Rice TSV2 encoding threonyl-tRNA synthetase is needed for early chloroplast development and seedling growth under cold stress

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Abstract

Threonyl-tRNA synthetase (ThrRS), one of the aminoacyl-tRNA synthetases (AARSs), plays a crucial role in protein synthesis. However, the AARS functions on rice chloroplast development and growth were not fully appraised. In this study, a thermo-sensitive virescent mutant tsv2, which showed albino phenotype and lethal after the 4-leaf stage at 20°C but recovered to normal when the temperatures rose, was identified and characterized. Map-based cloning and complementation tests showed that TSV2 encoded a chloroplast-located ThrRS protein in rice. The Lys-to-Arg mutation in the anticodon-binding domain hampered chloroplast development under cold stress, while the loss of function of the ThrRS core domain in TSV2 fatally led to seedling death regardless of growing temperatures. In addition, TSV2 had a specific expression in early leaves. Its disruption obviously resulted in the downregulation of certain genes associated with chlorophyll biosynthesis, photosynthesis, and chloroplast development at cold conditions. Our observations revealed that rice nuclear-encoded TSV2 plays an important role in chloroplast development at the early leaf stage under cold stress.

Keywords: albino phenotype; chloroplast development; cold stress; rice; threonyl-tRNA synthetase

Introduction

Aminoacyl-tRNA synthetase (AARS) is an important component of protein synthesis in ribosomes and is essential for translation in three different compartments of the plant cell: chloroplasts, mitochondria, and the cytosol (Berg et al. 2005; Sang et al. 2005). AARSs generally consist of catalytic core domain, binding zinc ion domain, insertion domain, anticodon-binding domain, and editing domain (Greenberg et al. 2008) and involves in the process of amino acid transfer to its homologous tRNA (O’Donoghue and Luthey-Schulten 2003). AARS and tRNA play a key role in the first step of protein synthesis. According to the homology and folding pattern, AARSs can be grouped into Class I and Class II, and each AARS class can be further divided into three subclasses: A, B, and C (Martinis et al. 1999).

Previous studies have shown that AARS not only plays a critical role in protein synthesis but also participates in and regulates various biological processes, including RNA transcription and splicing, protein translation, signal transduction, and cell apoptosis (Hausmann and Ibba 2008). In Arabidopsis thaliana, several mutants involving in AARS genes have been identified. In the edd1 mutant, inactivation of GLYRS encoding glycyl-tRNA synthetase causes embryonic development to stagnate, ultimately leading to the death (Uwer et al. 1998). Mutations in the PRORSI encoding prolyl-tRNA synthetase in Arabidopsis thaliana lead to abnormal transcription levels of the photosynthesis and plastid-synthesis genes, also leading to seedling death (Pesaresi 2006). The mutations of NbERS and NBR5, encoding glutamyl-tRNA synthetase and serine-tRNA synthetase, respectively, resulted in abnormal chloroplast and reduction of chlorophyll content, and finally etiolated leaf phenotype (Kim 2005). As for ThrAS in A. thaliana, though the phenotype (albinism and embryo death) observation of the mutant and specific expression localization has been carried out, its molecular mechanism is, however, still unclear (Berg et al. 2005; Duchêne et al. 2005).

Rice is the most important food crop in Asia and has been set up as a model species for genome study. Low temperature is a serious abiotic stress in rice production, which hinders a broad spectrum of cellular components (e.g. chloroplast), metabolisms (e.g. photosynthesis), plant growth, and yield. In spite of the facts that some indispensable genes for chloroplast development/seedling growth in rice at low temperatures have been identified, such as TCD9 (Jiang et al. 2014), V1 (Kusumoto et al. 2010b), V2 (Sugimoto et al. 2007), V3 (Yoo et al. 2009), OsV4 (Gong et al. 2014), TCD5 (Wang et al. 2016a), TCD10 (Wu et al. 2016), TCD11 (Wang et al. 2017), TSV3 (Lin et al. 2018), TCM12 (Lin et al. 2019), TCD33 (Wang et al. 2020), and TCD3 (Lin et al. 2020), the molecular mechanism of cold resistance and its impact factors are not well understood.
Materials and methods
Plant materials and growth conditions
The rice thermo-sensitive virescent mutant tsv2 was discovered in our mutant pool from Jiahua 1 [wild type (WT), japonica variety] treated with Cobalt-60 gamma rays. The mutant phenotype was distinguishable from normal green at Hainan Island, China (winter season, subtropical climate) and Shanghai, China (spring season, temperate climate) under local conditions during the early seedling stage. The tsv2 mutant was crossed with Peiai64S ( indica variety) and the obtained F2 seeds were used for genetic analysis and gene cloning. WT and tsv2 plants were cultured in incubators under controlled 12 h of light and 12 h of dark at a constant temperature of 20, 24, 28, and 32°C, respectively, for phenotypic characterization, photosynthetic pigment analysis, and DNA and RNA extractions.

Measurement of photosynthetic pigments and transmission electron microscopy
For the photosynthetic pigment analysis, 200 mg of fresh leaves were taken from the 3-leaf-stage seedlings cultured at 20, 24, 28, and 32°C and incubated with 5 mL of extraction buffer (ethanol:acetone:water = 5:4:1) at 4°C in the dark for 18 h. Using spectrophotometer (Beckman Coulter, Danvers, MA, USA), chlorophyll a, b, and Car contents were measured according to the modified methods of Arnon (1949) and Wellburn (1994). This experiment was carried out in three biological replicates.

To observe ultrastructure of chloroplast, tissues from the 3rd leaves of the 3-leaf-stage WT and tsv2 seedlings, grown at 20 and 32°C, respectively, were sampled and treated with the mixed solution of 3% glutaraldehyde and 2.5% paraformaldehyde. The observation of chloroplast was performed following Jiang et al. (2014). Samples were viewed under a Hitachi765 (Hitachi, Tokyo) transmission electron microscope (TEM).

Mapping and cloning of TSV2 gene
Total genomic DNA from rice fresh leaves were extracted by CTAB method described in Murray and Thompson (1980). DNA-specific fragments were amplified by EDC-810 PCR instrument (Eastwin, Shanghai, China). PCR products were transferred to 2–4% agarose gel containing ethidium bromide for electrophoresis. The bands were observed and recorded under UVP imager. The F2 population of 1308 individuals with the mutant phenotype was used for fine mapping of the TSV2 locus. First of all, ninety-two SSR primers based on the Gramene database (http://www.gra.mene.org) were used to investigate the chromosome of the target gene, and then developed SSR and InDel markers (Supplementary Table S1) were used for fine-mapping of TSV2. Next, DNA fragments of the candidate genes were PCR amplified and sequenced (SinoGenoMax, Shanghai, China). Lastly, the function and open reading frames of the candidate genes were obtained from TIGR (http://rice.plantbiology.wsu.edu/cgi-bin/gbrowse/rice/) and conserved domain structures were predicted using SMART (http://smart.embl-heidelberg.de/).

Complementation experiment
First, the WT genomic DNA fragment covering the entire TSV2 (LOC_Os02g33500) coding region (3.9 kb), plus a 1.4-kb upstream region and a 0.6-kb downstream sequence was amplified using the specific primers, pF: 5′-TACGAACTGAGCTCGGTACTTCCACCAAAGTTACGAGGCC-3′ (KpnI) and pR: 5′-GCTGTGAAGACCCGCCCTATGACGAGTGAGGCATGAAG-3′ (SalI). The underlined sequences stand for cleavage sites of the restriction enzymes. Then, the amplified fragment was cloned onto vector pCAMBIA1301 (CAMBIA, http://www.cambia.org.au), the pCAMBIA1301-TSV2 plasmids were transferred into Agrobacterium EHA105 and introduced into the tsv2 mutant by Agrobacterium tumefaciens-mediated transformation (Hei et al. 1994), except that the temperature used for in vitro plant differentiation was set at 20°C. The genotype of transgenic plants was determined using PCR amplification of the hygromycin phosphotransferase gene (HPT) with primers HPTF (5′-GGACACATTATAGCCGGGAGT-3′) and HPTR (5′-GGTTATACCGACCTTTGCACTG-3′) and GUS gene with primers GUSF (5′-GGGACATTACCGAGCCCAAGT-3′) and GUSR (5′-GCCGACAGGCAAGGTTCAT-3′) as selection. In addition, all T1 seedlings were cultured at 20°C and were used to examine the segregation of the mutant phenotype.

Targeted mutation of TSV2 gene
To determine if the novel allelic mutants in TSV2 display the similar or more severe phenotype compared with the tsv2 mutant, CRISPR/Cas9 technique was used for the targeted mutation. First of all, adaptor primers (pF1: 5′-GCCGGCGCTCGCTCGGTCACCTTCTGGT-3′ and pR1: 5′-AAACCACGAGACCGAGCTGAGC-3′; pF2: 5′-GCGGGGACATTACCGAGCCCAAGG-3′ and pR2: 5′-AAACCTTGCGCTGCGTGACCCGCGCC-3′) were designed by CRISPR Primer Designer (Naito et al. 2015). The sequence was inserted into the region between the OsU6 promoter and the gRNA scaffolds, from pYLgRNA-OsU6vector, of Cas9 expression backbone vector (pYLCRISPR/Cas9-MH) at the BsaI site according to the previous methods (Ma et al. 2015). The vector was transformed into the WT callus through A. tumefaciens-mediated transformation (Hei et al. 1994), except that the temperature used for in vitro plant differentiation was set at 20°C. The genotype of T0 transgenic plants was determined using PCR amplification of the hygromycin phosphotransferase gene (HPT) with primers HPTF (5′-GGGACACATTATAGCCGGGAGT-3′) and HPTR (5′-GGTTATACCGACCTTTGCACTG-3′). Then, the edited-edited T0 and T1 plants were determined by sequencing using PCR amplification of TSV2 gene with primers TSV2F (5′-ACCTCCACC AAGATTTACGAGGCC-3′) and TSV2R (5′-CATGTCGACCTCCAGG CATGCAAG-3′). Lastly, all T1 edited seedlings were grown at both 20 and 32°C, respectively, to observe the phenotype and the segregation.

Subcellular localization of TSV2
To determine the subcellular localization of TSV2, the cDNA fragment of TSV2 gene was amplified by PCR using the corresponding primer pairs (pF: 5′-GAAGATCTATGGCAGGCGCTCGCTGC-3′...
and pR: 5’-GGGGTACCCCGGTGTGGCGGATGCGGAGCA-3’). The PCR products were cloned into the pMON530-GFP vector, which was transformed into tobacco (Nicotiana tabacum) mesophyll cells through Agrobacterium-mediated infection. Meanwhile, empty pMON530-GFP vector was used as control. The analysis was carried out based on previously described method (Jiang et al. 2014).

Sequence and phylogenetic analyses
The protein domain prediction website InterPro was used to analyze the sequence and domain of TSV2, and Phyre2 (http://www.sbg.bio.ic.ac.uk/phyre2/html/) was used to predict the three-dimensional structure of protein. NCBI (http://www.ncbi.nlm.nih.gov) was employed to query the homologous proteins of TSV2 in other species, and MEGA6 and DNAMAN were used for phylogenetic tree analysis and homologous sequence alignment.

Transcriptional expression analysis
Total RNA was isolated from fresh tissues (root, stem, leaf, panicle) of WT and tsv2 plants using an RNA Prep Pure Plant Kit (TianGen, Beijing, China). The transcription of chlorophyll synthesis, chloroplast development, photosynthesis-associated genes (Cab1R, CAO, HEMA, YGL1, LHCF11, PORA, PsaA, PshA, RpoC, RpoC21, 16S RNA, 23S RNA) and temperature-sensitive genes for chloroplast development (TCD5, TCD9, TCD10, TCD11) (Supplementary Table S2), in rice was assessed using quantitative real-time PCR (RT-qPCR). The specific primers for qPCR are listed in Supplementary Table S2. A SYBR Premix Ex Taq™ RT-PCR Kit (Takara, Japan) was performed according to manufacturer’s instructions. ABI-7500™ Real-Time PCR System (Applied Biosystems; http://www.appliedbiosystems.com) was used to perform the analysis. The relative quantification of gene expression data was analyzed as previously described (Livak and Schmittgen 2001). Actin was used as an internal control. This experiment was carried out with four biological replicates.

Results
Phenotypic characterization of the tsv2 mutant
The growths of tsv2 and WT seedlings were observed under four growing temperatures (20, 24, 28, and 32°C) (Figure 1). All WT seedlings expectedly displayed green normal phenotype, regardless of temperatures and leaf-stage. However, the tsv2 mutant was albino phenotype from the beginning and died after 4-leaf stage at 20°C (Figure 1A). At 24°C, the tsv2 mutant turned to yellow but not lethal (Figure 1B). At 28°C, leaves of tsv2 were close to green (Figure 1C). Interestingly, at 32°C, the tsv2 mutant exhibited the normal green as WT plants (Figure 1D). The observations suggest that tsv2 is a low-temperature sensitive lethal mutant.

Consistent with the phenotype, the contents of photosynthetic pigments in tsv2 mutants were the lowest at 20°C (Figure 2A), and gradually upraised to the WT level (Figure 2, B–D) with increase of temperatures, indicating that chlorophyll accumulation in tsv2 mutants was suppressed under low temperature. By TEM observation, WT mesophyll cells were found to contain a lot of uniform chloroplasts, irrespective of temperatures (Figure 3, A–D). Nevertheless, the majority of cells in tsv2 mutants at 20°C contained only few chloroplasts and the structure was abnormal with the grana in the inner capsule decreased obviously (Figure 3, E and F).

![Figure 1](https://example.com/figure1.png)

**Figure 1** Phenotypic characterization of the tsv2 mutants. Seedlings of Jiahua1 (WT, left) and tsv2 mutant (right) at the 2-, 3-, and 4-leaf stages grown at (A) 20°C, (B) 24°C, (C) 28°C, and (D) 32°C.
Interestingly, at 32°C, the tsv2 chloroplasts were not obviously different from WT plants (Figure 3, G and H). It was then speculated that aberrant chloroplast of tsv2 resulted in the accumulation alleviation of chlorophyll and mutant phenotype under cold stress.

Map-based cloning of TSV2

To understand the molecular mechanism underlying the tsv2 mutant phenotype, map-based cloning was performed to identify the TSV2 locus. In view of an approximately 3 (green):1 (albino) ratio (χ² = 0.488 < χ²₀.₀₅ = 3.84) in F₂ segregating population (Supplementary Table S3), consisting of 231 green plants and 70 albino plants at 20°C, this showed the mutant phenotype was controlled by a recessive nuclear gene (tsv2). First, ninety-two F₂ mutant individuals were used for initial mapping, and the target gene TSV2 was located between ID12613 and MM3298 molecular markers on chromosome 2 (Figure 4A). Subsequently, the mapping F₂ population was expanded to 1308 individuals, and the TSV2 gene was narrowed to 131 kb between ID12947 and ID13097, including eight candidate genes (Figure 4B). Sequence and
analyze of all candidate genes found that, only a A-to-G mutation in the last exon of LOC_Os02g33500, which contains eight exons (http://rice.plantbiology.msu.edu/index.shtml) occurred, resulting in Lys(K)-to-Arg(R) mutation of the 614th site in TSV2 (Figure 4C).

**Complementation of the tsv2 mutants**

To assert that the mutation of LOC_Os02g33500 was responsible for the tsv2 phenotype, an expression plasmid containing the entire TSV2 (LOC_Os02g33500) driven by its endogenous promoter was constructed and transformed into the tsv2 mutants. To speed up, we intentionally induced the differentiation of rice callus at 20°C accompanying the uninfected calli as a control. Resultantly, 15 T0 transgenic seedlings harboring pCAMBIA1301: TSV2 were apparently green as WT plants, while control seedlings remaining albino phenotype (Figure 4D). This acknowledges that LOC_Os02g33500 can rescue the mutant phenotype. In addition, the segregation of albino phenotype in the transgenic T1 population was found at 20°C (Figure 4D). All those together assure that LOC_Os02g33500 is TSV2.

**Characterization of TSV2 protein**

Bioinformatic assay revealed that TSV2 encodes ThrAS protein, consisting of 675 amino acids, with a molecular mass of ~76.9 kDa, which belongs to Class A in Class II of AARS family. The ThrAS contains at least two domains, including core ThrAS domain and anticodon-binding domain (Figure 4C, Supplementary Figure S1). It was noted that the Lys-to-Arg mutation in tsv2 mutant occurred in the anticodon-binding domain (Figure 4C, Supplementary Figure S1).

Orthologs of rice TSV2 were identified in Zea mays, Sorghum bicolor, Setaria italic, Aegilops tauschii, Brachypodium distachyon, A. thaliana, Cicer arietinum, Nicotiana sylvestris, Vitis vinifera, and Ziziphus jujuba. TSV2 was very similar to many species and highly conserved in different higher plants (Figure 5A). Phylogenetic analysis showed that the evolutionary relationships of TSV2 homologs were consistent with the taxonomy (Figure 5B). Also, the mutated amino acid (Lys, K) in TSV2 was highly conserved among higher plants (Figure 5A) and its Lys-to-Arg mutation created the change of a β sheet structure (Supplementary Figure S2). This demonstrated the importance of this site for the functional integrity of the TSV2 protein.

**Expression pattern and subcellular localization of TSV2**

To clarify the expression pattern of TSV2 in rice, a semi-quantitative RT-PCR was carried out with various tissues (Figure 6). Consistent with the rice gene expression profiling data in the RiceXPro database (Supplementary Figure S3), TSV2 was highly expressed in the 1st and 2nd leaves, and weakly signaled in flag-leaf, root stem, and panicle (Figure 6A), showing the tissue-specific expression of TSV2. In addition, the TSV2 protein was predictively localized in the chloroplast using TargetP program (http://www.cbs.dtu.dk/services/TargetP/) (Emanuelsson...
Figure 5 Phylogenetic analysis of TSV2 protein. (A) Amino acid sequence alignment of TSV2 with the eleven homologous proteins from amino acids fully or partially conserved are shaded black and gray, respectively. (B) Phylogenetic tree of TSV2 and homologous proteins. Scale represents percentage substitution per site. Statistical support for the nodes is indicated.

Figure 6 Expression pattern and subcellular localization of TSV2. (A) Analysis of expression of TSV2 in different tissues by RT-PCR. OsActin was used as a control (the cycle number for OsActin was 28, the cycle number for TSV2 was 35). (B) Empty GFP vector without a specific targeting sequence. (C) TSV2-GFP fusion. The scale bar represents 20 μm.
et al. 2000). To verify the actual subcellular localization, the pMON530:CaMV3SS:TSV2-GFP plasmid was introduced into tobacco cells in the transient expression assay, with the pMON530:CaMV3SS-GFP vector as control. Observationally, the GFP fluorescence was co-localized with chlorophyll auto-fluorescence (Figure 6, B and C), affirming the localization of TSV2 in the chloroplast.

Versatility for TSV2 function

Zhang et al. (2017) early reported the existence of an allelic albino-lethal mutant (las) regardless of temperatures in rice, which is significantly different from the tsv2 phenotype reported in this study. This might imply the versatility for the function of LOC_Os02g33500(TSV2/LAS), depending on where its mutation site located. We performed targeted mutation of TSV2 gene in WT plants by CRISPR/Cas9 system. Surprisingly, all homozygous T0-edited transgenic seedlings, with two different edited-sites (5 and 7 base deletion, respectively) on the 1st exon of TSV2 gene (Supplementary Figure S4), displayed albino phenotype and eventually died when grown at 20°C (Figure 7A). In addition, all T1-edited homozygous seedlings from T0-edited heterozygous transgenic seedlings displayed all albino phenotype and finally died when grown at 20 and 32°C (Figure 7B). Owing to the exon 1 sequence determined the ThrRS core domain (Figure 4C, Supplementary Figure S1); thus, the loss of function in the core domain of LOC_Os02g33500 fatally led to albino death, regardless of temperatures. This is consistent with allelic las mutant (Zhang et al. 2017). Therefore, the expression of the core domain is co-regulated by other domains and temperatures, indicating the versatility for TSV2 function for chloroplast development and seedling growth.

The disruption of TSV2 alters the transcript levels of associated genes

To elucidate the effect of the tsv2 mutation on the expression of genes related to chloroplast development and to explore the regulating pathway, we performed RT-qPCR analysis of 20 genes involved in chlorophyll biosynthesis, photosynthesis, chloroplast development, and temperature sensitivity in rice (Supplementary Table S2). It was observed that transcript levels of all tested genes...
for Chl biosynthesis, i.e., chlorophyllide a oxygenase1 (CAO1), glutamyl tRNA reductase (HEMA), NADPH-dependent protochlorophyllide oxidoreductase (PORA), and Chl synthetase (YGL1), and for photosynthesis (Cab1R, RbcL, PsA, PsbA, LhcbII) were significantly downregulated under low temperatures (20°C) (Figure 8, A and B), aligned with the observed albino phenotype of tsv2 mutants (Figure 1); Also, at 20°C, the transcript levels of 16SrRNA (small subunit) and 23SrRNA (large subunit) involved in ribosome assembly and Rps20, encoding the small ribosomal subunit S20, in chloroplast, were greatly blocked (Figure 8C). Furthermore, at 20°C, except for TCD9 encoding chloroplast chaperone protein OsCpn60a subunit (Jiang et al. 2014), three temperature-sensitive genes (TCD5, TCD10, and TCD11) for chloroplast development (Wang et al. 2016a, 2017; Wu et al. 2016) showed a significant downward trend of transcripts, in particular, TCD10, encoding PPR protein, and TCD11, encoding plastid ribosomal protein S6 (Figure 8D). Conversely, all transcripts of the reduced genes at 20°C in the tsv2 seedlings recovered to or even exceeded WT levels (within twofold range) when grown at 32°C (Figure 9, A–D), coincident with the recovery of leaf-color (Figure 1D) and chloroplast development (Figure 3, G and H) in tsv2 at 32°C. As such, our data revealed that the tsv2 mutation leads to dramatic downregulation of many genes for chlorophyll biosynthesis, photosynthesis, and chloroplast development, under cold stress. In addition, the TSV2 influences chloroplast ribosome assembly thereby affecting the process of chloroplast development and accumulation of photosynthetic pigment.

**Discussion**

In this study, we identified and characterized rice tsv2 mutants with imperfect chloroplasts and albino lethal phenotype at low temperatures, resulting from the abnormal expression of genes associated with chlorophyll biosynthesis, photosynthesis, and chloroplast development. The tsv2 mutant phenotype was caused by the Lys-to-Arg mutation in anticodon binding domain in TSV2 gene. However, the loss of function in core domain for TSV2 fatally led to seedling death regardless of temperatures. Our data evidenced that TSV2 plays a critical role in chloroplast development and plant growth in rice.

**Incomplete of anticodon-binding domain in TSV2 leads to the albino phenotype under cold stress**

To date, a lot of temperature-sensitive seedling-color mutants of rice similar to tsv2 phenotype have been reported, with phenotypic white or yellow leaves at low temperatures while become normal green at high temperatures. Mutant v1 and v2 showed yellow leaf at 20°C and recovered to WT plants at 30°C (Iba et al. 1991; Kusumi et al. 1997). The coding protein of V1 gene participates in the regulation of plastid RNA metabolism and protein translation (Kusumi et al. 1997). Guanylate kinase encoded by V2 gene plays a special important role in the early development of chloroplasts (Sugimoto et al. 2007). Also, at 22°C, osv5a leaves yellowed and whitened, but turned to green phenotype over 28°C. OsV5A gene encodes chaperone protein interacting with PORA and PORB and stabilizing PORB protein (Zhou et al. 2013; Liu et al. 2016; Zhou et al. 2018).
Besides, the mutant dua1 showed a pale phenotype at 19°C, PPR protein DUA1 is essential for a chloroplast ribosome development under cold stress (Cui et al. 2019). Moreover, our research group also reported some thermo-sensitive leaf color mutants of rice. For example, tcd9 mutant presented albino phenotype before the 3-leaf stage over 24°C and TCD9 encodes chloroplast chaperone protein OsCpn60α subunit (Jiang et al. 2014). Mutants of tcd3, tcd5, tcd10, tcd11, tcd33, tsv3, and osv4 exhibited albino or white phenotype and malformed chloroplast at 20°C, but return to normal phenotype at 32°C (Gong et al. 2014; Jiang et al. 2014; Wu et al. 2016; Wang et al. 2016a,b, 2007, 2020; Lin et al. 2018, 2020). More interestingly, TCD5, TCD10, and OsV4 encode respective novel PPR protein (Gong et al. 2014; Wang et al. 2016b; Wu et al. 2016). Notwithstanding, different from the aforementioned mutants, the edited-mutant seedlings for TSV2 gene in this study did not present temperature-sensitivity for leaf color but died at high temperatures (Figure 7B). Thus, it is interesting to reasonably explore why allelic las mutant (Zhang et al. 2017) and the TSV2-edited mutants showed different phenotypes from the tsv2 mutant. This is because that, whether edited-transgenic plants or als mutants, their mutation sites occur in the core ThrRS domain (Supplementary Figures S1 and S4). However, the mutation site of tsv2 was located on the terminal exon 8, which only destroyed an original β sheet of the anticodon-binding domain in TSV2 (Supplementary Figure S2). Among AARSs, anticodon-binding domain is found in histidyl, glycyll, threonyl, and prolyl tRNA synthetases (Wolf et al. 1999) and involved in anticodon stem-loop binding and recognition (Aberg et al. 1997). In this study, the destruction of the core domain will lead to the full loss of the function of TSV2, as its mutant phenotype has nothing to do with temperatures. Interestingly, the Lys-to-Arg mutation in the anticodon-binding domain only resulted in whitening under cold stress and the increase of temperature can make up for the defect of its domain function. Therefore, the mutations of different domains in TSV2 will result in different phenotypes. Conclusively, the incomplete of anticodon-binding domain is responsible for the tsv2 albino phenotype under cold stress.

**TSV2 functions at the first step of chloroplast development**

It is well known that chloroplast development is divided into three steps (Kusumi et al. 2010a,b; Kusumi and Iba, 2014). The first step involves the activation of plastid replication and plastid DNA synthesis. The second step involves the establishment of chloroplast genetic system. The last step involves the high expression of plastid and nuclear targets encoding the photosynthetic apparatus, resulting in the synthesis and assembly of the photosynthetic apparatus. At present, through utilization of thermo-sensitive leaf-color rice mutants, it was shown that V3 (Yoo et al. 2009), TCD9 (Jiang et al. 2014), TCD10 (Wu et al. 2016), TCD11 (Wang et al. 2017), TSV3 (Lin et al. 2018a), TCD33 (Wang et al. 2016) and involved in anticodon stem-loop binding and recognition (Aberg et al. 1997). In this study, the destruction of the core domain will lead to the full loss of the function of TSV2, as its mutant phenotype has nothing to do with temperatures. Interestingly, the Lys-to-Arg mutation in the anticodon-binding domain only resulted in whitening under cold stress and the increase of temperature can make up for the defect of its domain function. Therefore, the mutations of different domains in TSV2 will result in different phenotypes. Conclusively, the incomplete of anticodon-binding domain is responsible for the tsv2 albino phenotype under cold stress.

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of three ribosome subunit genes (Rps20/ASL1 in chloroplasts. With the increase in temperature, all expressions of 16SrRNA and 23SrRNA responses to temperatures. In this study, the expression of Rps6/TCD11, the second step (TCD5), and the third step (Cabr1, Rbcl, Psaa, Pheh, Ucplf) of chloroplast development in tsu2 albino leaves under stress were observed in this study. Obviously, the transcript levels of the first step TCD10 and TCD11 genes were largely downregulated (Figure 8D), indicating that TSV2 functions in the first step of chloroplast development. However, TCD9, which also involved in the first step, was still highly expressed (jiang et al. 2014). This was probably attributed to that TSV2 is localized between the downstream of TCD9 and the upstream of TCD10 and TCD11 and was not regulated by TCD9. In addition, those inhibitions (TCD10, TCD11) caused by TSV2 dysfunction in the first step, would definitely lead to the inhibition of certain associated-genes in its second (TCD5) and third (Cabr1, Rbcl, Psaa, Pheh, Ucplf) stages (Figure 8, A and B), resulting in the reduction of chlorophyll accumulation and photosynthesis, consequently, seedling death.

Possible role of TSV2 in chloroplast ribosome assembly and protein synthesis

Chloroplast development is a complex biological process, in which many key proteins are translated and formed in chloroplasts. The plastid ribosomal proteins are crucial to ribosome biosynthesis, plastid protein biosynthesis, chloroplast development, and ribosome assembly (Zhao et al. 2016). In the past, several genes for chloroplast ribosome assembly were reported in rice, such as WLP1 encoding ribosome protein L13, Rpl13 (Song et al. 2014), ASL1 encoding ribosomal protein S20, RPS20 (Gong et al. 2013), ASL2 encoding ribosomal protein L21, RPL21 (Lin et al. 2015), and TCD11 encoding plastid ribosomal protein S6, RPS6 (Wang et al. 2017). As we have known, the mutation of TCD11/Rps6 (Wang et al. 2017) and WLP1/Rpl13 (Song et al. 2014) produced the thermo-sensitive leaf-color as the tsu2 mutants, but the mutation of ASL1/Rps20 (Gong et al. 2013) and ASL2/Rppl21 (Lin et al. 2015) led to seedling death, regardless of temperatures, indicating that plastid ribosome subunits have different functions or responses to temperatures. In this study, the expression of Rps20/ASL1, Rpl21/ASL2, and Rps6/TCD11 in tsu2 mutants at low temperatures decreased greatly, evidently suggesting that TSV2 affected chloroplast ribosome synthesis under cold stress. Notably, 16SrRNA and 23SrRNA were severely hampered (Figure 8C), definitely resulting in impaired translation and protein synthesis in chloroplasts. With the increase in temperature, all expressions of three ribosome subunit genes (Rps20/ASL1, Rpl21/ASL2, Rps6/TCD11) gradually returned to normal levels, especially for the expression of chloroplast ribosome 16SrRNA and 23S rRNA (Figure 9C). It is recognized that chloroplast translation occurs in 30S and 50S ribosomes; both are important components and their changes will directly affect ribosome assembly. Therefore, in the light of the reduced transcripts of 16SrRNA, 23SrRNA, Rps20, Rpl21, TCD11/Rppl6 in albino tsu2 seedlings (Figure 8, C and D), this once again proved that TSV2/LAS is involved in the chloroplast ribosomes assembly and protein synthesis, consistent with Zhang et al. (2017). In conclusion, our data clearly revealed that the TSV2 encodes a rice ThrRS, which is located in chloroplasts and is closely related to the assembly of chloroplast ribosomes and functions at the first step of chloroplast development. Mutation in the anticodon-binding domain of TSV2 causes chloroplast ribosomes to fail to assemble normally, resulting in impaired chloroplast development and lethal phenotype of rice seedlings under cold stress, but the loss of function in the core domain fatally led to seedling death, regardless of temperatures. Further work is warranted to explore from mechanism the roles of TSV2 in chloroplast development and plant growth.

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Data availability

All data generated or analyzed in this study are included in this article and its supplementary information files.

Supplementary material is available at G3 online.

Conflicts of interest

The authors declare that they have no competing interests.

Authors’ contributions

DL and YD provided the mutant seeds and generated F1 and F2 seeds for genotyping and phenotyping. DL, WZ, JS, YW, XP and YD performed the experiments of phenotype assays and molecular analysis. DL, XP and YD designed and discussed the research. DL, YD and XP wrote the manuscript. All authors approved the manuscript.

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