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Isolation and characterization of the aconitate hydratase 4 (*Aco4*) gene from soybean

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Abstract: Aconitase catalyzes the reversible isomerization of two tricarboxylic acids, citrate and isocitrate, during the Krebs cycle. Five aconitase genes, namely, *Aco1*, *Aco2*, *Aco3*, *Aco4*, and *Aco5*, have been identified in soybean. Previously, *Aco4* was mapped on chromosome 11. The purpose of this investigation was to isolate and sequence the candidate gene for *Aco4*. We mapped the *Aco4* gene to a 148 kb region on chromosome 11 that contained 19 predicted genes. One of these, *Glyma.11G080600*, codes for aconitate hydratase. Sequencing of two isozyme variants (A-line and B-line) for *Glyma.11G080600* revealed three synonymous and two non-synonymous substitutions. Perhaps, the two non-synonymous substitutions resulted in a variable isozyme pattern between the variants. *Glyma.11G080600* contains a catalytic domain and a swivel domain that are known to catalyze isomerization of citrate to isocitrate and swiveling conformational change in the enzyme mechanism, respectively. Conservation of both the domains implies the role of *Glyma.11G080600* in the interconversion of citrate and isocitrate. *Glyma.11G080600* is expressed in most tissues, with maximum expression in leaves. Phylogenetic analysis of 25 genes from different species displayed three major clusters. *Glyma.11G080600* (*Aco4*) and *Glyma.01G162800*, which are conserved in *Glycine max* (L.) Merr. and *Glycine soja* Siebold. & Zucc., may have common ancestry before *G. max* and *G. soja* split.

Key words: soybean, aconitase, linkage mapping, isozyme, *Aco4*.

Résumé : Pendant le cycle de Krebs, l'aconitase catalyse l'isomérisation réversible de deux acides tricarboxyliques (le citrate et l'isocitrate). Chez le soja, cinq gènes codent l'aconitase : *Aco1*, *Aco2*, *Aco3*, *Aco4* et *Aco5*. Jusqu'à présent, on situait *Aco4* sur le chromosome 11. Les chercheurs voulaient isoler et séquencer le gène correspondant à *Aco4*. Celui-ci se trouve dans une section du chromosome 11 de 148 kb sur laquelle figurent 19 gènes. L'un de ces derniers, *Glyma.11G080600*, code l'aconitase hydratase. En séquençant *Glyma.11G080600* chez deux variants de l'isozyme (lignée A et lignée B), les auteurs ont découvert trois substitutions synonymiques et deux qui ne le sont pas. Il se pourrait que la variation de l'isozyme chez ces lignées résulte des deux substitutions non synonymiques. *Glyma.11G080600* comprend un domaine catalytique et un domaine pivot, le premier catalysant l'isomérisation du citrate en isocitrate et le second commandant les changements de conformation dans ce mécanisme. La conservation des deux domaines signifie que *Glyma.11G080600* joue un rôle dans la conversion du citrate en isocitrate, et vice-versa. Le gène *Glyma.11G080600* s'exprime dans la plupart des tissus, mais son expression est à son maximum dans les feuilles. L'analyse phylogénétique de 25 gènes de l'espèce révèle l'existence de trois grandes grappes. *Glyma.11G080600* (*Aco4*) et *Glyma.01G162800*, qu'on retrouve chez *Glycine max* (L.) Merr. et *Glycine soja* Siebold & Zucc., pourraient avoir des ancêtres communs précédant la différenciation des deux espèces. [Traduit par la Rédaction]

Mots-clés : soja, aconitase, cartographie des liaisons, isozyme, *Aco4*.

Introduction

The aconitase gene produces an essential enzyme that plays a key role in the Krebs cycle (Gangloff et al. 1990;

Zhou and Ragan 1995), which is an important process that is required by all aerobic organisms to produce adenosine triphosphate, or energy, which is required to

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power cells. Aconitase catalyzes the reversible isomerization of two tricarboxylic acids, citrate and isocitrate (Chenicek and Hart 1987; Gangloff et al. 1990; Peyret et al. 1995; Zhou and Ragan 1995). Citrate is converted into isocitrate via an intermediate product, cis-aconitate. The enzyme is a single cluster or labile made up of iron and sulfur (Rouault and Klausner 1996). The inactive form of aconitase is 3Fe–4S, which prohibits the enzyme from converting tricarboxylic acids (Zhou and Ragan 1995; Sadka et al. 2000). The active form has a single Fe added that changes the structure to 4Fe–4S and allows the conversion of citrate to isocitrate (Zhou and Ragan 1995; Rouault and Klausner 1996; Sadka et al. 2000). Aconitase protein is one of the few proteins that contains an iron–sulfur cluster but does not contain an electron transport function (Zhou and Ragan 1995).

Aconitases are present in a wide variety of living organisms and are highly conserved (Gangloff et al. 1990). Considerable work has been done on studying the multitude of purposes and uses of aconitase genes. There are two different isoforms of aconitase found in cells: mitochondrial aconitase and cytosolic aconitase. Mitochondrial aconitases are a fundamental part of the Krebs cycle (Sadka et al. 2000; Schnarrenberger and Martin 2002; Moeder et al. 2007; Terol et al. 2010). Although cytosolic aconitases are under-researched as compared with mitochondrial aconitases, they are thought to be involved in numerous processes such as cytosolic citrate metabolism and the glyoxylate cycle (Hayashi et al. 1995; Eprintsev et al. 2015). Mitochondrial and cytosolic aconitases in plants have similar kinetic properties and are believed to be encoded by the same genes (Carrari et al. 2003; Moeder et al. 2007).

Isozymes are enzymes that differ in sequences but catalyze the same chemical reaction. Aconitase isozymes are codominant markers and have been employed in mapping studies in soybean (Griffin and Palmer 1987; Espinosa et al. 2015). Several aconitase genes have been isolated and characterized in various plant species (Peyret et al. 1995; Zhou and Ragan 1995; Sadka et al. 2000; Carrari et al. 2003; Moeder et al. 2007). The knock-out mutants for the three aconitase genes, *At4g35830*, *At4g26970*, and *At2g05710*, in *Arabidopsis thaliana* (L.) Heynh. did not show a morphological phenotype, but were more tolerant to oxidative stress (Moeder et al. 2007). The reduction in expression of *Aco1* in *Solanum pennellii* Correll. caused several noticeable alterations, such as the darker pigmentation of leaves due to an increase in chlorophyll content, increase in total starch and starch accumulation, less root matter, higher sucrose content, and higher rate of electron transport (Carrari et al. 2003).

Five different aconitase genes have been identified in soybean and are named as *Aco1*, *Aco2*, *Aco3*, *Aco4*, and *Aco5* (Doong and Kiang 1987; Griffin and Palmer 1987; Rennie et al. 1987; Kiang and Bult 1991; Espinosa et al. 2015). Inheritance studies showed that they assort independently of each other (Griffin and Palmer 1987;

Rennie et al. 1987; Kiang and Bult 1991). Of these five genes, *Aco3* and *Aco4* have been mapped to soybean chromosomes 6 and 11, respectively (Griffin and Palmer 1987; Espinosa et al. 2015). *Aco4* was flanked by two simple sequence repeat (SSR) markers, BARCSOYSSR_11_323 and BARCSOYSSR_11_336, to a ~292 kb region (Espinosa et al. 2015), which made it an excellent candidate for map-based cloning. The objectives of this research were to fine map the chromosomal region containing *Aco4* and to identify and sequence the candidate gene for *Aco4*.

Materials and Methods

Plant material

Two diverse parents, BSR 101 (PI 548519; *Aco4-aa*) and Noir I (PI 290136; *Aco4-bb*), that differ in their aconitase pattern were crossed to generate a mapping population. The F₁ plants were individually threshed and the F₂ seeds were planted at the Bruner Farm near Ames, IA. The F₂ mapping population consisted of 94 individuals. Samples were taken from the individual F₂ plants for aconitase activity and for DNA analysis.

Aconitase isozyme analysis

Aconitase activity [aconitate hydratase, enzyme commission (EC) 4.2.1.3] was determined as previously described (Espinosa et al. 2015). Isozyme patterns at the *Aco4* locus were determined for the parents and the F₂ population using starch gel electrophoresis as previously described (Espinosa et al. 2015). After electrophoresis, gels were sliced horizontally into 1.5 mm thick pieces to allow analysis of several isozymes from one gel. For visualization of the aconitase activity, the gel slices were incubated at 37 °C in 100 mL 0.2 mol L⁻¹ tris(hydroxymethyl)aminomethane (Tris) – HCl (pH 8.0), 200 mg cis-aconitic acid, 100 mg MgCl₂, 20 mg β-nicotinamide adenine dinucleotide phosphate, 20 mg methyl thiazolyl tetrazolium bromide, 4 mg phenazine methosulfate, and 40 units isocitrate dehydrogenase.

Molecular mapping

In a previous study, the *Aco4* gene was located on chromosome 11 (Espinosa et al. 2015). Twelve SSR markers from chromosome 11 were used on the parents to test for polymorphism. SSR primers were synthesized based on the sequence information from <http://soybase.org/resources/ssr.php> (Song et al. 2004, 2010). For the polymerase chain reactions (PCR), 50 ng DNA was used in 10 µL reaction containing 1× reaction buffer (10 mol L⁻¹ Tris–HCl, 50 mol L⁻¹ KCl, pH 8.3), 0.25 µM of each primer, 200 µM of each deoxynucleotide, 2.0 mol L⁻¹ MgCl₂, and 0.25 units of Biolase DNA polymerase enzyme (Bioline USA, Inc., Tauton, MA). Thermocycler conditions were as follows: 94 °C for 3 min, 35 cycles of 94 °C for 30 s, 46 °C for 30 s, and 72 °C for 1 min, and a final cycle of 72 °C for 10 min. The PCR products were visualized on 4% agarose gel containing ethidium bromide.

Fig. 1. Genetic linkage map and sequence-based physical map of soybean chromosome 11 showing locations of SSR markers close to the *Aco4* gene. Genetic distances are shown in centiMorgans (cM) and physical distances are shown in base pairs (bp). [Colour online.]

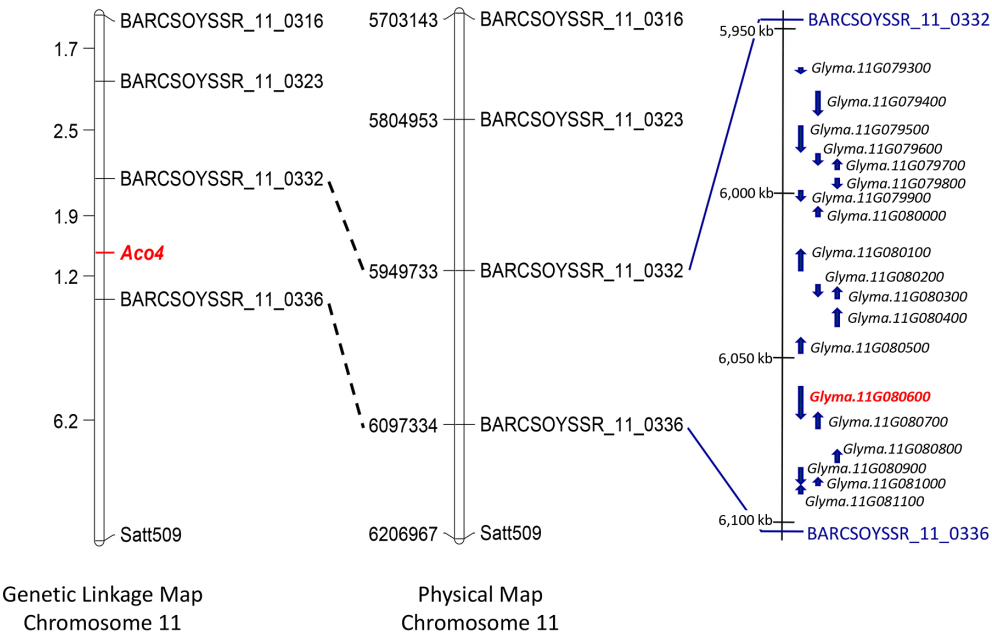


Table 1. Predicted genes present in the *Aco4* region.

Gene	Start position	End position	Predicted protein; function
<i>Glyma.11G079300</i>	5960613	5960978	Copper chaperone; metal ion transport
<i>Glyma.11G079400</i>	5968738	5974185	GDSL esterase/lipase
<i>Glyma.11G079500</i>	5979267	5985576	PLAC8 family
<i>Glyma.11G079600</i>	5987638	5989759	SCF ubiquitin ligase, Skp1
<i>Glyma.11G079700</i>	5991289	5992851	PPR repeat family
<i>Glyma.11G079800</i>	5995314	5996613	Trm112p-like protein
<i>Glyma.11G079900</i>	5999169	6000451	Ring finger domain containing
<i>Glyma.11G080000</i>	6005401	6006947	ATBZIP transcription factor related
<i>Glyma.11G080100</i>	6018320	6023418	Lupus la protein related
<i>Glyma.11G080200</i>	6027758	6029576	Putative methyltransferase
<i>Glyma.11G080300</i>	6030076	6031938	Peroxidase; heme binding
<i>Glyma.11G080400</i>	6036330	6040326	Aspartyl protease; proteolysis
<i>Glyma.11G080500</i>	6045490	6048516	Ring finger protein
<i>Glyma.11G080600</i>	6058457	6066818	Aconitate hydratase, aconitase
<i>Glyma.11G080700</i>	6068106	6071401	Prolyl 4-hydroxylase; oxidoreductase activity
<i>Glyma.11G080800</i>	6079536	6081820	None
<i>Glyma.11G080900</i>	6083715	6086922	Peroxidase; heme binding
<i>Glyma.11G081000</i>	6087915	6088796	None
<i>Glyma.11G081100</i>	6090248	6091010	NADH dehydrogenase; ATP synthesis coupled electron transport

Note: Name and predicted functions of the putative proteins encoded by 19 genes flanked by BARCSOYSSR_11_332 and BARCSOYSSR_11_336 are shown. The candidate gene of interest is shown in bold font.

Polymorphic markers were tested on the entire F₂ population and genetic distances were calculated. The mapping software Mapmaker 2.0 was used to determine genetic linkage, utilizing the Kosambi mapping function (Kosambi 1944; Lander et al. 1987). A logarithm of the odds threshold of 3.0 was used to determine order of

markers. Maps were drawn using the “MapChart” software (Voorrips 2002).

Long-range PCR

A long-range PCR approach was used to capture the entire gene for sequencing. Five forward and 4 reverse

Fig. 2. The amino acid sequence comparison of *Glyma.11G080600* between the A-line (BSR 101) and B-line (Noir I). The differences are due to two non-synonymous substitutions between A-line and B-line. The red font represents non-synonymous substitutions and the green font represents synonymous substitutions. [Colour online.]

A-Line	MATENPFNSILTTLEKPGGAGEFGKYFSLPALNDRRIDRLPYSVRILLESAIRNCDEFQV	60
B-Line	MATENPFNSILTTLEKPGGAGEFGKYFSLPALNDRRIDRLPYSVRILLESAIRNCDEFQV	60
A-Line	KSNDVEKIIDWENTSPKLGEIPFKPARVLLQDFTGVPVAVDLACMRDAMNKLGGDSNKIN	120
B-Line	KSNDVEKIIDWENTSPKLVEIPFKPARVLLQDFTGVPVAVDLACMRDAMNKLGGDSNKIN	120
A-Line	PLVPVDLVIDHSVQVDVARSENNAVQANMELEFQNRKERFGFLKWSNAFNMMVLVPPGSG	180
B-Line	PLVPVDLVIDHSVQVDVARSENNAVQANMELEFQNRKERFGFLKWSNAFNMMVLVPPGSG	180
A-Line	IVHQVNLEYLGRVVFNTNGVLYPDSVVGTDSTHTMIDGLGVAGWGVGGIEAEAAAMLGQPM	240
B-Line	IVHQVNLEYLGRVVFNTNGVLYPDSVVGTDSTHTMIDGLGVAGWGVGGIEAEAAAMLGQPM	240
A-Line	SMVLPGVGVGFKLLGKLRDGVATDVLVLTQMLRKHGTVGKVFVEFYEGMSELSLADRAT	300
B-Line	SMVLPGVGVGFKLLGKLRDGVATDVLVLTQMLRKHGTVGKVFVEFYEGMSELSLADRAT	300
A-Line	IANMSPEYGATMGFFPVDHVTQLRLTGRSDETVMIESYLANKMFVDYSEPVQERVY	360
B-Line	IANMSPEYGATMGFFPVDHVTQLRLTGRSDETVMIESYLANKMFVDYSEPVQERVY	360
A-Line	SSYLELNLEDVEPCVSGPKRPHDRVPLREMKVDWHACLNNKVGFKGFAVPKESQNKVAEF	420
B-Line	SSYLELNLEDVEPCVSGPKRPHDRVPLREMKVDWHACLNNKVGFKGFAVPKESQNKVAEF	420
A-Line	TFQGTPAHLRHGDVIAAITSTNTSNPSVMLGAALVAKKACELGLQVKPWIKTSLAPGS	480
B-Line	TFQGTPAHLRHGDVIAAITSTNTSNPSVMLGAALVAKKACELGLQVKPWIKTSLAPGS	480
A-Line	GVVTKYLRSGLQKYLNELGFNIVGYGCTTCIGNSGDINEAVASAITENDIVAAVLSGN	540
B-Line	GVVTKYLRSGLQKYLNELGFNIVGYGCTTCIGNSGDINEAVASAITENDIVAAVLSGN	540
A-Line	RNFEGRVHPLTRANYLASPLVAYALAGTVDIDFDTEPIGIGKDGTKIFFRDIWPSSEE	600
B-Line	RNFEGRVHPLTRANYLASPLVAYALSGTVDIDFDTEPIGIGKDGTKIFFRDIWPSSEE	600
A-Line	IANVVQSSVLPAMFRDTYNAITQGNPMWNNLSVPTGTLYAWDPTSTYIHEPPYFRDMSMS	660
B-Line	IANVVQSSVLPAMFRDTYNAITQGNPMWNNLSVPTGTLYAWDPTSTYIHEPPYFRDMSMS	660
A-Line	PPGSHGVKDAYCLNFGDSITTDHISPAGSIHKDSPAARYLIERGVDRRDFNSYGSRRGN	720
B-Line	PPGSHGVKDAYCLNFGDSITTDHISPAGSIHKDSPAARYLIERGVDRRDFNSYGSRRGN	720
A-Line	DEVMARGTFANIRIVNKFNGEVGPKTIHIPSGEKLSVFDAAEKYKSEGHDMIILAGAEY	780
B-Line	DEVMARGTFANIRIVNKFNGEVGPKTIHIPSGEKLSVFDAAEKYKSEGHDMIILAGAEY	780
A-Line	GSGSSRDWAAKGPMLLGVKAVIAKSFERIHRSNLVGMGIPLCFKPGDDADSLGLTGHER	840
B-Line	GSGSSRDWAAKGPMLLGVKAVIAKSFERIHRSNLVGMGIPLCFKPGDDADSLGLTGHER	840
A-Line	YTIDLPSNVNEIRPGQDVTVVTDAGKSFSVTLRFDEVELAYFNHGGILQYVIRNMVNAK	900
B-Line	YTIDLPSNVNEIRPGQDVTVVTDAGKSFSVTLRFDEVELAYFNHGGILQYVIRNMVNAK	900
A-Line	H	901
B-Line	H	901
	*	

primers were developed (Supplementary Table S1).¹ The forward and reverse long-range primers were designed within 1000 base pairs (bp) upstream and downstream of the gene, respectively. For long-range PCR, a LongAmp™ Taq PCR kit (New England Biolabs, Ipswich, MA) was used.

The PCR program consisted of initial denaturation at 94 °C for 30 s, followed by 40 cycles of 94 °C for 30 s, 53 °C for 30 s, and 65 °C for 7 min, and elongation was set at 65 °C for 10 min. PCR products were separated on 0.8% agarose gel and desired fragment bands were

¹Supplementary material is available with the article through the journal Web site at <http://nrcresearchpress.com/doi/suppl/10.1139/cjps-2016-0363>.

excised. DNA was purified from the gel fragments using an extraction kit (Clontech Laboratories, Inc., Mountain View, CA).

Sequencing

Sequencing was performed using 15 forward and 15 reverse primers along the length of the candidate *Aco4* gene using the long-range product as a template (Supplementary Table S2).¹ PCR conditions used for amplification were as follows: 94 °C for 3 min, 40 cycles of 94 °C for 30 s, 52 °C for 30 s, and 72 °C for 4 min, with final extension at 72 °C for 10 min. Sequencing was performed at Functional Biosciences, Inc. (Madison, WI) using Sanger sequencing. Vector NTI™ Express (Life Technologies Corporation, Carlsbad, CA) was used to align the DNA fragments.

Expression analysis of *Glyma.11G080600*

The soybean RNA sequencing dataset was used to compare the expression of *Glyma.11G080600* during different developmental stages (Severin et al. 2010). Fragments per kilobase of transcript per million mapped reads (FPKM) for different development stages of soybean were compared (<https://phytozome.jgi.doe.gov/pz/portal.html#!gene?search=1&detail=1&method=4433&searchText=transcriptid:30528935>).

Phylogenetic analysis

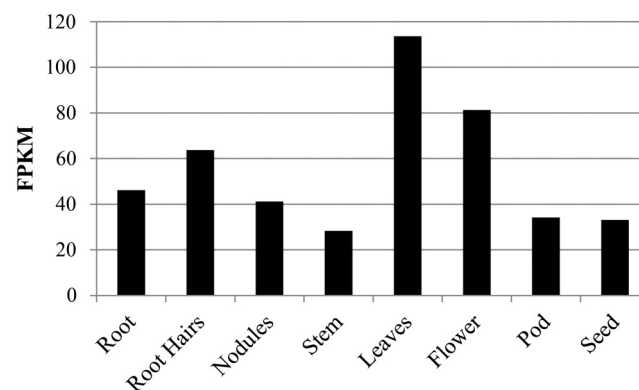
The protein sequence for *Glyma.11G080600* was used in a protein Basic Local Alignment Search Tool (BLAST®) search using the National Center for Biotechnology Information BLAST web service (<http://blast.ncbi.nlm.nih.gov/Blast.cgi>) to find homologous sequences from several plant species. The protein sequences were aligned in SeaView 4.5.4, a phylogenetic alignment software program (Gouy et al. 2010). The phylogenetic relationship between *Aco4* (*Glyma.11G080600*) and 24 homologous proteins was then constructed using the neighbor-joining method. The evolutionary distances were computed using the Poisson correction method and are in units of the number of amino acid substitutions per site.

Results and Discussion

Fine mapping

In a previous study, the *Aco4* gene was mapped on chromosome 11 between SSR markers BARCSOYSSR_11_323 and BARCSOYSSR_11_336 that flanked a 292 kb region (Espinosa et al. 2015). In this investigation, we wanted to narrow down the region, so the putative candidate gene for *Aco4* can be identified. We selected eight SSR markers between BARCSOYSSR_11_323 and BARCSOYSSR_11_336 and tested those for polymorphism between the parents. Only BARCSOYSSR_11_332 showed polymorphism; we then generated a new map including BARCSOYSSR_11_332. The *Aco4* gene mapped between

Fig. 3. RNA sequencing based expression analysis of the *Aco4* gene (*Glyma.11G080600*) in different tissues in soybean (<https://phytozome.jgi.doe.gov/pz/portal.html#!gene?search=1&detail=1&method=4433&searchText=transcriptid:30528935>). FPKM, fragments per kilobase of transcript per million mapped reads.

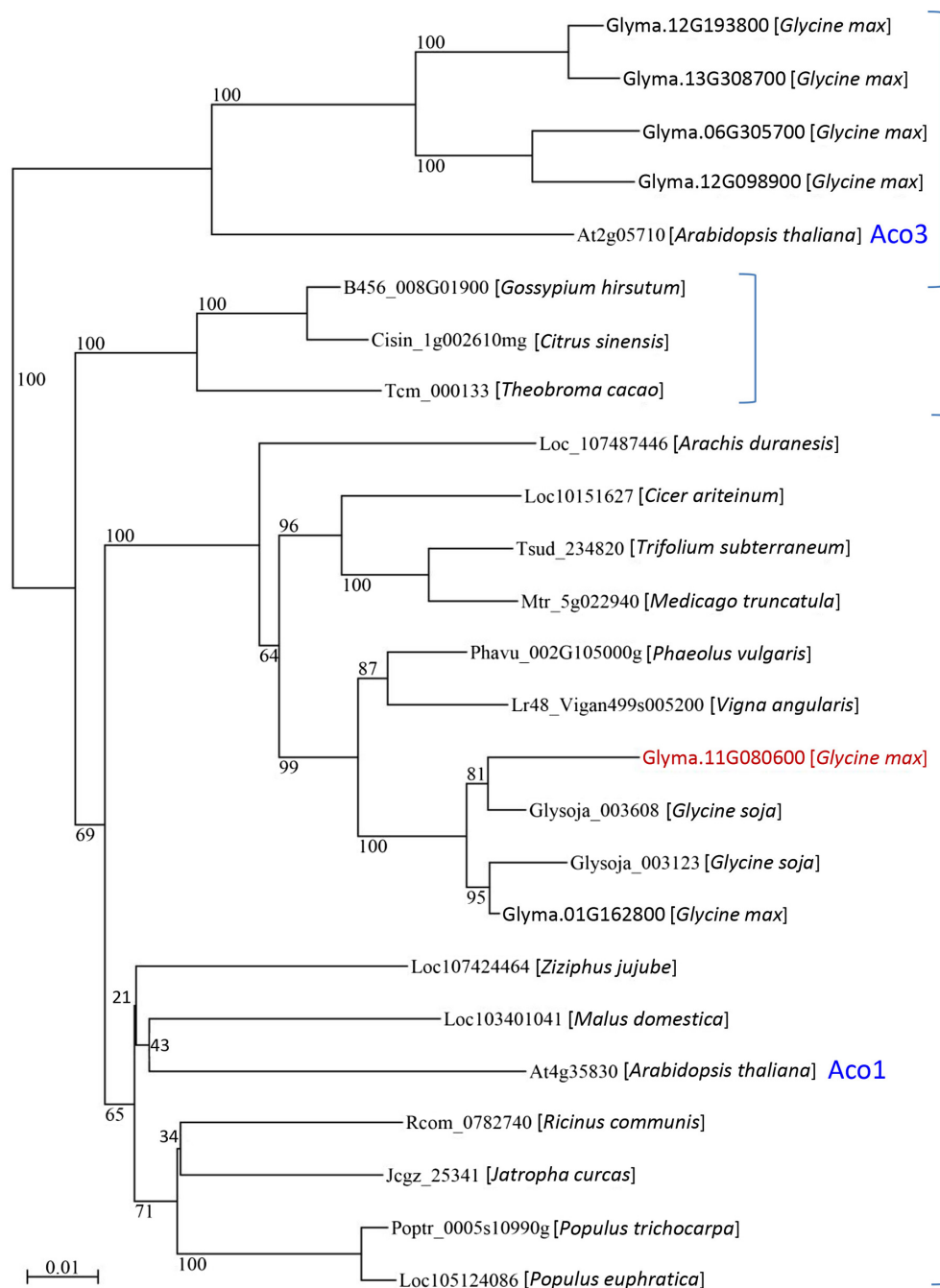


BARCSOYSSR_11_332 and BARCSOYSSR_11_336 (Fig. 1). This region is physically ~148 kb and there are only 19 predicted genes in this region (Table 1). Of these 19 genes, *Glyma.11G080600* is annotated to code for aconitate hydratase (Table 1; www.soybase.org), which makes *Glyma.11G080600* a perfect candidate for *Aco4*.

Isolation of the *Aco4* gene

To confirm if *Glyma.11G080600* is a viable candidate for *Aco4*, we sequenced the *Glyma.11G080600* locus in the parents BSR 101 (*Aco4-aa*; A-line) and Noir I (*Aco4-bb*; B-line) and homozygous F₂ plants *Aco4-aa* (pooled sample from 10 plants) and *Aco4-bb* (pooled sample from 10 plants). The genomic sequence of *Glyma.11G080600* is 8366 bp long with 21 exons and codes for a 901 amino acid-long protein (Supplementary Fig. S1).¹ Sequence comparison between the A-line and B-line showed 12 base substitutions and 7 indel variations in the introns (Supplementary Fig. S2).¹ In the exons, there were only five differences. Of these, three were synonymous substitutions of G with A, C with T, and G with C at the positions 766 (G766A), 1718 (C1718T), and 2750 (G2750C), respectively (Fig. 2; Supplementary Fig. S2).¹ Positions of the substitutions are described based on the sequence of the A-line. The substitutions G766A, C1817T, and G2750C did not affect the amino acids glutamic acid, histidine, and serine, respectively, in the protein sequence. Synonymous substitutions most likely were not the cause of variation in the isozyme pattern in the A-line and B-line. The other two differences between the A-line and B-line were non-synonymous substitutions. The substitution of T with A at position 1197 (T1197A) resulted in a change from valine to glycine and substitution G5424T caused a change from alanine to serine in the protein sequence (Supplementary Fig. S2¹; Fig. 2). The new amino acids create slightly

Fig. 4. Phylogenetic tree of the homologs of *Glyma.11G080600* (*Aco4*). The tree was generated using the neighbor-joining method in the SeaView software (Gouy et al. 2010). Numbers on the nodes represent percent bootstrap support in a 1000-replicate bootstrap test with a total of 838 sites in the final dataset. The tree is drawn to scale, with branch lengths in the same units as those of the evolutionary distances used to infer the phylogenetic tree. [Colour online.]



different proteins in the A- and B-lines, which possibly will have a different structure and lead to a different isozyme pattern.

Glyma.11G080600 contains two major domains including aconitase A catalytic domain (amino acids 87–571) and aconitase A swivel domain (amino acids 675–845) (Supplementary Fig. S1).¹ Aconitase catalytic domain

catalyzes isomerization of citrate to isocitrate and vice versa during the Krebs cycle (Marchler-Bauer et al. 2015). The swivel domain is known to play an important role in swiveling conformational change in the enzyme mechanism (Marchler-Bauer et al. 2015). Conservation of both domains suggests that the *Glyma.11G080600* protein is able to catalyze the interconversion of isocitrate and citrate.

Expression of *Glyma.11G080600* in different soybean tissues

Publicly available RNA sequencing data were used to compare the expression of *Glyma.11G080600* in different soybean tissues (Severin et al. 2010). *Glyma.11G080600* is expressed in most plant tissues with maximum expression in leaves and minimum expression in the stem (Fig. 3). Expression in leaves is more than 4-fold higher than stem, which may be due to active involvement of leaves in metabolism.

Phylogenetic analysis of aconitate hydratase in different plant species

To analyze the evolutionary history of the aconitate hydratase family members in different plant species, the *Glyma.11G080600* protein sequence was used in BLAST analysis to identify homologous proteins in different plant species. Twenty-five different proteins were compared in a multiple alignment and phylogenetic analysis was performed. The phylogenetic tree displayed three major clusters (Fig. 4). The *Aco* family in soybean displays multiple sequences associated with a specific aconitase gene, perhaps due to the paleoploid nature of the soybean genome. Four soybean genes, *Glyma.12G193800*, *Glyma.13G308700*, *Glyma.06G305700*, and *Glyma.12G098900*, cluster together with the *Arabidopsis Aco3* gene, *At2g05710* (Fig. 4). In a previous study, the soybean *Aco3* gene was mapped on chromosome 6 in the vicinity of *Glyma.06G305700* (Griffin and Palmer 1987; Kiang and Bult 1991). It is likely that *Glyma.06G305700* is an ortholog of *Arabidopsis Aco3*. The other three soybean genes may be paralogs of *Glyma.06G305700*, which may have developed due to duplication events. Two soybean genes, *Glyma.11G080600* and *Glyma.01G162800*, fall in the same cluster with *Arabidopsis Aco1*. In this investigation, we have shown that *Glyma.11G080600* is the most suitable candidate for *Aco4*. It seems that the *Aco1* gene has evolved into two closely associated genes, *Glyma.11G080600* (*Aco4*) and *Glyma.01G162800* (*Aco1*), in soybean. The short evolutionary distance between the genes suggests an occurrence of a recent duplication event. Each of these two *Glycine max* (L.) Merr. genes has a closely associated orthologous gene in *G. soja* Siebold. & Zucc., which indicates that the duplication event happened prior to the *G. max* and *G. soja* split. Further research on the characterization of the *Aco* genes in soybean may shed light on the need of several variants in plant development, survival, and reproductive success.

Conclusion

We have mapped the *Aco4* gene to about a 148 kb region on chromosome 11 in soybean. Of the 19 predicted genes present in this region, *Glyma.11G080600* codes for aconitate hydratase. Sequencing of *Glyma.11G080600* in the lines that vary for their aconitase pattern displayed five differences in their exon sequences, of which, only two were non-synonymous

substitutions, which most likely are the cause of different isozyme patterns in the two variants. These observations indicate that *Glyma.11G080600* is the most suitable candidate for *Aco4*. Phylogenetic analysis suggested that soybean *Aco4* and *Aco1* are very closely associated and may have emerged from a single progenitor gene in the recent past.

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References

- Carrari, F., Nunes-Nesi, A., Gibon, Y., Lytovchenko, A., Loureiro, M.E., and Fernie, A.R. 2003. Reduced expression of aconitase results in an enhanced rate of photosynthesis and marked shifts in carbon partitioning in illuminated leaves of wild species tomato. *Plant Physiol.* **133**(3): 1322–1335. doi:[10.1104/pp.103.026716](https://doi.org/10.1104/pp.103.026716). PMID:[14551334](https://pubmed.ncbi.nlm.nih.gov/14551334/).
- Chenicek, K.J., and Hart, G.E. 1987. Identification and chromosomal locations of aconitase gene loci in Triticeae species. *Theor. Appl. Genet.* **74**(2): 261–268. doi:[10.1007/BF00289978](https://doi.org/10.1007/BF00289978). PMID:[24241574](https://pubmed.ncbi.nlm.nih.gov/24241574/).
- Doong, J.Y.H., and Kiang, Y.T. 1987. Inheritance of aconitase isozymes in soybean. *Genome*, **29**(5): 713–717. doi:[10.1139/g87-122](https://doi.org/10.1139/g87-122).
- Eprintsev, A.T., Fedorin, D.N., Nikitina, M.V., and Igamberdiev, A.U. 2015. Expression and properties of the mitochondrial and cytosolic forms of aconitase in maize scutellum. *J. Plant Physiol.* **181**: 14–19. doi:[10.1016/j.jplph.2015.03.012](https://doi.org/10.1016/j.jplph.2015.03.012). PMID:[25933369](https://pubmed.ncbi.nlm.nih.gov/25933369/).
- Espinosa, K., Boelter, J., Lolle, S., Hopkins, M., Goggi, S., Palmer, R.G., and Sandhu, D. 2015. Evaluation of spontaneous generation of allelic variation in soybean in response to sexual hybridization and stress. *Can. J. Plant Sci.* **95**(2): 405–415. doi:[10.4141/cjps-2014-324](https://doi.org/10.4141/cjps-2014-324).
- Gangloff, S.P., Marguet, D., and Lauquin, G.J. 1990. Molecular cloning of the yeast mitochondrial aconitase gene (*AC01*) and evidence of a synergistic regulation of expression by glucose plus glutamate. *Mol. Cell. Biol.* **10**(7): 3551–3561. doi:[10.1128/MCB.10.7.3551](https://doi.org/10.1128/MCB.10.7.3551).
- Gouy, M., Guindon, S., and Gascuel, O. 2010. SeaView version 4: a multiplatform graphical user interface for sequence alignment and phylogenetic tree building. *Mol. Biol. Evol.* **27**(2): 221–224. doi:[10.1093/molbev/msp259](https://doi.org/10.1093/molbev/msp259). PMID:[19854763](https://pubmed.ncbi.nlm.nih.gov/19854763/).
- Griffin, J.D., and Palmer, R.G. 1987. Inheritance and linkage studies with five isozyme loci in soybean. *Crop Sci.* **27**: 885–892. doi:[10.2135/cropsci1987.0011183X0027000500010x](https://doi.org/10.2135/cropsci1987.0011183X0027000500010x).
- Hayashi, M., De Bellis, L., Alpi, A., and Nishimura, M. 1995. Cytosolic aconitase participates in the glyoxylate cycle in etiolated pumpkin cotyledons. *Plant Cell Physiol.* **36**(4): 669–680. PMID:[7640891](https://pubmed.ncbi.nlm.nih.gov/7640891/).
- Kiang, Y.T., and Bult, C.J. 1991. Genetic and linkage analysis of aconitate hydratase variants in soybean. *Crop Sci.* **31**: 322–325. doi:[10.2135/cropsci1991.0011183X003100020020x](https://doi.org/10.2135/cropsci1991.0011183X003100020020x).
- Kosambi, D.D. 1944. The estimation of map distance from recombination values. *Ann. Eugen.* **12**: 172–175. doi:[10.1111/j.1469-1809.1943.tb02321.x](https://doi.org/10.1111/j.1469-1809.1943.tb02321.x).
- Lander, E.S., Green, P., Abrahamson, J., Barlow, A., Daly, M.J., Lincoln, S.E., and Newburg, L. 1987. MAPMAKER: an interactive computer package for constructing primary genetic

- linkage maps of experimental and natural populations. *Genomics*, **1**(2): 174–181. doi:[10.1016/0888-7543\(87\)90010-3](https://doi.org/10.1016/0888-7543(87)90010-3).
- Marchler-Bauer, A., Derbyshire, M.K., Gonzales, N.R., Lu, S., Chitsaz, F., Geer, L.Y., Geer, R.C., He, J., Gwadz, M., Hurwitz, D.L., Lanczycki, C.J., Lu, F., Marchler, G.H., Song, J.S., Thanki, N., Wang, Z., Yamashita, R.A., Zhang, D., Zheng, C., and Bryant, S.H. 2015. CDD: NCBI's conserved domain database. *Nucleic Acids Res.* **43**(D1): D222–D226. doi:[10.1093/nar/gku1221](https://doi.org/10.1093/nar/gku1221). PMID:[25414356](https://pubmed.ncbi.nlm.nih.gov/25414356/).
- Moeder, W., Del Pozo, O., Navarre, D.A., Martin, G.B., and Klessig, D.F. 2007. Aconitase plays a role in regulating resistance to oxidative stress and cell death in *Arabidopsis* and *Nicotiana benthamiana*. *Plant Mol. Biol.* **63**(2): 273–287. doi:[10.1007/s11103-006-9087-x](https://doi.org/10.1007/s11103-006-9087-x).
- Peyret, P., Perez, P., and Alric, M. 1995. Structure, genomic organization, and expression of the *Arabidopsis thaliana* aconitase gene. Plant aconitase show significant homology with mammalian iron-responsive element-binding protein. *J. Biol. Chem.* **270**(14): 8131–8137. doi:[10.1074/jbc.270.14.8131](https://doi.org/10.1074/jbc.270.14.8131).
- Rennie, B.D., Beversdorf, W.D., and Buzzell, R.I. 1987. Genetic and linkage analysis of an aconitate hydratase variant in the soybean. *J. Hered.* **78**(5): 323–326. doi:[10.1093/oxfordjournals.jhered.a110396](https://doi.org/10.1093/oxfordjournals.jhered.a110396).
- Rouault, T.A., and Klausner, R.D. 1996. Iron-sulfur clusters as biosensors of oxidants and iron. *Trends Biochem. Sci.* **21**(5): 174–177. doi:[10.1016/S0968-0004\(96\)10024-4](https://doi.org/10.1016/S0968-0004(96)10024-4).
- Sadka, A., Dahan, E., Cohen, L., and Marsh, K.B. 2000. Aconitase activity and expression during the development of lemon fruit. *Physiol. Plant.* **108**(3): 255–262. doi:[10.1034/j.1399-3054.2000.108003255.x](https://doi.org/10.1034/j.1399-3054.2000.108003255.x).
- Schnarrenberger, C., and Martin, W. 2002. Evolution of the enzymes of the citric acid cycle and the glyoxylate cycle of higher plants. A case study of endosymbiotic gene transfer. *Eur. J. Biochem.* **269**(3): 868–883. doi:[10.1046/j.0014-2956.2001.02722.x](https://doi.org/10.1046/j.0014-2956.2001.02722.x).
- Severin, A.J., Woody, J.L., Bolon, Y.T., Joseph, B., Diers, B.W., Farmer, A.D., Muehlbauer, G.J., Nelson, R.T., Grant, D., Specht, J.E., Graham, M.A., Cannon, S.B., May, G.D., Vance, C.P., and Shoemaker, R.C. 2010. RNA-Seq Atlas of *Glycine max*: a guide to the soybean transcriptome. *BMC Plant Biol.* **10**: 160. doi:[10.1186/1471-2229-10-160](https://doi.org/10.1186/1471-2229-10-160). PMID:[20687943](https://pubmed.ncbi.nlm.nih.gov/20687943/).
- Song, Q., Jia, G., Zhu, Y., Grant, D., Nelson, R.T., Hwang, E.-Y., Hyten, D.L., and Cregan, P.B. 2010. Abundance of SSR motifs and development of candidate polymorphic SSR markers (BARCSOYSSR_1.0) in soybean. *Crop Sci.* **50**(5): 1950–1960. doi:[10.2135/cropsci2009.10.0607](https://doi.org/10.2135/cropsci2009.10.0607).
- Song, Q.J., Marek, L.F., Shoemaker, R.C., Lark, K.G., Concibido, V.C., Delannay, X., Specht, J.E., and Cregan, P.B. 2004. A new integrated genetic linkage map of the soybean. *Theor. Appl. Genet.* **109**(1): 122–128. doi:[10.1007/s00122-004-1602-3](https://doi.org/10.1007/s00122-004-1602-3).
- Terol, J., Soler, G., Talon, M., and Cercos, M. 2010. The aconitate hydratase family from *Citrus*. *BMC Plant Biol.* **10**: 222. doi:[10.1186/1471-2229-10-222](https://doi.org/10.1186/1471-2229-10-222). PMID:[20958971](https://pubmed.ncbi.nlm.nih.gov/20958971/).
- Voorrips, R.E. 2002. MapChart: software for the graphical presentation of linkage maps and QTLs. *J. Hered.* **93**(1): 77–78. doi:[10.1093/jhered/93.1.77](https://doi.org/10.1093/jhered/93.1.77).
- Zhou, Y.H., and Ragan, M.A. 1995. Characterization of the nuclear gene encoding mitochondrial aconitase in the marine red alga *Gracilaria verrucosa*. *Plant Mol. Biol.* **28**(4): 635–646. doi:[10.1007/BF00021189](https://doi.org/10.1007/BF00021189). PMID:[7647296](https://pubmed.ncbi.nlm.nih.gov/7647296/).