Pyruvate transporter BnaBASS2 impacts seed oil accumulation in *Brassica napus*

Shan Tang1,2, Ning Guo1,2, Qingqing Tang1,2, Fei Peng1,2, Yunhao Liu1,2, Hui Xia1,2, Shaoping Lu1,2 and Liang Guo1,2,3,4,*

1National Key Laboratory of Crop Genetic Improvement, Huazhong Agricultural University, Wuhan, China
2Hubei Hongshan Laboratory, Wuhan, China
3Shenzhen Institute of Nutrition and Health, Huazhong Agricultural University, Wuhan, China
4Shenzhen Branch, Guangdong Laboratory for Lingnan Modern Agriculture, Genome Analysis Laboratory of the Ministry of Agriculture, Agricultural Genomics Institute at Shenzhen, Chinese Academy of Agricultural Sciences, Shenzhen, China

Summary

Bile acid: sodium symporter family protein 2 (BASS2) is a sodium-dependent pyruvate transporter, which transports pyruvate from cytosol into plastid in plants. In this study, we investigated the function of chloroplast envelope membrane-localized BnaBASS2 in seed metabolism and seed oil accumulation of *Brassica napus* (*B. napus*). Four BASS2 genes were identified in the genome of *B. napus*. BnaA05.BASS2 was overexpressed while BnaA05.BASS2 and BnaC04.BASS2-1 were mutated by CRISPR in *B. napus*. Metabolite analysis revealed that the manipulation of BnaBASS2 caused significant changes in glycolysis-, fatty acid synthesis–, and energy-related metabolites in the chloroplasts of 31 day-after-flowering (DAF) seeds. The analysis of fatty acids and lipids in developing seeds showed that BnaBASS2 could affect lipid metabolism and oil accumulation in developing seeds. Moreover, the overexpression (OE) of BnaA05.BASS2 could promote the expression level of multiple genes involved in the synthesis of oil and the formation of oil body during seed development. Disruption of BnaA05.BASS2 and BnaC04.BASS2-1 resulted in decreasing the seed oil content (SOC) by 2.8%–5.0%, while OE of BnaA05.BASS2 significantly promoted the SOC by 1.4%–3.4%. Together, our results suggest that BnaBASS2 is a potential target gene for breeding *B. napus* with high SOC.

Keywords: *Brassica napus*, pyruvate transporter, BASS2, seed oil content, lipids, metabolites.

Introduction

Pyruvate is a central intermediate of carbohydrate metabolism in plants. During seed germination of oil plants, pyruvate is an important intermediate metabolite involved in the conversion of lipids to sugars through gluconeogenesis and β-oxidation (Sattler *et al.*, 2004). Moreover, pyruvate is the output of glycolysis of glucose. On one hand, pyruvate produced from glycolysis is transported across the mitochondrial membrane by the mitochondrial pyruvate carrier (MPC; Bricker *et al.*, 2012; Herzig *et al.*, 2012). Under the catalysis of mitochondrial pyruvate dehydrogenase, pyruvate is converted into acetyl-CoA, which further participates in the tricarboxylic acid cycle, connecting the metabolism of sugars, proteins, and lipids (Le *et al.*, 2021; Tovar-Mendez *et al.*, 2003). On the other hand, pyruvate produced from glycolysis is involved in cellular metabolic flux in the plastid. Under the catalysis of plastidal pyruvate dehydrogenase, pyruvate is converted into acetyl-CoA, which acts as a metabolic precursor for many plastid-localized biosynthetic pathways, such as methyl erythritol phosphate (MEP) pathway, branched-chain amino acids and fatty acid (FA) pathway (Furumoto *et al.*, 2011; Prabhakar *et al.*, 2010; Weber and Linka, 2011). Thus, pyruvate is a vital intermediate connecting sugar, lipid, and amino acid metabolism of plants.

In plants, there are two pathways for supplying pyruvate to the plastid. The first way is to catalyse phosphoenolpyruvate (PEP) to produce pyruvate by plastidic pyruvate kinase (PKp) in the plastid (Andre *et al.*, 2007; Baud *et al.*, 2007b; Prabhakar *et al.*, 2010). Another way is to convert PEP into pyruvate by cytosolic pyruvate kinase (PKc) in the cytosol. Cytosolic pyruvate is then transported into the plastid via the pyruvate transporter (Eastmond and Rawsthorne, 2000; Prabhakar *et al.*, 2010; Weber and Linka, 2011). Plastidial pyruvate plays an important role in FA and other metabolic pathways during seed development. In Arabidopsis, PKp consists of three subunits α, β1, and β2. Disruption of PKp resulted in defects in seed oil accumulation, FA synthesis, embryo elongation, and seed germination (Andre *et al.*, 2007; Andre and Benning, 2007; Baud *et al.*, 2007b). In rice, OsPKp consists of four subunits α1, α2, β1 and β2, mutation of OsPKpα1 would affect endosperm starch/FA biosynthesis, granule formation, grain filling, and seed germination (Cai *et al.*, 2018a,b). In *Brassica napus* (*B. napus*), plastids were isolated from embryos of 23, 32, and 41 day-after-anthesis (DAA). Metabolic fluxes for oil synthesis were measured and the activity of the plastidial pyruvate transporter was increased by 19-fold and the utilization of plastidial pyruvate for FA synthesis was increased by 25-fold at 41 DAA (Eastmond and Rawsthorne, 2000). Thus, the above advances reveal that plastidial pyruvate plays an important role in seed development, especially in oil accumulation.

Bile acid: sodium symporter family protein 2 (BASS2) is a chloroplast envelope membrane-localized sodium-dependent pyruvate transporter that transports pyruvate from cytosol to plastid. This process is involved in the photosynthesis of C4 plants
and the MEP pathway of C3 plants (Furumoto, 2016; Furumoto et al., 2011). In Arabidopsis, disruption of BASS2 resulted in mevastatin-sensitive phenotype, and the phenotype could be recovered by constitutive expression of BASS2 in wheat and Arabidopsis, indicating that BASS2 functions in supplying pyruvate to the MEP pathway (Furumoto et al., 2011; Zhao et al., 2016). In addition to the role in the MEP pathway, supplying pyruvate to plastid from cytosol via BASS2 is reported to impact FA biosynthesis and seed oil accumulation. Seed-specific overexpression (OE) of BASS2 resulted in 10%–37% more oil in Arabidopsis seeds (Lee et al., 2017). Nevertheless, beyond the confirmation that BASS2 affects the phenotype of Arabidopsis seed oil accumulation, BASS2’s impact on pyruvate content in plastid and seed metabolism has not been revealed.

In this study, four plastidial BASS2 proteins were identified in B. napus. The gene function and molecular mechanism of BnaBASS2 on seed oil accumulation were investigated using OE and CRISPR mutants of BnaBASS2. These results suggest that BnaBASS2 has a significant impact on seed metabolism and seed oil accumulation in B. napus.

**Results**

**Generation of BnaBASS2 OE and CRISPR mutant lines**

To investigate the function of BASS2s in B. napus, the amino acid sequences of Arabidopsis BASS2 were used for BLAST analysis in BnTIR database (http://yanglab.hzau.edu.cn/BnTIR; Liu et al., 2021). The results show that there are four homologous copies of BASS2 (BnaA05G06697002S, designated as BnaA05.-; BnaBASS2; BnaC04G00719002S, designated as BnaC04.BASS2-1; BnaA04G042485002S, designated as BnaA04.BASS2; and BnaC04G05637002S, designated as BnaC04.BASS2-2), and they all exhibited more than 85% homology with AtBASS2. Among the four genes, BnaA05.BASS2 is the most homologous to BnaC04.BASS2-1 with a homology of 98%. BnaA04.BASS2 is the most homologous to BnaC04.BASS2-2 with a homology of 97% (Figures 1a and S1). Next, the expression level of BnaBASS2 in different tissues was collected from the BnTIR database (Liu et al., 2021). The results show that the expression level of BnaA04.BASS2 and BnaC04.BASS2-2 is very low in different tissues, while BnaA05.BASS2 and BnaC04.BASS2-1 have higher expression in multiple tissues, especially in developing siliques and seeds (Figure 1a).

BASS2 was predicted to be a membrane protein with nine transmembrane regions (Figure S1). In this research, BnaA05.-BASS2 fused with GFP at its C-terminus was transiently expressed in the epidermal cells of tobacco and protoplasts of Arabidopsis. The green fluorescence signal is found to encapsulate the autofluorescence of chloroplast membrane, suggesting that BnaA05.-BASS2 is localized on the chloroplast membrane (Figures 1b and S2). The result was consistent with previous studies (Furumoto et al., 2011).

BnaA05.BASS2, having a much higher expression level than the other three homologous genes in developing siliques and seed, was overexpressed in B. napus driven by 35S promoter. Four transgenic lines with increased expression of BnaA05.-BASS2 in the seeds at 24 day-after-flowering (DAF) are successfully obtained (Figure 1c). Western blot using Flag antibody confirms that positive OE plants are acquired (Figure 1d). OE2 and OE5 having high expression at transcription and protein levels were used for further analysis in this study. CRISPR/Cas9 was utilized to create mutants of BnaBASS2s. As a result, two homozygous mutants CR253 and CR313 with mutations in BnaA05.BASS2 and BnaC04.BASS2-1 are identified by sequencing and the mutations in BnaA05.BASS2 and BnaC04.BASS2-1 caused premature termination (Figures 1e and S3).

**BnaBASS2 has significant effects on seed metabolism**

It has been hypothesized and partially verified that pyruvate produced by glycolysis could be transported into the chloroplast via BASS2 (Furumoto et al., 2011; Schwender et al., 2004; Weber and Linka, 2011). A brief model of carbon influx from the glycolysis pathway into FA synthesis is shown in Figure 2a. To explore the impact of pyruvate transporter BASS2 on seed metabolism of B. napus, metabolites were extracted from 31 DAF seeds and determined by liquid chromatography with tandem mass spectrometry (LC–MS/MS) method. The results show that significant changes occur in most glycolytic and FA synthesis metabolites in the seeds of OE and mutant lines (Figure 2b). In the whole seeds, the contents of the metabolites of glucose-6-phosphate (G6P), fructose-6-phosphate (F6P), fructose-1,6-diphosphate (FBP), dihydroxy-acetone-phosphate (DHAP), 3-phosphoglycerate (3-PGA), PEP and pyruvate (Pyr) related to glycolysis are significantly lower in CR253 than that in WT. However, most of them are higher in OE5 than that in WT, except that the content of G6P, F6P, and 3-PGA have no difference between OE5 and WT. Acetyl-CoA and malonyl-CoA are the precursors for FA synthesis. Compared with WT, the contents of acetyl-CoA and malonyl-CoA are lower in CR253 than that in WT. Nevertheless, the content of acetyl-CoA is higher, but there has been no change in malonyl-CoA in OE5 (Figure 2b). Additionally, intact chloroplasts were separated from 31 DAF seeds by the sucrose gradient method (Figure S4). In chloroplast, the contents of the metabolites of G6P, F6P, FBP, DHAP, 3-PGA, PEP, and Pyr related to glycolysis, and acetyl-CoA and malonyl-CoA related to FA synthesis are significantly lower in CR253 than that in WT. Whereas, the level of those metabolites are higher in OE5 except G6P and PEP (Figure 2c).

The synthesis of seed oil is an energetically costly process, which requires ATP, NADH, and NADPH for carbon flux. Glycolysis is a significant source of ATP, NADH, and NADPH. In chloroplast, light energy can be utilized for ATP and NADPH synthesis, which is required by FA synthesis (Judge and Dodd, 2020; Pfeil et al., 2014; Rawsthorne, 2002). The levels of ATP, ADP, NAD(P), and NAD(P)H in whole seeds and isolated chloroplasts of 31 DAF seeds were determined by LC–MS/MS method. In whole seeds, the contents of ATP, NAD, and NADPH are significantly increased in OE5 but decreased in CR253 compared with WT. However, the content of ADP, NAD and NADPH are significantly decreased in OE5 but elevated in CR253 except no significant change of NADP in CR253 compared with WT (Figure 3a–c). In chloroplast, compared with WT, the contents of ATP, NADH, and NADPH are significantly increased in OE5 but decreased in CR253 except no significant change of NADP in CR253 compared with WT (Figure 3a–c). In chloroplast, compared with WT, the contents of ATP, NADH, and NADPH are significantly increased in OE5 but decreased in CR253 except no significant change of ATP and NADH in CR253. The contents of ADP, NAD, and NADPH are significantly decreased in OE5 but elevated in CR253. However, there is no significant change in NADP in OE5 and CR253 (Figure 3d–f).

The ratios of energy metabolites were used to evaluate the energy charge and reducing power in WT, OE, and mutant lines. In whole seeds, ATP/ADP, NADH/NAD, and NADPH:NAD are
much higher in OE5 but lower in CR253, except for NADPH/NADP in CR253, which is comparable with WT. In chloroplast, ATP/ADP, NADH/NAD, and NADPH/NADP are also much higher in OE5 but lower in CR253, except for NADH/NAD in CR253, which is comparable with WT. Interestingly, the ATP/ADP, NADH/NAD, and NADPH/NADP in the whole seeds of OE5 are changed from 3.8, 2.6, and 2.1 to 16.9, 0.03, and 4.3 in the chloroplast of OE5. ATP/ADP, NADH/NAD, and NADPH/NADP in the whole seeds of CR253 are changed from 0.8, 0.4, and 1.1 to 0.6, 0.008, and 0.5 in the chloroplast of CR253, respectively (Figure 3a–f). The above results indicate that BnaBASS2 affects multiple metabolic pathways including glycolysis, FA synthesis, and energy metabolism in the seeds.

**BnaBASS2** promotes oil accumulation during seed development

Developing seeds were collected and FAs at 21, 27, 31, 36, 41, and 55 DAF were extracted and quantified by the gas chromatography-flame ionization detector (GC-FID) method. Compared with WT, OE5 has a higher level of oil, while the mutant CR253 exhibits a lower level in the process of seed oil accumulation (Figure 4a). The data of seed weight and oil per seed were obtained by calculating the total number of seeds and seed weight. The results revealed that compared with WT, OE5’s seed weight has no significant difference, but the mutant CR253 shows a decreased weight, especially at the seed maturation stage (Figure 4b).

Lipidome analysis of 31 DAF seeds showed that triacylglycerol (TAG) is increased by 57.1% in OE5 but decreased by 51.1% in CR253 compared with WT. The content of DAG is increased by 18.0% in OE5 while has no obvious difference in CR253 compared with WT. Moreover, compared with WT, the content of other lipids MGDG, DGDG, PC, PE, and PA has no obvious difference between WT, OE5, and CR253 (Figure 4c). The content of TAG species -52:2, -52:3, -52:5, -54:2 to -54:7, -56:2, -56:3, and -56:4 is increased in OE5, but the content of TAG-52:2, -52:3, -52:5, -54:2 to -54:7, -56:2, -56:3, and -56:4 is decreased in CR253 (Figure 4d). Compared with WT, the content of DAG species -34:0, -34:2, -34:4, -36:1, -36:3, -36:4, -36:5, and -36:6 is increased in OE5, but the content of DAG-36:3 is decreased in CR253 (Figure 5).

Furthermore, the expression of some key genes related to FA synthesis, TAG synthesis, and oil body (OB) formation was analysed in 31 DAF seeds of WT, OE5, and CR253. The results showed that the expression of *WRINKLED1* (WRI1), a master regulator in the glycolytic pathway, and FA biosynthesis (Adhikari et al., 2016, Baud et al., 2009, Kong and Ma, 2018, Kong et al., 2019, To et al., 2012, Yang et al., 2022), is elevated by 25.8% in OE5 but reduced by 31.8% in CR253 compared with WT. Compared with WT, the expression of FA synthesis-related genes malonyltransferase (MCMT), enoyl-ACP reductase (ENR), and acyl-ACP thioesterase A (FATA) is decreased by 34.3%, 68.2%, and 28.9% in CR253. In OE5, MCMT and FATA are increased by 22.9% and 30.8%, but ENR shows no change. The
expression of genes related to TAG synthesis such as glycerol-3-phosphate dehydrogenase (GPDH), glycerol-3-phosphate acyltransferase 9 (GPAT9), lysophosphaticid acid acyltransferase 1 (LPAAT1), lysophosphaticid acid acyltransferase 2 (LPAAT2), diacylglycerol acyltransferase 1 (DGAT1), diacylglycerol acyltransferase 2 (DGAT2), lysophosphatidylcholine acyltransferase (LPCAT) and CDP-choline:diacylglycerol cholinephosphotransferase (CPT) is lower in CR253 than that in WT, while they have higher expression in OE5. Moreover, the expression level of genes related to OB formation such as oil body oleosin 1 (OBO1), oil body oleosin 2 (OBO2), oil body oleosin 3 (OBO3), and caleosin (CAO) is also determined. The data show that all of them had lower expression in CR253. In detail, OBO1, OBO2, OBO3, and CAO are decreased by 82.0%, 94.9%, 76.1%, and 85.1% in CR253 than that in WT, while they have increased by 22.9%, 91.6%, and 46.2% in OE5 than that in WT except that there is no difference in OBO2 (Figure 4e). The expression level of several genes in the glycolytic pathway, including hexokinase 1 (HK1), phosphofructokinase 1 (PFK1), plastidic pyruvate kinase (pPK1 and pPK3), and cytosolic pyruvate kinase (cPK1 and cPK2), which control rate-limiting steps of the glycolytic pathway, were determined (Figure S6). The results show that the expression levels of all of the above genes are significantly increased in BnaBASS2 mutant CR253. However, except for HKK, the expression of other genes is down-regulated in the OE5. Thus, the above results suggest that BnaBASS2 promotes oil accumulation and expression level of genes involved in seed oil synthesis during seed development.

BnaBASS2 impacts seed oil content in mature seeds

Seed oil content-related traits of mature seeds were also investigated. Three-year (Year 2019, 2020, and 2021) of near-
infrared spectrometer (NIRS) data display that SOC of OE is increased by 1.4%–3.4%, while SOC of two CRISPR mutants is reduced by 2.8%–5.0% compared with WT (Figure 5a). To confirm the results of NIRS, FAs in mature seeds were analysed by GC-FID. Compared with WT, SOC of OE2 and OE5 is elevated by 2.1% and 4.7%, while CR253 and CR313 are reduced by 4.3% and 6.7% (Figure 5b). In addition, the FA composition shows that the content of palmitic acid (C16:0), stearic acid (C18:0), and linoleic acid (C18:2) is slightly reduced, and oleic acid (C18:1) is slightly increased in OE, while C18:2 is slightly increased and C20:1 is reduced in the mutants (Figure S7).

TAG is the predominant lipid in mature seeds. The content of TAG is increased by 12.2% in OE5 and reduced by 11.4% in CR253, compared with WT (Figure 5c). Compared with WT, the content of TAG species -52:3, -52:4, -54:5 to -54:7, -56:2 to -56:5 is increased in OE5, but the content of TAG-52:2 to -52:3 and -54:2 to -54:5 is decreased in CR253 (Figure 5d).

Oil bodies are the place for TAG storage. Compared with WT, the density of OBs is obviously less in mutant CR253, while OE5 exhibits the opposite trend. The area of protein bodies is larger in CR253 but smaller in OE5 (Figure S7).

Moreover, other seed-related traits such as protein, soluble sugar, and starch content were measured. Compared with WT, the protein content in OE2 and OE5 is noticeably decreased by 1.7% and 1.8%, while it is increased by 2.7% and 2.2% in CR253 and CR313 (Figure S8a). The soluble sugar is decreased in OE lines but increased in mutants (Figure S8b). The starch content is decreased in mutants, but there is no change in OE lines (Figure S8c). Taken together, the above results suggest that BnaBASS2 is important for seed oil accumulation in B. napus.

The effects of BnaBASS2 on plant growth

The seeds of WT, OE, and mutant lines were sown in the pots in the growth chamber and their growth phenotypes at 20, 40, and 80 days were observed. It shows that both OE and mutant lines are grown normally, and no significant growth phenotypic differences are observed (Figure S9a–c). To further evaluate the growth, several growth-related traits were measured for 40-day-old plants. The content of chlorophyll a (Chla), chlorophyll b (Chlb), and carotenoid (Car) is comparable between WT, OE, and mutant lines (Figure S10a). Compared with WT, the leaves of OE lines have a higher level of soluble sugars, whereas the mutants CR253 has more OBs. The total area of OBs is 100 µm² and the size per OB are larger in OE5 but smaller in CR253 (Figure 5f–h).

Moreover, other seed-related traits such as protein, soluble sugar, and starch content were measured. Compared with WT, the protein content in OE2 and OE5 is noticeably decreased by 1.7% and 1.8%, while it is increased by 2.7% and 2.2% in CR253 and CR313 (Figure S8a). The soluble sugar is decreased in OE lines but increased in mutants (Figure S8b). The starch content is decreased in mutants, but there is no change in OE lines (Figure S8c). Taken together, the above results suggest that BnaBASS2 is important for seed oil accumulation in B. napus.

**Figure 3** Metabolic impact of BnaBASS2 on ATP/ADP, NADH/NAD, and NADPH/NADP in developing seeds. (a) ATP content, ADP content, and ATP/ADP in 31 DAF seeds. (b) NADH content, NAD content, and NADH/NAD in 31 DAF seeds. (c) NADPH content, NADP content, and NADPH/NADP in 31 DAF seeds. (d) ATP content, ADP content, and ATP/ADP in separated chloroplasts of 31 DAF seeds. (e) NADH content, NAD content, and NADH/NAD in separated chloroplasts of 31 DAF seeds. (f) NADPH content, NADP content, and NADPH/NADP in separated chloroplasts of 31 DAF seeds. Values are mean ± SD (n = 6–8). Different letters represent significant differences at P < 0.05, based on an ANOVA analysis with the Fisher LSD test.
show the opposite trends (Figure S10b). Nevertheless, there is no difference in the starch content in the leaves among WT, OE, and mutant lines (Figure S10c). The biomass determination shows that there is no significant difference in fresh and dry weight among WT, OE, and mutant lines (Figure S10d). In addition, the photosynthetic efficiency was monitored. The results show that OE lines have a higher photosynthetic rate, transpiration rate, stomatal conductance, and intercellular CO2 concentration than that in WT. However, mutant lines show the opposite (Figure S10e–h). These results suggest that BnaBASS2 impacts sugar accumulation and plant photosynthesis in the leaves.

The impact of BnaBASS2 on agronomic traits of B. napus was also evaluated. Plants were sown in the field and harvested at the mature stage. The agronomic traits including plant height, thousand seed weight and plant yield, etc. were investigated (Figure S11, Table S1). The results show that the value of most of the agronomic traits has no significant change between OE lines and WT, except a slight decrease in the number of effective branches in OE2. Most of the agronomic traits are significantly lower in mutants than that in WT, except no obvious difference in plant height and number of effective branches. In detail, the silique number per plant, seed number per silique, thousand seed weight, and yield per plant of the mutants are reduced by 32.7%, 17.0%, 7.8%, and 26.8%, respectively. These results suggest that BnaBASS2 is necessary for plant growth, and disruption of BnaBASS2 can reduce rapeseed yields.

**Discussion**

The function of BASS2 is transporting pyruvate from cytosol to the plastid (Furumoto et al., 2011). It’s reported that the plastidial pyruvate is used for the MEP pathway and FA
biosynthesis (Furumoto et al., 2011; Prabhakar et al., 2010; Weber and Linka, 2011). Knockout out BASS2 caused mevastatin-sensitive phenotype (Furumoto et al., 2011; Zhao et al., 2016), but it is not clear whether FA biosynthesis or oil accumulation is affected in BASS2 mutant. In this study, it is the first time to report that the knockout of BASS2 could reduce the SOC by 2.8%–5.0% in B. napus. Metabolites of seed plastid were determined and glycolysis- and FA-related metabolites were significantly decreased in the mutants of BnaBASS2. Meanwhile, a lower level of energy-related metabolites ATP and NAD(P)H was detected in the mutant (Figures 2 and 3). Previous studies found that BASS2 showed sodium-dependent pyruvate uptake activity, and the sodium influx was balanced by sodium:proton antiporter (NHD1), which was required for exporting sodium from the chloroplast to maintain the sodium gradient required for the import of pyruvate (Furumoto et al., 2011). We hypothesize that disruption of BnaBASS2 can block the supply of pyruvate to the plastid, which affects the process of photosynthesis, glycolysis pathway, etc. These processes are closely related to energy. Due to the decreased level of pyruvate and energy, FA synthesis in the plastid was reduced and the formation of TAG is suppressed. Multiple studies showed that WRI1 is a master regulator in the glycolytic pathway, FA biosynthesis, and lipid metabolism (Adhikari et al., 2016; Baud et al., 2009; Kong et al., 2019, Figure 5 BnaBASS2 promotes seed oil accumulation in mature seeds. (a) Oil content in mature seeds. Data were measured by NIR. Values are means ± SD (n = 4–6). (b) Oil content in mature seeds. Data were determined by GC-FID. Values are means ± SD (n = 4–6). (c) Contents of neutral lipids TAG. Values are means ± SD (n = 6–8). (d) Triacylglycerol species content. Values are mean ± SD (n = 6–8). (e) OBs of mature seeds were examined by scanning transmission electron microscopy. Bottom is a zoom-in on the red box of upper. (f–h) Number of OBs per 100 μm², total area of OBs in 100 μm², and size per OB in (e). Image J software was used to calculate the number and size of OBs. Values are mean ± SD (n = 12–16). CW, cell wall; OB, oil body; PB, protein body. Different letters represent significant differences at P < 0.05, based on an ANOVA analysis with the Fisher LSD test.
Kong and Ma, 2018; To et al., 2012; Yang et al., 2022). In our research, genes involved in the FA and oil biosynthesis of seed were significantly increased in OE lines and decreased in mutants (Figure 4f). Increased SOC of OE lines and decreased SOC of mutant lines might result from the regulation of WRI1, which affects the glycolysis, FA synthesis, TAG synthesis, and OB formation in the process of seed oil accumulation. Hence, BnaBASS2 could impact multiple pathways to regulate seed oil accumulation.

The metabolite level of the whole seeds was also significantly reduced in mutants, which may be caused by photosynthesis. Reduced photosynthesis in the mutants leads to a reduction in carbon sources and thus a decrease in metabolite level. In this study, reduced photosynthesis was detected only in the leaves of mutants (Figure S10e–h), whether the photosynthesis of seeds is affected in mutants needs to be determined in the future. Source, sink, and flow are three important factors that determine crop yield, and high yield can only be achieved when the development of crops reaches sufficient source, large sink, and smooth flow (Adachi et al., 2020; Wang et al., 2021). In our research, photosynthesis was significantly reduced in the mutants, that is, the reduction in the source. Moreover, the number of silique number per plant, seed number per silique, and thousand seed weight were significantly reduced in mutants, that is, the reduction in the sink. Therefore, the yield reduction in the mutants may be caused by insufficient source and small capacity of the sink, which work together to limit the effects of push (source) and pull (sink). It is worth noting that, ultimately, the contents of sugar and protein are lower in OE seeds, while higher in mutants. And the content of starch is lower in mutants (Figure S8a–c). It indicates that more carbon flows into seed oil synthesis in OE lines.

In plants, there are several translocators that transport glycolysis metabolites from cytosol to plastid. Glucose 6-phosphate translocator (GPT), triose phosphate translocator (TPT), xylulose 5-phosphate translocator (XPT), and phosphoenolpyruvate translocator (PPT) transport G6P, 3-PGA, xylulose 5-phosphate, and PEP, respectively, from cytosol to plastid (Weber and Linka, 2011). Previous studies revealed that several plastidic phosphate translocators had an effect on plant growth and chloroplast development. GPT1 was essential for pollen maturation and normal embryo sac development, the mutation of GPT1 was lethal (Andriotis et al., 2010; Niewiadomski et al., 2005; Rolletschek et al., 2007). The combined loss of TPT and XPT showed severe growth retardation (Hilgers et al., 2018a, 2018b). Knockout of PPT1 caused slower growth, yellowing leaf, and abnormal development of chloroplast (Li et al., 1995; Streitfeld et al., 1999). However, in our study and previous studies, BASS2-deficient plants did not exhibit a visible vegetative growth phenotype under normal growth conditions (Figure S9). We speculate the main reason is that PEP metabolized by PKp in the plastid is the main source of plastidial pyruvate. In Arabidopsis, disruption of PKp could cause 60% reduction in seed oil, retarded embryo elongation, and seed germination (Andre et al., 2007; Andre and Benning, 2007; Baud et al., 2007b). At different stages of seed development in B. napus, the activity of PKp is slightly higher than that of PKc (Eastmond and Rawsthorne, 2000). In our study, although knockout of BnaBASS2 resulted in blocked pyruvate transport from cytosol to plastid, another source of plastidial pyruvate could compensate for the pyruvate import deficiency in the BnaBASS2 mutant. PPT1 transports PEP into the plastid, and PKp catalyses the synthesis of plastidial pyruvate, which is used for normal chloroplastic membrane lipid synthesis, thus maintaining normal growth. Our results suggest that loss of BnaA05.BASS2 and BnaC04.BASS2-1 resulted in a moderate yield penalty and a decrease in SOC in B. napus. Together, these results indicate that conversion of PEP to pyruvate by PKp provides a major source of pyruvate for metabolisms in the plastid.

Brassica napus is an important oil crop in the world, and improving oil production is one of the important goals of rapeseed breeding (Hua and Wang, 2016; Zou et al., 2022). Biotechnology has been broadly used to improve the oil content in oil seeds. At present, metabolic engineering strategies to improve seed oil accumulation can be divided into upregulation of FA biosynthesis (Push), increasing TAG assembly (Pull), and increasing TAG storage/preventing breakdown. For example, overexpression of transcription factors such as WRI1, LEC1, and LEC2 could improve seed oil accumulation by regulating genes involved in FA biosynthesis and functioning to ‘Push’ FA de novo synthesis (Baud et al., 2007a; Liu et al., 2010; Manan et al., 2017; Mu et al., 2008; Santos Mendoza et al., 2005; Shen et al., 2010; Tan et al., 2011). Overexpression of DGAT improved seed oil by ‘Pull’ much more acyl chains to DAG for TAG biosynthesis (Jako et al., 2001; Liu et al., 2015a; Vanhercke et al., 2013; Zhang et al., 2018). Gene silencing of Sugar-Dependent 1 (SDP1) could enhance seed oil by blocking the degradation of TAG (Kelly et al., 2013; Kim et al., 2014). In our study, overexpression of BnaBASS2 could significantly enhance the SOC in B. napus and had no obvious negative effect on agronomic traits. Thus, BnaBASS2 is a potential gene for breeding B. napus with high oil content.

Materials and methods

Analysis of BASS2 in B. napus

The genomic sequence, amino acid sequence, and different tissue expression data of BnaBASS2 were downloaded from BrnTIR (http://yanglab.hzau.edu.cn/; Liu et al., 2021). Phylogeny and amino acid sequence alignment of BnaBASS2 were performed by MEGA7 and CLUSTALW (https://www.genome.jp/tools-bin/clustalw).

Subcellular localization

The CDS of BnaA05.BASS2 was amplified and ligated into the pmDC83 vector, which contains a 2 x 35S promoter and a C-terminal GFP tag. The recombinant BnaA05.BASS2::GFP vector was transformed into the mesophyll protoplasts of Arabidopsis (Yoo et al., 2007) and epidermal cells of tobacco (Yang et al., 2021). The fluorescence pictures were observed using a Leica confocal microscope at 12 h after transfection.

Vector construction and plant transformation

To construct the OE vector of BnaBASS2, the CDS of BnaA05.BASS2 was cloned and ligated into p35S-FAST. To create the mutants of BnaBASS2, CRISPR/cas9-mediated gene editing technique was utilized (Xing et al., 2014). Briefly, two sgRNAs, which could both target BnaA05.BASS2 and BnaC04.BASS2-1, were designed on the website (http://cbi.hzau.edu.cn/cgi-bin/CRISPR; Lei et al., 2014). The amplified product by PCR including two sgRNAs was linked to pKSE401 vector using Golden Gate Assembly (Gao et al., 2013).

The OE and mutant vectors of BnaBASS2 were transformed into Agrobacterium tumefaciens GV3101, respectively. Genetic transformation of rapeseed was carried out by agrobacterium-
mediated method as described previously (Dai et al., 2020). Westar was utilized as a transformation receptor. The obtained OE plants were detected by PCR using specific primers.

RNA extraction and quantitative real-time PCR

30–50 mg developing seed was collected and total RNA was extracted using Eastep®Super Total RNA Extraction Kit (Promega Corporation, Madison, WI) (catalogue number LS1040). The quality of isolated RNA was determined by NanoDrop (Thermo Scientific, Waltham, MA) and 1.5% agarose gel electrophoresis, respectively. Then, 1–3 μg total RNA was used to synthesize cDNA with cDNA Synthesis SuperMix (TransGen, Beijing, China; catalogue number AE311). The synthesized cDNA was diluted 10 times, and then quantitative real-time PCR (qPCR) was conducted with an ABI 7500 system (ABI, Waltham, MA) using TransStart® Green qPCR SuperMix (TransGen; catalogue number AQ101). BnaACTIN7 was utilized as the internal control to normalize expression data according to the ∆∆CT Method (Livak and Schmittgen, 2013).

Western blot

For Western blot, the total soluble protein was extracted from 300 to 500 mg 24 DAF seeds using buffer A containing 50 mM Tris–HCl 7.5, 10 mM KCl, 1 mM EDTA, 0.5 mM phenylmethylene-sulfonyl fluoride (PMSF), 2 mM dithiothreitol (DTT) and protease inhibitors (Thermo Scientific, Waltham, MA; catalogue number PF200523). 50 μg protein was separated by 10% SDS-PAGE gel and the protein was transferred to the PVDF membrane. The membrane was immunoblotted with an anti-Flag antibody (Sigma, St. Louis, MO, catalogue number A3687). At last, the target BASS2 protein of 45 KD was detected and positive OE lines were obtained.

Identification of Cas9-edited mutants

To obtain mutants of BnaBASS2, primers were used to detect Cas9 in T0 transformed individual plants first, then the PCR product including sgRNA target sequence was amplified and sent to a sequence. Eventually, Cas9 edited mutations were analysed and acquired by DSDecode (Liu et al., 2015b). After multiple generations of breeding and identification, homozygous mutants of BnaBASS2 were obtained. Notably, the identification of mutants showed that only one target site on the eighth exon was edited and another target had no editing.

Procedure of developing seed collection

In full bloom, select a well-grown plant and pinch off the 2–3 flowers and then gently tie a wire between the open flower and the flower bud (Figure S12a). Four to five individual plant replicates were selected for WT, OE, and mutant lines, and each plant was labelled with the main branch and three branches. The individual plants were bagged and selfed, and the selfing bag was removed after 4 days (Figure S12b). The seeds close to the wire are the rapeseed seeds we want to take for the specified development days.

Chloroplast isolation from developing seeds

Intact chloroplasts of 31 DAF seeds were isolated according to the previous study with modifications (Zhang, 2010; Figure S4). One gram of rapeseed seeds were placed in 4 mL of well-prechilled organelle isolation buffer (300 mM sorbitol, 50 mM Tris–HCl PH 8.0, 50 mM EDTA PH8.0) and ground into a homogenate. The homogenate of seeds was filtered by a 300-mesh sieve (Figure S4a). The homogenate was centrifuged at 2000 g at 4 °C for 10 min, and the resulting precipitate was the crude chloroplast extract (Figure S4b). The crude extract was re-suspended in 300 μL of organelle isolation buffer and overlaid on a sucrose gradient (320 μL of 52% sucrose at the bottom and 280 μL of 30% sucrose at the top; Figure S4c). After centrifugation at 13,800 g at 4 °C for 1 h, the green bands at the interface between 52% sucrose and 30% sucrose were intrachloroplasts (Figure S4d). Chloroplasts were collected and chloroplasts examination was performed under an optical microscope to check the integrity (Figure S4e). Finally, isolated chloroplasts were precipitated and collected for the following experiments (Figure S4f).

Metabolite extraction and analysis

The metabolites involved in glycolysis and FA pathways were extracted and determined by LC–MS/MS according to previous research with modifications (Guo et al., 2014; Luo et al., 2007). Briefly, whole seeds and isolated chloroplasts of 31 DAF seeds were collected and freeze-dried (Millrock Technology, Kingston, NY). Five milligram of samples was weighed, and a mix of 1.8 mL methanol:chloroform (7:3, v/v) was added. Samples were incubated for 2 h at –20 °C and gently mixed several times during the incubation. 1.6 mL ddH2O was added and samples were vortexed and centrifuged at 100 g for 5 min. The upper was transferred into a new tube and dried using nitrogen. The extracts were re-dissolved with 200 μL ddH2O and then filtered with 0.45 μM cellulose acetate centrifuge tube filters. The re-dissolved extracts were used for metabolites analysis by LC–MS/MS (QTRAP 6500 plus). Mass signal intensity normalized by dry weight was calculated for comparing primary metabolites.

FA extraction and GC-FID analysis

5–10 mg oven-dried developing seeds or mature seeds were weighed and placed in a glass tube with Teflon-lined screw caps. 2.0 mL methanol with 5% (v/v) H2SO4 and 0.01% butylated hydroxyl toluene (BHT) was added. The seeds were crushed with a glass pestle, and then, 50 μL 16.2 μM heptadecanoic acid (17:0, used as an internal to quantify the FA) was added and the tube was capped tightly. Tubes were incubated at 85 °C for 2 h. Cool the samples, 2.0 mL H2O and 2.0 mL hexane were added, vortex briefly, then centrifuged at 1000 rpm for 5 min. The upper phase (around 1 mL) was transferred to a glass vial for GC-FID analysis. The GC-FID instrument and analytical parameters were set as described previously (Tang et al., 2020).

Lipid extraction and analysis

Lipids were extracted and analysed according to the previous report (Lu et al., 2019). 5–10 mg freeze-dried seeds of 31 DAF were weighed and dropped into 4 mL isopropanol at 75 °C with 0.01% BHT for 15 min. Samples were cooled, 3 mL chloroform and 1.2 mL ddH2O were added, and then, the samples were vortexed and agitated at room temperature for 1 h. Lipid extracts were transferred into a new glass screw-cap tube. 3 mL chloroform/methanol (2:1) with 0.01% BHT was added, and then, the tube was shaken for 30 min. The lipid extract was transferred into the glass screw-cap tube and this step is repeated several times until the seeds become white. 2 mL 1 μKCl was added to the lipid extracts and vortexed. Centrifuge at 1000 rpm for 5 min and then the upper phase was discarded. 2 mL ddH2O was added and vortexed. Centrifuge at 1000 rpm for 5 min and discard the upper phase. The whole extracts are dried with N2 and re-
dissolved in chloroform to 5 mg/mL. The samples were prepared for analysis as follows: 210 μL 300 mM ammonium acetate-methanol (1:20, v/v), 60 μL 5 mg/mL extracts; then, the volume was filled with chloroform to 300 μL. The lipids were quantified by LC–MS (TripleTOF 5600) and the level was calculated by mass signal intensity normalized by dry weight.

**Plant growth conditions and traits survey**

In this study, *B. napus* cultivar (cv.) Westar was used for gene cloning and genetic transformation. The seeds of WT, OE, and mutant were sown in the genetically modified field to set a plot experiment. The growth phenotypes were investigated, and the whole plants were harvested at maturity. Agronomic traits especially seed-related traits, such as silique length, seed number per silique, thousand seed weight, and the content of seed oil and protein were measured. The content of seed oil and protein was measured with a near-infrared reflectance spectroscopy (Foss NIRSystems 5000; Gan et al., 2003).

**Measurement of photosynthetic efficiency**

Seeds of WT, OE, and mutant were planted in the growth chamber PGW40 (Conviron, Surrey, BC, Canada) and its growth phenotype was observed indoors under a condition of 16 h light/8 h dark, 22/20 °C, and a relative humidity of 60%. The photosynthetic efficiency was measured using a portable photosynthesis system (LI-6800; Li-Cor, Lincoln, NE) with the parameters of 600 μmol s⁻¹ flow rate, 60% relative humidity, 400 μmol mol⁻¹ CO₂, 1000 rpm fan speed, 1200 μmol m⁻² s⁻¹ light intensity, 20 °C.

**Measurement of soluble sugar and starch content**

Total soluble sugar and starch of seeds and leaves were measured by the antherone method with some modification (Hong et al., 2018). 80–100 mg seeds or leaves were collected and added into 25 mL ddH₂O, heating at 95 °C for 30 min. 50 μL of supernatant was added to 400 μL 2 mg/mL anthrone sulfuric acid, followed by heating at 95 °C for 5 min. The mix of heated supernatant and anthrone sulfuric acid was measured at the wavelength of 625 nm. The concentration of soluble sugar was calculated according to the glucose standard curve. Content of soluble sugar (mg/g) = concentration × volume ÷ dry weight (DW).

The residues were dried and ground into powder. 5 mL 0.2 M KOH was added and the powder was ground into a homogenate. Then, the samples were transferred to a tube and heated at 95 °C for 30 min. 1 mL 1 M acetic acid was added and centrifuged at 12 000 rpm for 5 min. Finally, 50 μL of supernatant was used to perform the remaining steps as the measurement for soluble sugar. The concentration of hydrolysis product glucose was calculated according to the glucose standard curve. Content of starch (mg/g) = 0.9 × concentration × volume ÷ DW.

**OB observation of seeds by transmission electron microscope**

Peel the rapeseed seeds carefully and the inner cotyledons were collected. Then, the cotyledons were cut into 1 × 1 mm and fixed in 2.5% glutaraldehyde solution overnight. The preparation, sectioning, and observation of samples were completed by the electron microscope platform of Huazhong Agriculture University. A transmission electron microscope (TEM; H-7650, HITACHI, Tokyo, Japan) was used in the research. The number and size of OBs in WT, OE5, and CR253 were counted and measured by Image J.

**Primers**

All primers used in the study were listed in Table S2.

**Analysis of SOC and protein content**

In this study, both SOC and protein data are % of dry weight, and when the data are compared, the difference was directly used as the increase (decrease).

**Imaging software**

In this study, the heatmap, bar chart line chart, and data difference analysis are all completed by Origin software.

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**Author contributions**

L.G. conceived and supervised the study. S.T., N.G., F.P., Q.T., Y.L., and H.X. performed the experiments. S.T. analysed the data. S.T. wrote the manuscript. L.G. and S.L. revised the manuscript. All authors read and approved the manuscript.

**Conflict of interest**

The authors declare no conflict of interest.

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Supporting information

Additional supporting information may be found online in the Supporting Information section at the end of the article.

Figure S1 Multiple sequence alignment of BASS2 proteins and structure similarity in B. napus and Arabidopsis.

Figure S2 BnaA05.BASS2 is a chloroplast membrane protein by protoplast transformation of Arabidopsis thaliana.

Figure S3 Sequence analysis of CRISPR target genes BnaA05.-BASS2 and BnaC04.BASS2-1.

Figure S4 Procedures of seed plastid isolation.

Figure S5 DAG species content in 31 DAF seeds.

Figure S6 Expression levels of genes involved in glycolytic pathway of seeds.

Figure S7 Fatty acid composition of mature seeds was determined by GC-FID.

Figure S8 The protein (a), soluble sugar (b), and starch (c) content in mature seeds.

Figure S9 Growth phenotype in the growth chamber.

Figure S10 Phenotypic measurement of 40-day-old plants.

Figure S11 Growth phenotype at mature stage.

Figure S12 Procedures of developing seed collection.

Table S1 Measurement of agronomic traits after plants were harvested.

Table S2 Primers used in this study.