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**Engineering heat tolerance in potato by temperature-dependent expression of a specific allele of *HEAT SHOCK COGNATE 70*.**

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1. Summary

For many commercial potato cultivars, tuber yield is optimal at average day time temperatures in the range of 14-22°C. Further rises in ambient temperature can reduce or completely inhibit potato tuber production, with damaging consequences for both producer and consumer. The aim of this study was to use a genetic screen based on a model tuberisation assay to identify Quantitative Trait Loci (QTL) associated with enhanced tuber yield. A candidate gene encoding HSc70 was identified within one of the three QTL intervals associated with elevated yield in a Phureja-Tuberosum hybrid diploid potato population (06H1). A particular HSc70 allelic variant was linked to elevated yield in the 06H1 progeny. Expression of this allelic variant was much higher than other alleles, particularly on exposure to moderately elevated temperature. Transient expression of this allele in Nicotiana benthamiana resulted in significantly enhanced tolerance to elevated temperature. An TA repeat element was present in the promoter of this allele, but not in other HSc70 alleles identified in the population. Expression of the HSc70 allelic variant under its native promoter in the potato cultivar Desiree resulted in enhanced HSc70 expression at elevated temperature. This was reflected in greater tolerance to heat stress as determined by improved yield under moderately elevated temperature in a model nodal cutting tuberisation system and in plants grown from stem cuttings. Our results identify HSc70 expression level as a significant factor influencing yield stability under moderately elevated temperature and identify specific allelic variants of HSc70 for the induction of thermotolerance via conventional introgression or molecular breeding approaches.

2. Introduction

Potato is the third most important food crop in the world after rice and wheat. More than a billion people worldwide eat potato, and global crop production exceeds 300 MT per annum. Yet this crop is particularly vulnerable to increased temperature, which is considered to be the most important uncontrollable factor affecting growth and yield (Levy and Veilleux, 2007). Potato (Solanum tuberosum L.) originated in the Andes of South America from regions with cool temperatures and most cultivated germplasm is highly sensitive to elevated temperature. For most commercial cultivars, yield is optimal at average day time temperatures in the 14-22°C range, above which yield falls off sharply (Van Dam et al., 1996).

Elevated temperatures are known to affect numerous physiological processes in potato plants. Heat strongly suppresses tuberisation and reduces the proportion of assimilated carbon partitioned to tuber starch (Ewing, 1981; Wolf et al., 1991; Hancock et al., 2014). Compounding this problem, photosynthetic performance is also adversely affected by high
temperatures that reduce chlorophyll levels and CO₂ fixation rates (Reynolds et al., 1990). In addition, high temperature has a negative effect on potato tuber dormancy causing premature sprouting or secondary growth (Bodlaender et al., 1964). A significant interaction is observed between photoperiod and temperature (Menzel, 1985). During longer day lengths, lower temperatures are required for optimal tuberisation. However, effects of high temperature depend on the plant developmental stage; warm conditions can be beneficial during early growth phases while during tuber induction, cool temperatures especially during the dark period are essential (reviewed in Levy and Veilleux, 2007).

Whilst cultivated potato is generally a cool climate crop, there is significant variation in response to heat stress between cultivars (Marinus and Bodlaender, 1975; Mendoza and Estrada, 1979; Menzel, 1985; Levy, 1986; Levy et al., 1991; Midmore and Prange, 1991), in land races and wild potato species (Mendoza and Estrada, 1979; Hetherington et al., 1983; Reynolds and Ewing, 1989) and in progeny clones from heat tolerant parents (Mendoza and Estrada, 1979; Haynes and Haynes, 1983; Morpurgo et al., 1985; Veilleux et al., 1997).

Despite reported variation, we are unaware of any reports that identify QTLs for heat tolerance in potato. In contrast, in other crop species, QTL mapping studies have proven useful in identifying markers linked to heat stress tolerance. Multiple loci for heat tolerance have been identified in wheat (Paliwal et al., 2012) and maize (Messmer et al., 2009). A major QTL for high temperature germination and an additional QTL having smaller effects were identified in a genetic analysis of lettuce seed thermo-inhibition (Argyris et al., 2008).

A major difficulty in screening for complex abiotic stress tolerance traits is control of environmental parameters whilst growing sufficient numbers of genotypes under replication for genetic analysis. In potato, segregation for earliness of tuberisation is a confounding factor in heat stress screening and has led to the development of screens based on nodal cuttings (Ewing and Wareing, 1978; Van den Berg et al., 1990). This technique uses an excised potato leaf and its subtended axillary bud, that when maintained in moist compost, tuberises rapidly. Leaf cuttings from induced plants produce tubers (in ~14 days) at the axillary bud, whereas non-induced plants fail to tuberise. The nodal cutting assay therefore provides a convenient, space and time-saving high throughput method to study tuberisation and dry matter partitioning. The use of a model system for analysis of traits that are impacted on by multiple factors has the potential to simplify the trait and allows the investigator to focus on specific components.

In this study we have exploited the well-characterised 06H1 biparental diploid potato population (Prashar et al., 2014) to identify a locus associated with tuber yield at both conventional and mildly elevated temperature. Interestingly this diploid population does not
segregate for variation at the maturity locus on chromosome 5 (Kloosterman et al., 2013) showing fairly late uniform foliage maturity. A candidate gene encoding \( HSc70 \) was identified within the QTL interval and a particular allelic variant was linked to elevated yield in the 06H1 progeny. Expression of a transgene containing \( HSc70 \) under its native promoter in the potato cultivar Desiree resulted in enhanced \( HSc70 \) expression at elevated temperature leading to improved yield under moderately elevated temperature. Thus our results identify \( HSc70 \) expression level as a significant factor influencing yield stability under moderately elevated temperature and we identify specific allelic variants of \( HSc70 \) for the induction of thermotolerance via conventional introgression or molecular breeding approaches.

3. Results

Assessing tuber yield using the nodal cutting assay and QTL analysis.

Nodal cutting tuberisation assays were carried out for 170 genotypes from the 06H1 population. For each genotype, six nodal cuttings were prepared from two plants, using the 2\( ^{nd} \), 3\( ^{rd} \) and 4\( ^{th} \) node from the apex. Two independent experiments were conducted. Nodal cutting tuber yields (based on a total of 12 cuttings) from the 06H1 population grown at 22\(^\circ\)C and 28\(^\circ\)C ranged from 0.0002 to 3.674 g (22\(^\circ\)C) and 0.0002 to 2.277 g (28\(^\circ\)C), with means of 1.017 and 0.285 respectively, the data being considerably skewed towards zero (Figure 1). These data were used for QTL analysis using the MapQTL 6 software (Van Ooijen, 2011) and the SNP linkage map previously reported (Prashar et al., 2014). This linkage map comprises 2,157 mapped SNP markers generated using an Illumina SNP platform comprising a total of 8303 SNP markers. The 06H1 tuber yield data were subjected to a non-parametric Kruskal-Wallis (KW) analysis in MapQTL6 which essentially carries out a one way analysis of variance (ANOVA) to test for association between each mapped marker and the analysed trait. QTL effects were detected on three linkage groups, 1, 4 and 9. At 22\(^\circ\)C the only QTL effect detected was around the maternal marker c2_11487 at 11.53 cM (\( K^*=14.21 \)). At 28\(^\circ\)C QTLs were detected on linkage group 1 (biparental marker c2_49910 at 76.85 cM, \( K^*=21.32 \)), on linkage group 4 (maternal marker c2_11487 at 11.53 cM, \( K^*=14.22 \)) and on linkage group 9 (paternal marker c2_3948 at 11.64 cM, \( K^*=16.243 \)).

Identification of \( HSc70 \) as a candidate gene for tuber yield in the tuber nodal cutting assay

The genes encoded in the region of the QTLs on linkage groups 1, 4 and 9 were examined in the Potato Genome Browser hosted by Michigan State University (http://solanaceae.plantbiology.msu.edu/cgi-bin/gbrowse/potato/). For the linkage group 4 QTL inspection of the genomic region corresponding to ~5 cM either side of the QTL peak
revealed a strong candidate gene encoding HSc70 (PGSC0003DMG400027750). Previously, genetic variation in HSc70 has been suggested to underpin heat tolerance in cabbage (Park et al., 2013) and variation in heat shock protein levels have been correlated with heat tolerance in potato (Ahn et al., 2004). Inspection of a similar sized genomic region on linkage groups 1 and 9 revealed no obvious candidate genes. The presence of the HSc70 gene in the QTL region on linkage group 4 allied to results from previous studies with this gene led to our targeting it for further functional analysis.

**Identification of HSc70 alleles**

A region of the HSc70 gene was amplified from the parents of the 06H1 population (HB171(13) and 99FT1b5) by PCR (Figure S1). Sequence analysis of the PCR products identified four distinct alleles with the HB171(13) parent containing alleles designated A1 and A2 and 99FT1b5 containing alleles A3 and A4. A cleaved amplified polymorphic sequence (CAPS) assay (Konieczny and Ausubel, 1993) was designed to discriminate between the four HSc70 alleles (Figure 2) and the allelic complement of the genotypes from the 06H1 population was determined. For each genotypic class present in the population, the mean tuber fresh and dry weight yields from nodal cutting experiments were determined. Genotypes containing the A2 allele yielded the highest fresh weight at both 20°C and 28°C (p < 0.05), with the A2A3 genotype having the highest yields and the A1A4 combination the lowest yield (Table 1). The ‘phase’ of the marker data around the position of the gene on linkage group 4 are consistent with a maternally inherited QTL effect caused by action of the HSc70 gene, the A2 allele being linked in coupling with the segregating maternal SNP alleles. The deduced amino acid sequences encoded by the four alleles were also compared (Figure S2). This analysis indicated that the amino acid sequences are similar and are only different in the C-terminal substrate-binding domain (SBD) (Mayer, 2013) (Figure S2). In the sequences encoded by the A2 and A3 alleles there are 6 additional amino acids resides (KIEEVD) in this region compared with the proteins encoded by the A1 and A4 alleles. No amino acid sequence variation was specific to only one of the four alleles. To further dissect any functional significance of sequence variation, amplification of up to 1KB upstream of the presumed ATG start codon was achieved for each allele by designing further PCR primers (Table S1). Comparison of the promoter sequences of the four alleles identified sequence differences specific to the A2 allele (Figure 3). In particular, at ca. 495 bp upstream of the ATG start codon, the A2 allele uniquely has an extended run of ten TA repeats compared with four repeats in the other three alleles (Figure 3).
Expression levels of *HSc70* in genotypes with contrasting response to moderately elevated temperature

*HSc70* expression level was compared by qRT-PCR in four genotypes that contained the A2A3 allelic combination and gave good tuber yield in the nodal cutting assay at 28°C and with four low yielding genotypes containing A1A4 alleles (Figure 4). In both tubers and leaves *HSc70* expression was significantly higher in the A2A3 genotypes than in the A1A4 when maintained at the optimal temperature of 22°C (*p* < 0.05 for both organs). Furthermore, following transfer of cuttings to 28°C for four hours genotypes containing the A2A3 alleles exhibited a further significant increase in the abundance of the *HSc70* transcript in both leaves and tubers whereas there was no significant change in abundance (*p* = 0.8 to 0.3) of the same transcript in genotypes containing the A1A4 alleles (Figure 4).

In order to determine which *HSc70* allele was up-regulated on exposure to elevated temperature a semi-quantitative RT-PCR expression assay was carried out and gave similar results to those shown in Figure 4 (Figure S3a). The PCR products from the RT-PCR assay were sequenced and from 8 cloned products, sequence analysis showed that all were transcripts arising from the A2 allele (Figure S3b).

**Transient expression of *HSc70*** in *Nicotiana benthamiana* leaves

The function of the A2 *HSc70* allele was characterized in transient expression experiments conducted in *Nicotiana benthamiana*. A binary construct containing the A2 coding sequence and 1KB of sequence upstream of the ATG start codon was engineered and introduced into leaves of *Nicotiana benthamiana* by agroinfiltration. Following agroinfiltration, plants were placed in a chamber at 45°C under 12h light. After 24 hours, the *HSc70* expression level was compared in agro-infiltrated plants and mock inoculated plants. Those agro-infiltrated with the *HSc70* construct expressed the *HSc70* gene at levels ca. 7-fold higher than controls. Furthermore, plants agro-infiltrated with the *HSc70* construct exhibited significantly lower levels of cell membrane injury compared with controls (60% compared with 90%, *p* < 0.05, Figure 5). The values were consistent with the visual assessment of plant damage which was greater in controls (Figure S4).

**Promoter deletion analysis of the A2 *HSc70* allele**

The A2 allele was differentiated from the other three *HSc70* alleles by the presence of ten TA repeats in the promoter in a region 495 nucleotides upstream of the deduced start codon, whereas in the A1, A3 and A4 alleles only four TA repeats were present in this region (Figure 3). In order to test whether this repeat sequence was connected to the elevated
*HSc70* expression level observed for the A2 allele on exposure to elevated temperature, promoter deletions were tested. Promoter constructs with 4 to 10 TA repeats were engineered in a binary construct upstream of the A2 coding sequence. Transient expression of *HSc70* was measured on agroinfiltration of these constructs into *Nicotiana benthamiana* leaves following exposure to elevated temperature. *HSc70* expression level was significantly higher (6-fold) in plants agro-infiltrated with constructs containing 8 or 10 TA repeats than in those agro-infiltrated with constructs containing 4 or 6 repeats (Figure 6a). Furthermore, the cell membrane injury level was greater in the plants agro-infiltrated with constructs containing 4 or 6 repeats (Figure 6b), providing evidence that the TA repeat sequence is important for *HSc70* expression and reinforcing the link between *HSc70* expression and heat tolerance. The values were consistent with the visual assessment (Figure S7).

**Characterisation of transgenic potato lines over-expressing the A2 *HSc70* allele**

Transgenic potato lines expressing the A2 *HSc70* allele under its native promoter were generated in the cultivar Desiree and independent transgenic lines were screened for *HSc70* expression level in leaves (Figure 7). In leaves harvested from plants grown at 22°C, *HSc70* expression level was low in both transgenic lines and controls although one transgenic line (line 56) did exhibit significantly higher *HSc70* expression at this temperature. In plants subjected to 4 hours of elevated temperature (28°C) there was a dramatic increase in *HSc70* expression level particularly in line 56, where *HSc70* transcripts were 60-fold (p < 0.05) more abundant than in wild type controls. Lines 33 and 48 exhibited significantly greater expression levels (4 and 8-fold respectively) than wild type after elevated temperature treatment. Lines 33, 48 and 56 were therefore selected for further analysis.

Wild type and selected A2 *HSc70* expressing lines were exposed to elevated temperature (40°C) for 24 hours. Leaf cell membrane injury was assessed using an electrolyte leakage assay. After four hours, *HSc70* expression was greatly enhanced by up to 50-fold in all transgenic lines relative to wild type. However, following 24 hours exposure, there was no significant difference in expression level between any of the lines tested, with expression level in the transgenic lines declining to the levels prior to heat exposure (Figure 8a). No changes in electrolyte leakage from leaves were observed after 4 hours high temperature exposure in any of the genotypes however, after 24 hours there was significant increase in electrolyte leakage from wild-type leaves that was not observed in any of the *HSc70* transgenic lines (Figure 8b).
Protection of tuber yield at elevated temperature in A2 HSc70 expressing lines

Having established that elevated transient expression of A2 HSc70 provided a temperature-dependent protective effect as assessed by the electrolyte leakage assay, we wished to determine whether expression of the A2 allele could also protect tuber formation and yield at moderately elevated temperature. We therefore performed nodal cutting tuber yield assays. At 20°C yield was not significantly (p < 0.05) different between the wild type and any of the three transgenic lines tested (Figure 8). However, in Desiree WT, yield decreased by 75% at 28°C compared to 22°C (p < 0.05). Yield reductions were also observed in transgenic lines over-expressing HSc70 at higher temperatures however, these were not as extensive as observed in Desiree wild-type (Figure 9). For example in lines 48 and 56, fresh weight yield at 28°C was ca. 2-fold greater than in wild type controls at the same temperature (p < 0.05). Dry weight yield values also showed the same significant pattern with higher yield in the transgenic lines at elevated temperature (Table S3). Tuber yield was also measured in plants grown from stem cuttings. At maximum day temperature of 20°C, two of the three transgenic lines exhibited no significant difference in tuber yield between the over-expressing lines and wild type control while line 48 exhibited a significant increase in both fresh and dry tuber weight. In contrast, at 28°C tuber yield was significantly higher (p < 0.05) in all of the transgenic lines relative to the controls on both a fresh weight and dry weight basis (Table 2).

4. Discussion

Genetic analysis of tuber yield using nodal cuttings

For many years, a simplified model of the potato plant based on nodal cuttings has been used as a convenient system for studying tuberisation (Kumar and Wareing, 1972; Kumar and Wareing, 1974). It has also been suggested that cuttings can be used as a screen for heat tolerance (Ewing and Wareing, 1978). In addition, nodal cutting assays were developed to investigate the effects of temperature on tuber second growth (Van den Berg et al., 1990) where the negative effects of temperature on tuberisation were clearly demonstrated. Heat tolerance is dependent on both the ability to produce haulm that grows vigorously at high temperature and on maintaining carbon partitioning and tuber development. For genetic analysis it is useful to dissect complex traits into their components. In view of recent advances in potato genetics such as the availability of bi-parental populations and dense genetic maps (Prashar et al., 2014) we considered it timely to investigate the genetic control of tuber yield by deploying a nodal cutting system on a segregating population.
**HSc70 identification from genetic analysis**

In the diploid 06H1 population QTLs for tuber yield were identified on linkage group 4 and two other linkage groups. The linkage group 4 QTL for yield was detected at both temperatures tested (22°C and 28°C). Inspection of the genome browser indicated that there were a large number of genes in the linkage group 4 QTL region and so the approach did not yield definitive genetic evidence of the causative gene. Nevertheless, it did focus our attention on the set of candidate genes amongst which, a gene encoding HSc70 was considered a good candidate for yield protection particularly at the elevated temperature.

Heat shock cognate protein 70 (HSc70) is a constitutively expressed molecular chaperone which belongs to the heat shock protein 70 (HSP70) family (Al-Whaibi, 2011). In contrast to heat shock proteins (HSPs), heat shock cognate proteins are not generally induced by elevated temperature (McCallister et al., 2015). Despite this difference in expression response, no amino acid motifs have been defined that distinguish heat shock cognate proteins from HSPs. Phylogenetic analysis confirms that all four alleles of the presumed HSc70 gene in this study are more closely related to other HSc70 genes in Arabidopsis and tomato than to HSP70 sequences (Figure S5). Nevertheless the A2 allele identified here is clearly heat inducible, illustrating the difficulties in annotating this gene family. All organisms respond to heat stress by inducing heat-shock proteins (HSPs). HSP70s are the most abundant type of HSP, having important roles in preventing newly synthesized proteins from misfolding and aggregating. HSP70s are encoded by multi-gene families that can be divided into four subfamilies based on sub-cellular localization: cytosol, endoplasmic reticulum (ER), plastids, and mitochondria (Sung and Guy, 2003). In addition, *Nicotiana tabacum* contains a nuclear-localized HSP70, NtHSP70-1, which helps to prevent the fragmentation and degradation of nuclear DNA during heat stress (Cho and Choi, 2009). Although many studies have elucidated the molecular functions of individual family members, genome-wide analysis of this family is still limited, especially for crop species (Jung et al., 2013).

**HSc70 and protection from elevated temperature**

HSc70s are characterised by highly conserved ATPase and substrate-binding domains (SBDs) (Mayer, 2013). ATP binding by the ATPase domain induces conformational changes in the SBD, facilitating transient association with hydrophobic stretches in peptides. When stimulated by both substrate binding and the J-domains of co-chaperone proteins, HSc70s hydrolyse ATP to ADP, triggering a SBD conformation change resulting in capture of the hydrophobic substrate. Release of ADP by nucleotide exchange factors (NEFs) causes the SBD to return to the open conformation, releasing the substrate.
In Arabidopsis overexpression of HSc70 correlates with the acquisition of thermotolerance as well as an increase in tolerance to water deficit and salt stress (Lee and Schöffl, 1996; Leborgne-Castel et al., 1999; Alvim et al., 2001; Sung and Guy, 2003). Attempts to decrease HSc70 expression levels using a transgenic approach were unsuccessful implying that reduced HSc70 expression is lethal. Constitutive over-expression resulted in transgenic lines with impaired developmental phenotypes characterised by dwarfism and altered root structure. Nevertheless, these transgenic lines also exhibited a greater tolerance to heat shock (44°C for 10 minutes) than controls (Sung and Guy, 2003). These results led to the conclusion that it is necessary to tightly control HSc70 expression during development to avoid pleiotropic effects whilst having good tolerance to abiotic stresses. In the present study, we examined allelic variation both in the coding region and in the upstream promoter sequence of HSc70. Comparison of the deduced amino acid sequence encoded by the four alleles revealed differences in the coding region (Figure S2) although none specific to the A2 encoded protein. The A2 and A3 alleles contain the C-terminal sequences KIEEVD which are missing in the A1 and A4 alleles. The conserved C-terminal EEVD sequences of Hsp70 and Hsp90 mediate interactions with specialized tetratricopeptide repeat (TPR) domains in Hop and other related co-chaperones (Brychzy et al., 2003). The absence of this motif in the A1 and A4 alleles could be of functional significance and could account for the particularly low level of heat tolerance (based on yield at elevated temperature, Table 1) in the A1A4 genotype class. However, the A1A3 genotype class, containing one allele with the C-terminal domain also exhibits lower yields than genotypes containing the A2 allele.

Interestingly, the protective A2 allele exhibited a unique extended TA repeat approximately 495 bp upstream of the putative ATG start codon, implying that the A2 phenotype may be due to differences in expression patterns at elevated temperature. The best combination of alleles for enhanced yield arises from the A2A3 combination, where both alleles contain the KIEEVD domain as well as the high level of expression from the A2 allele and so it is possible that a combination of the A2 promoter sequence and the presence of the KIEEVD domain in both alleles is required for optimal yield.

In Desiree lines transformed with the construct in which expression was driven by the A2 promoter HSc70 expression was similar in wild-type and transgenic lines grown at 22°C however, on transfer to elevated temperature A2 expression was rapidly and transiently induced in transgenic lines. Transgenic lines did not exhibit any developmental abnormalities and our work therefore supports previous observations that appropriate HSc70 expression is essential to provide protection against abiotic stress while preventing adverse pleiotropic phenotypes. Our work further highlights the A2 promoter as a tool to drive heat inducible expression of a range of gene constructs in potato and other Solanaceae.
We are unaware of any previous reports that identify alleles of \textit{HSc70} genes that may confer different expression properties and hence impact on yields particularly under abiotic stress conditions. However, in this respect, reports of heat tolerance in cabbage genotypes are of relevance (Park et al., 2013). Heat-responsive gene expression profiles in four heat-tolerant and four heat-sensitive genotypes were compared. Significantly higher expression levels (ca. 8-fold) of an \textit{HSc70} gene were measured in all heat-tolerant lines than in the heat-tolerant lines, implying that \textit{HSc70} may be a marker for heat tolerance. Differential expression of small HSPs has been reported for several plant species including potato, where genotypic differences in heat tolerance correlated with small HSP expression. Two thermotolerant potato varieties expressed higher levels of small HSPs than two more sensitive varieties (Ahn et al., 2004). Similarly, in common bean, heat tolerant varieties express small HSPs at higher levels on exposure to heat stress (Simões-Araújo et al., 2003).

\textbf{Promoter motif associated with the differential response of the A2 allele}

In order to account for the different expression characteristics of the A2 allele of \textit{HSc70}, the promoter sequences of the four different alleles in the parents of the 06H1 population were compared. The most striking difference between the promoter sequences was an extended TA repeat motif in the A2 promoter which contained 10 repeats compared with 4 in the other alleles. In yeast it has been demonstrated that tandem TA repeats in promoter regions can result in enhanced expression levels (Vinces et al., 2009) due to effects on local chromatin structure. Whilst we are unaware of a specific effect of TA tandem repeats in plants in relation to heat stress, TA rich regions in plant promoters do often enhance gene expression (Sandhu et al., 1998). In this study, promoter deletions clearly demonstrate the relationship between the number of TA repeats and the \textit{HSc70} expression level on exposure to elevated temperature in transient assays in \textit{Nicotiana benthamiana}. It will be interesting to investigate the occurrence of the TA repeat allele in other potato genotypes and wild species.

\textbf{5. Conclusion}

Overall our results demonstrate that the presence of the A2 \textit{HSc70} allele confers tolerance to elevated temperature, both in the 06H1 population and transgenic lines where \textit{HSc70} expression is driven by the A2 promoter. In the nodal cutting system and in plants generated from stem cuttings, the presence of the A2 allele results in enhanced yield at elevated temperature. In transgenic lines, the yield was up to 2-fold greater than in wild type plants at 28\degree C but with no significant effect on yield at 20\degree C. In 06H1 genotypes however, yield was enhanced at both temperatures tested. In both cases we propose the rapid response of the A2 allele to temperature perturbation underlies the yield effect. Deployment of the A2 allele
in potato breeding may provide a strategy for enhancing yield either under temperate conditions or under periods of abiotic stress.

6. Experimental procedures

06H1 population

The mapping population used here (06H1) is a full-sibling progeny (n = 186) of a cross between two highly heterozygous diploid potato clones (HB171(13) and 99FT1b5), both of which result from crosses between diploids of Solanum tuberosum group Tuberosum and Solanum tuberosum group Phureja (Prashar et al., 2014).

Growth of plant material

06H1 clones were grown from cores (6mm diameter, excised from tubers, each containing a single bud) in 10 cm diameter pots containing standard compost mix. Plants were raised in a glasshouse maintained at a daytime temperature of 20°C and a nocturnal temperature of 15°C. Light intensity (photosynthetic photon flux density) ranged from 400 to 1000 μmol m² s⁻¹. Single nodal cuttings (Ewing and Wareing, 1978), were taken from 7-8 week old plants and the base of the petiole was placed in 50/50 coir/sand mix. A cutting consists of a fully extended leaf and its subtended bud. Cuttings were left in glasshouse conditions for 24h then moved to growth rooms set at 70% humidity, 12h photoperiod (light intensity of 400 μmol m⁻² s⁻¹) and various temperature regimes. Cuttings were watered daily with prewarmed water. Tubers were harvested after 3 weeks.

Four heat sensitive and four heat tolerant genotypes of the 06H1 population were grown from cores excised from tubers as described above. Plants were raised under glasshouse conditions for 8 weeks. Then, the plants were acclimated for 2 weeks under controlled environment conditions (20°C day/16°C night). Plants were moved to a cabinet at high temperature (28°C) and low temperature (22°C) for 4 hours, at a light intensity of 400 μmol m⁻² s⁻¹. After 4 hours, leaves and tubers were collected, immediately frozen in liquid nitrogen, and then stored at −80°C until use.

HSc70 overexpressing lines were grown from in vitro propagated tissue culture plants in 10 cm diameter pots containing standard compost mix. Plants were grown in a glasshouse (15-20°C, 12h light, 400 to 1000 μmol m⁻² s⁻¹) for 7-8 weeks. Single nodal cuttings were prepared as described above.
QTL analysis

QTL mapping of nodal cutting tuber yield data (22°C and 28°C) was performed using MapQTL® 6.0 (Van Ooijen, 2011) and Genstat 15.1 (VSN International Ltd.) software. The non-parametric Kruskal–Wallis (KW) test supported in MapQTL version 6.0 was performed. In this method, a single marker analysis was used to test the association of a marker with the trait at significance \( p \leq 0.001 \).

Allele mining and cleaved amplified polymorphic sequence (CAPS) assay

Genomic DNA was isolated from leaves using the AquaGenomic™ DNA isolation solution (http://www.aquaplasmid.com) according to manufacturer's instructions. PCR primers (Table S1) were designed to amplify a partial fragment from parental DNA templates. PCR was performed on 25 to 50 ng DNA in 50 µl reaction volume using 2.5 U of Platinum high fidelity DNA polymerase and buffer (www.invitrogen.com) containing 1.25 mM MgSO₄. Gene specific primers and deoxynucleotides (dNTPs) were used at a concentration of 0.4 µM and 200 µM, respectively. Thermal cycling conditions were: 2 min denaturation at 95°C followed by 25 to 40 cycles (30 sec at 95°C, 30 sec annealing at the appropriate Tm, 1 to 2 min extension at 68°C) followed by 5 min final extension step at 68°C. Polymorphisms were present in the parental sequences at positions that resulted in the presence or absence of a HindIII restriction site (Figure 2). PCR products were digested with HindIII prior to being analysed by electrophoresis on agarose gels and visualised following staining with ethidium bromide. Distinct restriction digestion patterns were observed for the four alleles identified (Figure 2).

qRT-PCR

RNA was extracted from potato leaves and tubers as described (Ducreux et al., 2008). The first strand cDNA templates were generated by reverse transcription, using random hexamers as primer and SuperScript II reverse transcriptase (Invitrogen Life Technologies, Carlsbad, California, USA). Potato elongation factor1-alpha (EF1α) primers were used as a control. The expression level of HSc70 was analyzed using the StepOnePlus Real-Time PCR system (Applied Biosystems) and StepOne Software version 2.3 (Applied Biosystems). Gene-specific primers and Universal probe Library (UPL, Roche Life Science) probes (Table S2) were used at a concentration of 0.2 µM and 0.1 µM, respectively. Thermal cycling conditions were: 10 min denaturation at 95°C followed by 40 cycles of 15s at 94°C, 60 s at 60°C. Relative expression levels were calculated and the primers validated using the Ct method (Livak, 1997). To normalize the values, an alternative method for calculating relative
quantification was used (Pfaffl, 2001). The HSc70 expression level in transgenic lines was determined using the same method.

**HSc70 Promoter analysis**

Genomic DNA was isolated and PCR performed as described for allele mining using the specific primers described in Table S1. PCR products were analysed by electrophoresis on agarose gels and DNA fragments eluted from the gel. The purified DNA fragments were ligated into pGEM-T and transformed into *E.coli* strain DH5α and the plasmid DNA from each clone was extracted using a DNA Plasmid Miniprep Kit (Promega, Madison, WI, USA) and sequenced with M13 universal primers on a 3730 automated DNA sequencer (Applied Biosystems: http://www.lifetechnologies.com) using a cycle sequencing protocol and the BigDye Terminator Cycle Sequencing Kit (version 3.1; Applied Biosystems). Analysis of sequences was performed using Sequencher software v.4.9 (http://genecodes.com/).

**Transgenic plants generation.**

The binary construct was designed using the GoldenBraid strategy (gb.cloning.org) (Sarrion-Perdigones et al., 2011). Specific primers for the HSc70-promoter from the A2 allele and the HSc70-A2 coding region containing the required BsmBI type II restriction sites were designed (Table 1). PCR amplification was performed using High Fidelity Polymerase (Invitrogen). PCR products were analyzed by agarose 1% gel electrophoresis and purified using a Promega kit. The PCR purified fragments were ligated into a pUPD entry vector using the BsmBI digestion-ligation reaction protocol of the GoldenBraid 2.0 cloning methodology. This method required repeated cycles of 37°C for 2 minutes and 16°C for 5 minutes. Once complete, the ligation reaction was transformed into *E.coli* strain DH5α (Invitrogen, USA). Cloned inserts were sequenced using M13 universal primers to verify the fragment sequence and the Type II restriction sites required for recombination in the expression vector. The HSc70 overexpression construct in pDGB3 was created by recombining GBparts (HSc70-promoter, HSc70-coding region, pTnos-terminator) as described by (Sarrion-Perdigones et al., 2011) using BsAl (New England Biolabs, Ipswich, MA, USA) restriction enzyme and T4 DNA ligase in 25-cycle digestion/ligation reactions. The binary construct was transformed into *Agrobacterium tumefaciens* strain AGL1 by electroporation (Curtis, 2004) Transformation of potato cv. Desiree using this construct was as described (Ducreux et al., 2005).

**Electrolyte leakage assay**

Cell membrane injury was assessed using an electrolyte leakage assay as described in (Campos et al., 2003). Four replicate samples of three 10mm leaf discs were punched from
each leaf sample assayed and placed in a test tube. The discs were washed twice with deionised water then 5ml of deionised water was added to each tube and samples were shaken for 1h in a 29°C incubator. The leachate was transferred to a 50 ml tube, 25 ml deionised water was added and the initial conductivity was measured using a conductivity meter. Samples were autoclaved, and total conductivity determined after cooling to room temperature. The extent of cell membrane injury was calculated as follows: [Initial conductivity x 100 / total conductivity].

**Transient expression in N. benthamiana**

For the transient expression experiments, plasmids were transferred to *A. tumefaciens* strain AGL1 by electroporation. Agroinfiltration was performed as described previously (Wydro, 2006). Overnight-grown bacterial cultures were pelleted and re-suspended in agro-infiltration medium (10 mM MES, pH 5.6, 10 mM MgCl₂, and 200 mM acetylsyringone) to an optical density at 600 nm of 1.2. Infiltrations were carried out using a needle-free syringe in leaves of *N. benthamiana* plants (growing conditions: 24°C day/20°C night in a 16h-light/8h-dark cycle) and the plants were kept at 22°C overnight. The following morning, the agro-infiltrated plants were moved to 45°C. After 24h, 12h photoperiod with light intensity of 400 µmol m⁻² s⁻¹, leaves were harvested for gene expression analysis and cell membrane injury assays as described above.

**Promoter deletion assays**

In order to investigate the role of the TA repeats in the A2 promoter in the region 495 base pairs upstream of the start codon promoter deletions were engineered and the effects tested on transient expression in *Nicotiana benthamiana*. Sequences from the *HSc70*-promoter containing 4, 6, 8 or 10 TA repeats were engineered using GenArt™ Gene Synthesis (Invitrogen) as illustrated in Figure S6. The binary constructs were generated using the GoldenBraid strategy as described previously, and plasmids were transferred to *A. tumefaciens* strain AGL1 by electroporation. The transient expression experiments in *N. benthamiana* were performed as described previously and leaves were harvested for gene expression analysis and cell membrane injury assays as described above.

**Statistical Analysis**

All analysis of variance (ANOVA) and the Student’s t-test were conducted using GenStat, 18th edition (VSN International, Oxford, United Kingdom).
7. Acknowledgements

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9. Tables

**Table 1.** Genotype means after heat stress screening of 188 genotypes from the 06H1 population. Fresh weight (FW) and dry weight (DW) are measured in grams (gr)

| Trait measured | Genotypic mean for A1A3 (gr) | Genotypic mean for A2A3 (gr) | Genotypic mean for A1A4 (gr) | Genotypic mean for A2A4 (gr) |
|----------------|-------------------------------|-------------------------------|-------------------------------|-------------------------------|
| FW at 20°C     | 2.27±0.28<sup>a</sup>         | 3.58±0.31<sup>b</sup>         | 1.46±0.18<sup>a</sup>         | 3.13±0.32<sup>b</sup>         |
| FW at 28°C     | 0.51±0.13<sup>ab</sup>        | 1.24±0.20<sup>c</sup>         | 0.35±0.08<sup>a</sup>         | 0.92±0.15<sup>bc</sup>        |
| DW at 20°C     | 0.50±0.2<sup>ab</sup>         | 0.76±0.18<sup>b</sup>         | 0.31±0.22<sup>a</sup>         | 0.85±0.18<sup>b</sup>         |
| DW at 28°C     | 0.11±0.06<sup>ab</sup>        | 0.30±0.08<sup>c</sup>         | 0.06±0.01<sup>a</sup>         | 0.24±0.08<sup>bc</sup>        |

Data are presented as mean ± standard error and different letters indicate significant differences between genotypes (Fisher’s, p<0.05).

**Table 2.** Weight of tubers from single stem plants of A2 *HSc70* expressing lines. Fresh weight (FW) and dry weight (DW) are measured in grams (gr).

| Trait measured | Line 33 (gr) | Line 48 (gr) | Line 56 (gr) | WT (gr) |
|----------------|--------------|--------------|--------------|---------|
| FW at 20°C     | 11.9±4.0<sup>a</sup> | 14.7±4.9<sup>b</sup> | 11.2±3.8<sup>a</sup> | 10.6±2.1<sup>a</sup> |
| FW at 28°C     | 15.3±5.1<sup>b</sup> | 19.7±8.8<sup>b</sup> | 17.6±4.9<sup>b</sup> | 8.8±1.8<sup>a</sup>  |
| DW at 20°C     | 1.89±0.63<sup>a</sup> | 2.93±0.98<sup>b</sup> | 1.7±0.57<sup>a</sup> | 1.96±0.30<sup>a</sup> |
| DW at 28°C     | 2.7±0.90<sup>b</sup> | 4.80±1.00<sup>c</sup> | 2.90±0.73<sup>b</sup> | 1.39±0.27<sup>a</sup> |

Data are presented as mean ± standard error (n=9) and different letters indicate significant differences between lines (Fisher’s, p<0.05).
10. Figure Legends

**Figure 1.** Frequency distribution of tuber yield from genotypes of the 06H1 population using the nodal cutting assay.

**Figure 2.** *HSc70* cleaved amplified polymorphic sequence (CAPS) assay. Four alleles were identified in the diploid parents. HB171(13) parent contains alleles A1 and A2 and 99FT1b5 contains alleles A3 and A4. Vertical lines represent HindIII restriction enzyme sites and numbers refer to nucleotide position.

**Figure 3.** *HSc70* promoter sequence of the four different alleles. TA extension in A2 at 495 bp upstream of ATG start codon is marked in red.

**Figure 4.** Relative expression level of *HSc70* in 06H1 population. (a) Tubers from A2A3 heat tolerant genotypes (genotype number 50, 153, 289 and 295); (b) tubers from A1A4 heat sensitive genotypes (genotype number 202, 206, 242 and 278); (c) leaves from A2A3 heat tolerant genotypes (genotype number 50, 153, 289 and 295) and; (d) leaves from A1A4 heat sensitive genotypes (genotype number 202, 206, 242 and 278). Asterisk indicates statistical differences between temperatures as determined by Student’s t-test. Error bars represent the standard error of the mean (n=3) (p<0.05).

**Figure 5.** Agroinfiltration of *Nicotiana benthamiana* plants. (a) Relative expression level of *HSc70* after one day at 45°C in agro-infiltrated plants with *HSc70* construct versus agroinfiltrated plants with empty vector (Mock).(b) Membrane damage in agro-infiltrated *Nicotiana benthamiana* plants after 1 day at 45°C by electrolyte leakage assay compared with agro-infiltrated plants with empty vector (Mock). Asterisks indicate significant difference with wild type at high temperature as estimated using Student’s t-test (p<0.05). Error bars represent the standard error of the mean (n=6) (p<0.05).

**Figure 6.** Promoter deletion analysis by agroinfiltration of *Nicotiana benthamiana* plants. (a) Relative expression level of *HSc70* after one day at 42°C in agro-infiltrated plants with *HSc70*-promoter deletion constructs versus agro-infiltrated plants with empty vector (Mock). (b) Membrane damage in agro-infiltrated *Nicotiana benthamiana* plants after one day at 42°C by electrolyte leakage assay compared with agro-infiltrated plants with empty vector (Mock). Asterisks indicate significant difference. Error bars represent the standard error of the mean (n=3) (p<0.05).

**Figure 7.** Expression level of *HSc70* gene in transgenic potato lines A2 *HSc70* allele after 4h at 22°C or 28°C relative to wild-type plants Desiree cv (WT). Asterisks indicate significant difference. Error bars represent the standard error of the mean (n=3) (p<0.05).
Figure 8. Screening of overexpression transgenic lines at high temperature (40°C). (a) Expression level of HSc70 gene in transgenic potato lines A2 HSc70 allele and wild-type plants Desiree cv (WT). (b) Membrane damage by electrolyte leakage assay of A2 HSc70 expressing lines and wild-type plants Desiree cv (WT). Asterisk indicates statistical differences with wild type at high temperature as determined by Student’s t-test. Error bars represent the standard error of the mean (n=6).

Figure 9. Fresh weight of tubers in A2 HSc70 expressing lines at 22°C and 28°C. Asterisk indicates statistical difference with wild type at high temperature (Fisher’s, p<0.05). Error bars represent the standard error of the mean (n=3) (p<0.05).

11. Supporting Information Legends

Figure S1. Clustal Omega alignment of the genomic DNA sequences of HSc70 alleles A1, A2, A3 and A4 isolated from parents of the 06H1 population (HB171(13) and 99FT1b5). Intron is indicated by blue text. Quantitative PCR primers and probe binding sites are highlighted in grey. Reverse primers used to amplify promoter regions: HSC70PA1R CTGAACGAGAATCATGAATCT; HSC70PCOMMONR AGATGCGAAGCGATTAATTGGT; HSC70PA3R TATACCAAACATAAACTCAT; HSC70PA4R TCCTAGCTCCAATACTAAACA

Figure S2. CLUSTAL O (1.2.1) multiple sequence alignment of the predicted HSc70 amino acid sequences from the C-terminal region of 4 alleles.

Figure S3. Expression level of HSc70 by semi-quantitative RT-PCR. (a) Bands of semi-quantitative RT-PCR in agarose gel (1%). HT means heat tolerate genotype; HS means heat sensitive genotype; C means control, L means 4h at 20°C and H means 4h at 28°C. (b) Sequence analysis of clone products with HSc70-A2 sequence and HSc70-A3 sequence as model, where all transcripts show GATT region as A2.

Figure S4. Phenotype of HSc70 agro-infiltrated and Mock inoculated plants of Nicotiana benthamiana after 24h at 45°C.

Figure S5. Phylogenetic tree of heat shock protein sequences resulting from a BLASTP search against Arabidopsis, Potato and Tomato databases using the translated HSc70 A2 allele. The tree was generated using Phylogeny.fr web service (Dereeper et al., 2008). The scale bar represents amino acid substitutions per site i.e., the number of changes or 'substitutions' divided by the length of the sequence.

Table S1. Primer sequences used for PCR and binary construct.
Table S2. Primer sequences and probe used in qRT-PCR.

Table S3. Nodal cutting results of HSc70 overexpression lines. Fresh weight (FW), dry weight (DW).

1. Figures

Figure 1. Frequency distribution of tuber yields from genotypes of the 06H1 population using the nodal cutting assay.
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```
Hsc70-promoter-A2: TCAAGTATATATAATATAATTTTCTCTGTAATAATTAACCTCAATTTTACAAAGCTCACACAGATTATTTTT -400 bp
Hsc70-promoter-A3: TCAAGTATATATAATATAATTTTCTCTGTAATAATTAACCTCAATTTTACAAAGCTCACACAGATTATTTTT
Hsc70-promoter-A4: TCAAGTATATATAATATAATTTTCTCTGTAATAATTAACCTCAATTTTACAAAGCTCACACAGATTATTTTT
Hsc70-promoter-A1: TCAAGTATATATAATATAATTTTCTCTGTAATAATTAACCTCAATTTTACAAAGCTCACACAGATTATTTTT
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Figure 7. Expression level of HSc70 gene in transgenic potato lines A2 HSc70 allele after 4h at 22°C or 28°C relative to wild-type plants Desiree or (WT). Asterisks indicate significant difference. Error bars represent the standard error of the mean (n=3) (p<0.05).
Figure 8. Screening of overexpression transgenic lines at high temperature (40°C). (a) Expression level of HSc70 gene in transgenic potato lines A2 HSc70 allele and wild-type plants Desiree cv (WT). (b) Membrane damage by electrolyte leakage assay of A2 HSc70 expressing lines and wild-type plants Desiree cv (WT). Asterisk indicates statistical differences with wild type at high temperature as determined by Student’s t-test. Error bars represent the standard error of the mean (n=6).
Figure 9. Fresh weight of tubers in A2 HSc70 expressing lines at 22°C and 28°C. Asterisk indicates statistical difference with wild type at high temperature (Fisher’s, p<0.05). Error bars represent the standard error of the mean (n=3) (p<0.05).