced by cold, salt and drought stresses, and transgenic tobacco expressing \textit{FaNAC2} showed increased resistance to cold and salt stresses (Liang et al. 2020). Di19 proteins are a subset of the Cys2/His2 zinc finger protein family, which plays an important role in regulating multiple stress responses (Millà et al. 2006; Qin et al. 2014). In Arabidopsis, \textit{AtDi19-1} and \textit{AtDi19-3} could be rapidly induced by drought stress, while \textit{AtDi19-2} and \textit{AtDi19-4} could be induced by salt stress (Millà et al. 2006). In addition, \textit{GhDi19-1} and \textit{GhDi19-2} are involved in plant response to salt stress and ABA signals (Li et al. 2010).

Calcium plays a key role in plant development and abiotic stress response as a second messenger (Dodd et al. 2010; Kudla et al. 2018). Plants complete the signal transduction process by increasing the concentration of intracellular calcium, which activates a complex intracellular defense mechanism (Yang and Poovaiah 2003). The cold tolerance of transgenic tomato overexpressing \textit{SICML37} was significantly improved. The conformation of \textit{SICML37} induced by Ca$^{2+}$ could interact with SIUMP1; activated SIUMP1 could affect the activity of proteasome, and regulate the degradation of target proteins, which improve plant cold tolerance (Tang et al. 2021). The \textit{VaCAM7} obtained by our screening is also a Ca$^{2+}$ sensor, which may be closely related to cold stress in plants. Therefore, it is speculated that the interaction between \textit{VaCP17} and \textit{NAC72}, \textit{Di19} and \textit{CAM7} may affect the expression of downstream genes, and then jointly regulate the tolerance of plants to cold stress.

In conclusion, the results of this study indicate that \textit{VaCP17} can enhance the tolerance of transgenic Arabidopsis to cold stress. However, more work is needed to clarify the underlying molecular mechanism.

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Author contributions ZJX designed the experiments and revised the manuscript. SX performed experiments and wrote the manuscript. GB cloned VaCP17. ZHI participated in RNA extraction from the 35S::VaCP17-GFP lines. TYY participated in the determination of physiological and biochemical indicators.

Data availability The data sets supporting the results of this article are included within the article and its additional files.

Declarations

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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