Deficiency of a peroxisomal NADP-isocitrate dehydrogenase leads to dwarf plant and defect seed in upland cotton

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The NADP-isocitrate dehydrogenase-encoded gene GH_D13G1452 with a C-terminus tripeptide Proline-Lysine-Leucine was localized in the peroxisome. It was highly expressed in stems and ovules of 15 days post-anthesis and responded to multiple external stimuli in upland cotton. An upland cotton mutant (Ghpericdh) was identified by flanking sequence amplification and genome variation detection that exogenous sequence was inserted in the middle of the 12th intron of GH_D13G1452, resulting in the deficiency of gene expression. The GhperICDH mutant displayed a dwarf plant phenotype when grown under field or greenhouse conditions, and GH_D13G1452 functioned as an incomplete dominance on plant height. The germination rate of mutant seed from greenhouse-grown plants was dramatically lower than that from field-grown plants, which indicated that GhperICDH plays a critical role in seed maturation and germination. Therefore, GH_D13G1452 is indispensable in the development of stems and seeds and functions in the adaptability of cotton to the environment. The Ghpericdh mutant provides insight into the function of peroxisomal ICDH and may contribute to the genetic improvement in cotton.

KEYWORDS
perICDH, Gossypium, plant height, seed development, subcellular localization

Introduction

NADP-dependent isocitrate dehydrogenase (ICDH, EC. 1.1.1.42) catalyzes the oxidative decarboxylation of isocitrate to form 2-oxoglutarate and NADPH, which are involved integrally in ammonia assimilation and reactive oxygen species metabolism (Hodges et al., 2003). Through subcellular localization, various ICDH isozymes were identified, such as cytICDH localized in the cytoplasm, mitICDH in the mitochondrion, chlICDH in the chloroplast, and perICDH in the peroxisome, of which the activity of cytICDH accounts for more than 90% of overall isozymes (Mhamdi et al., 2010). In Arabidopsis, deficiency of cytICDH or perICDH does not
affect plant growth under normal conditions (Mhamdi et al., 2010; Mhamdi and Noctor, 2015). Deletion of perICDH results in stomata semi-closed without a detrimental effect on the overall ICDH activity and redox homeostasis in Arabidopsis (Mhamdi and Noctor, 2015; Leterrier et al., 2016). These studies in Arabidopsis suggested that perICDH was not necessarily involved in plant growth, which can be explained by 2-oxoglutarate which is mainly produced by mitochondrial NAP-dependent isocitrate dehydrogenase (EC. 1.1.1.41) (Behal and Oliver, 1998), and the NADPH could be compensated by the pentose-phosphate pathway and NADH phosphorylation in plants (Corpas et al., 1998; Waller et al., 2010).

Here, we identified a cotton peroxisomal ICDH (GhperICDH) that was highly expressed in stems and seeds, and deficiency of GhperICDH would lead to abnormal plant growth and seed development in upland cotton.

Materials and methods

Plant materials

Gossypium hirsutum cv. TM-1 and mutant Ghpericdh were used here. TM-1 is a standard genetic line of upland cotton, obtained from USDA-ARS, College Station, TX, United States. The Ghpericdh is a deficiency mutant of the GH_D13G1452 gene, generated from CCRI49 as a receptor overexpressing a glyphosate resistance gene g10evo (Tan, 2016), which was developed by our lab for five generations, and its wild-type (non-1007), a non-transgenic line with normal GH_D13G1452 separated from the selfing transgenic plant T0, were used in the experiment. CCRI49 is a conventional cotton cultivar, which was provided by the Cotton Research Institute, Chinese Academy of Agricultural Sciences. All cotton plants were grown in the Agricultural Station field from May to September (Zhejiang University, Hangzhou) or in a greenhouse at 28°C/25°C under a 14-h photoperiod with a light intensity of 35000 lx.

DNA and RNA extraction and PCR

DNA and total RNA were extracted from fresh young leaves. cDNA synthesis, RT-PCR, and qPCR were performed according to Cao et al. (2021). Fusion primer and nested integrated PCR (FPNI-PCR), used for flanking sequence amplification, were designed according to methods described previously (Li, 2016; Xu, 2017). Primers in this study are listed in Supplementary Table 1.

Southern blot

The genomic DNA of 30 µg was digested completely with Hinde, separated by 0.8% gel electrophoresis, transferred into a nylon membrane (Amersham, United Kingdom), and hybridized with digoxin-labeled DNA fragments of g10evo at 65°C overnight. The signaling was detected by the image analyzer FLA-5100 (FUJIFILM, Japan). Detailed procedures were as described in the DIG High Prime DNA Labeling and Detection Starter Kit II (Roche, Switzerland).

Subcellular localization

Full-length open-reading frames (ORFs) of GH_D13G1452 fused with a superfolder green fluorescent protein (sGFP) on its N-terminus or C-terminus and driven by CaMV35S (pCAMBIA1300 vector), transiently co-expressed with the known peroxisomal markers 984 fused with mCherry in epidermal cells of tobacco via Agrobacterium (Nelson et al., 2007). The GFP and mCherry fluorescence in epidermal cells of tobacco was detected and photographed by a laser confocal microscope (Olympus, Japan) after injection from 48 to 72 h.

Vector construction and genetic transformation in cotton

The complete cDNA of gene g10evo (Tan, 2016), amplified from Deinococcus Radiodurans, was ligated into the overexpression vectors pCAMBIA-1300 (driven by the CaMV35s promoter) and transformed into Gossypium hirsutum cv. CCRI49 according to methods described previously (Yan, 2011). The transformants were selected on a selective medium containing 2.0 M glyphosate.

Stress treatment

The TM-1 seedlings were transferred into a plastic bucket of 1 L with a 1/2 MS culture medium. When growing to a two-leaf stage, plants were stressed by 10⁻⁴ M IAA, 10⁻⁵ M GA, 10⁻⁷ M ABA, 10⁻⁴ M JA, 10⁻³ M SA, 15% PEG6000, and 0.1-M NaCl for 3, 6, and 24 h with the non-treatment group as control. The roots were used for extracting RNA. The tendency (T) of the gene relative expression level over time under different treatments was calculated as \[ T = 2^{-\Delta C_t} \] of the treatment group - 2⁻(ΔCt) of the control group.

Measurement for plant height and germination rate

The plant height was measured from the base of hypocotyl to the first fully expanded true leaf. Here, the cotyledon is defined as the first node, the first leaf of the main stem...
is the second node, and so on. The distance between two adjacent leaves is defined as internode length. Internode length = Plant height/node numbers (each line repeated 6 individual plants). The germination rate was computed as the proportion of the germinated seeds in 15 days to the total seeds used (three replicates with 50 seeds in each one). Seed germination was carried out under a greenhouse at 28°C/25°C under a 14-h photoperiod with a light intensity of 35,000 lx.

Measurement for stomatal aperture

The experiment was started in the morning after 10 h of the dark cycle. Cotyledon detached from cotyledon-stage seedlings was floated on the incubation medium (10-mM MES, 50-mM KCl, 100-μM CaCl₂, pH = 6.1) in Petri dishes under light for 2 h to make stomata open fully (Xi et al., 2019). The lower epidermis was peeled off, transferred to a drop of incubation medium on a glass slide, and immediately observed under a microscope (Nikon Eclipse Ni, Japan) for stomata and photographed. The long axis and short axis of the stomatal aperture were measured to calculate the opening degree (OD). OD = short axis/long axis. Three individuals were randomly selected from each genotype, and three visual fields were randomly selected from each cotyledon.

Sequence analysis and phylogenetic tree construction

Pairwise sequence comparisons were conducted using ClustalW (Thompson et al., 1994). Sequence similarities were analyzed with BioEdit (Tiddmann, 2004). The phylogenetic tree was constructed by the neighbor-joining method with a bootstrap replication of 500 by using MEAG 5 (Tamura et al., 2011).

Genome variation detection

Whole-genome resequencing of the Ghpericdh mutant was carried out with Oxford Nanopore Technologies (ONT) by Novogene (Tianjin, China) (Supplementary Table 2). Based on the resequencing data, the reads that could be mapped to both the g10evo gene and the cotton genome were extracted with NextGenMap-LR software (Sedlazeck et al., 2018). Then, the BAM files were obtained by comparing the extracted reads with the cotton genome. The VCR files were obtained by detecting the structural variation of the extracted BAM files using sniffles software (Sedlazeck et al., 2018). Based on the results of variation detection, the insertion sequence was extracted and compared with the g10evo gene for similarity analysis.

Results

Identification for a perlCDH in upland cotton

GH_D13G1452 has a high similarity of 84.4% with the peroxisomal ICDH from soybeans in the amino acid sequence (Table 1) and belongs to the peroxisomal ICDH group based on phylogenetic analysis (Figure 1A). The peroxisomal ICDHs possess the type-I peroxisomal targeting signal (PTS1), a tripeptide sequence typically found at the C terminus of peroxisomal proteins (Gould et al., 1989), such as SKL existed in perlCDHs of soybeans and SRL in Arabidopsis perlCDH, while Proline-Lysine-Leucine (PKL) was observed in the C-terminal of GH_D13G1452 (Figure 1B). Subcellular localization further confirmed that the peroxisomal targeting signal was located at the C-terminus of GH_D13G1452, while the peroxisomal signals disappeared when their C-terminus fused GFP (Figure 1C). Based on the above results, we named GH_D13G1452 as GhperICDH.

Transcripts Per Million (TPM) data of upland cotton transcriptome (Hu et al., 2019) downloaded from CottonFGD were used to analyze the gene expression pattern in different tissues and organs. The GhperICDH was expressed in different tissues and organs with a peak level in stem (30.39 TPM) and had variable levels at different stages of the ovule and developed fibers with peak level (31.7 TPM) in ovules of 15 days post-anthesis (DPA) (Figure 1D). Cis-elements analysis showed that a large number of environmental response elements are found in the 3-kb region upstream of GhperICDH, besides the core elements of the promoter (Figure 1E), which indicates that GhperICDH responds to a variety of stress conditions. Treating with IAA, GA, ABA, JA, SA, PEG, and salt for 3, 6, and 24 h, the expression trend of GhperICDH showed that this gene had obvious responses to ABA, JA, SA, PEG, and salt (Figure 1F).

Identification of a GhperICDH-deficiency mutant

A GhperICDH-deficiency mutant with glyphosate resistance was identified, and we named it Ghpericdh. Southern blotting displayed one copy of g10evo in the mutant (Figure 2A). Ghpericdh plants could grow normally under the recommended concentration of glyphosate isopropylamine in a field (1.37 kg·ai-hm⁻²), while became damaged and displayed leaf malformation under high concentrations (4.10 and 6.83 kg·ai-hm⁻²). Wild-type (WT) plants died under any concentration of glyphosate isopropylamine (Figure 2B). FPNI-PCR revealed the accurate position of g10evo, which was located in the middle of the 12th intron of GH_D13G1452 (Hu et al., 2019; Figure 2C). Genome variation detection of the
mutant found only one site on 45,469,231 nt of chromosome D13 of *G. hirsutum* cv. TM-1 genome containing an insertion sequence that possessed a high similarity of 92.4% with the g10evo gene, which was consistent with the result of FPNI-PCR. *GH_D13G1452* did not express in mutant *Ghpericdh* but did express in WT (Figure 2D). As controls, the homologous gene *GH_A13G1507* and the housekeeping gene *GhUBQ7* did normally express in both mutant and WT (Figure 2D and Supplementary Figure 1). Based on the genome information of mutant *Ghpericdh*, two specific molecular markers 80Ln/LBSP2 and Rb2b/80R only amplified in *Ghpericdh* were designed (Figure 2E and Supplementary Table 1).

**Phenotypic characteristics of *Ghpericdh* mutant**

The mutant *Ghpericdh* displays a dwarf phenotype (Figure 3A), but it can grow, flower, and bear fruits. Whether the plants growing naturally in the field or cultivated in the greenhouse, the plant height and the internode length of mutants were significantly lower than those of WT (Figure 3B). The seeds from the mutant plants were shorter than WT seeds (Figure 3C). The greenhouse condition caused a decline in seed size of the mutant, which kept a similar trend in the WT as well (Figure 3C). Interestingly, the germination rate of seeds from greenhouse-grown mutant plants was dramatically decreased from 97.5 to 84.8% (Figure 3C). An observation from the *Ghpericdh* kernel found that seeds from the greenhouse appeared black (Figure 3D, marked by the red arrow) that became necrotic gradually during germination (Figure 3E), which led to growth stagnation. Similar to the Arabidopsis *pericdh* mutant, most stomata of *Ghpericdh* kept semi-closed (OD = 0.25-0.5) under light, while those of WT stayed open (OD ≥0.5) (Supplementary Figure 2).

**Discussion**

The content of ICDH was higher in roots, stems, and vascular bundles than in other tissues in other plants (Boiffin et al., 1998; Popova et al., 2002; Pascual et al., 2008). Overexpression *cytICDH* in poplar made plants stronger and higher with increased expression of genes related to vascular differentiation (Pascual et al., 2018). The *perICDH* could be activated by cadmium chloride (Romero-Puertas et al., 2006) and contributed to natural senescence (Corpas et al., 1999). However, the absence of *perICDH* does not cause abnormality in plant growth, except for stomata semi-closed in Arabidopsis (*Leterrier et al.*, 2016). Interestingly, in addition to semi-closed stomata like Arabidopsis *pericdh*, cotton *Ghpericdh* appears more abnormal phenotypes, such as dwarf plants and developmental defect seeds, and functioned as an incomplete

| Accession number | Host organism | Subcellular localization | Amino acid identity (%) | References |
|------------------|---------------|--------------------------|-------------------------|------------|
| AF095445         | Soybean       | Peroxisome               | 84.4%                   | Hodges et al., 2003 |
| AF155333         | Rice          | Cytoplast                | 83.5%                   | Hodges et al., 2003 |
| At5g54340        | Arabidopsis   | Peroxisome               | 82.5%                   | Leterrier et al., 2016 |
| X77944           | *N. tabacum*  | Cytoplast                | 82.5%                   | Hodges et al., 2003 |
| AC007789         | Rice          | Peroxisome               | 82.2%                   | Hodges et al., 2003 |
| Q06197           | Soybean       | Cytoplast                | 81.5%                   | Hodges et al., 2003 |
| At1g5930         | Arabidopsis   | Cytoplast                | 81.3%                   | Leterrier et al., 2016 |
| AAR05796         | Poplar        | Cytoplast                | 81.3%                   | Pascual et al., 2018 |
| X969728          | *N. tabacum*  | Mitochondrion/chloroplast| 65.8%                   | Hodges et al., 2003 |
| X92486           | Potato        | Mitochondrion/chloroplast| 64.6%                   | Hodges et al., 2003 |
| At5g14590        | Arabidopsis   | Mitochondrion/chloroplast| 64.2%                   | Leterrier et al., 2016 |
dominance on plant height. Additionally, the absence of GhperICDH made the development of seed extremely sensitive to the growth environment. GhperICDH was also found to be expressed with peak levels in stems and ovules of 15 days post-anthesis in upland cotton and responded to multiple external stimuli. Therefore, GhperICDH is indispensable in the development of stems and seeds and functions in the adaptability of cotton to the environment.
In cotton, studies mainly focus on the mapping and cloning of genes related to fibers (Mao and Cao, 2018), and the regulation mechanism of plant height is little known. Dwarf and dense cultivation of cotton can resist lodging, facilitate nutrient utilization, and mechanized operation, which could effectively improve the unit yield and reduce the labor input (Lou et al., 2021). Therefore, the study on plant dwarfing is of great significance in the high-yield breeding of cotton. Mutants are important materials for functional genomics research. The artificial mutants of wheat (Botticella et al., 2011), corn (Parry et al., 2009), tomato (Shirasawa et al., 2016), and rice (Abe et al., 2012) have accelerated their process in gene function resolving and genetic breeding. In cotton, the GhACT17D was found to regulate fiber elongation and plant height based on the Li1 mutant (Thyssen et al., 2017; Sun et al., 2019; Cao et al., 2021). Therefore, the cotton dwarf mutant Ghpericdh...
FIGURE 3
Phenotypic characteristics of Ghpericdh. (A) Plants growing for 5 months in the field. Scale bar = 10 cm. (B) Comparison of plant height, node number, and internode length between genotypes in the field for 5-month growing and in the greenhouse for 3 months cultured. (C) The kernel length and the germination rate of seeds harvested from the field and the greenhouse. Scale bar = 1 cm. (D) The kernel characteristics harvested in different locations. Scale bar = 1 cm. (E) Comparison of greenhouse-seed germination between genotypes. Scale bar = 1 cm. (F) Seedling morphology (the left) and average plant height (the right) of $F_1$. $^*p < 0.05$, $^{**}p < 0.01$. 

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is useful to reveal the molecular mechanism of plant height regulation in upland cotton. More observations in cytology and physiology need to be carried out in the *Ghpericdh* mutant, and the regulation mechanism of *GhperICDH* in cotton plant height and seed development should be studied further.

**Data availability statement**

The data presented in this study are deposited in the BioProject database of NCBI repository, accession number: PRJNA872936.

**Author contributions**

TZ, SZ, and YC designed the experiments and wrote the manuscript. YC and WW performed the experiments. JC manipulated plant materials. All the authors have read and approved the final manuscript.

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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.2022.1000883/full#supplementary-material

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