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Nutrient partitioning and grain yield of *TaNAM*-RNAi wheat under abiotic stress

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Abstract

Aims — Decreased expression of *TaNAM* genes by RNAi results in delayed senescence and decreased grain protein, iron, and zinc concentrations. Here, we determined whether *NAM* expression level alters onset of senescence under stress conditions, whether delayed senescence in the *TaNAM*-RNAi line resulted in improved tolerance to post-anthesis abiotic stress, and determined the effects of post-anthesis abiotic stress on N and mineral remobilization and partitioning to grain.

Methods — Greenhouse-grown WT and *TaNAM*-RNAi wheat were characterized in two studies: three levels of N fertility or water limitation during grain fill. Studies were conducted under both optimal and heat stress temperatures. Senescence onset was determined by monitoring flag leaf chlorophyll.

Results — Under optimal temperatures, *TaNAM*-RNAi plants had a yield advantage at lower N. *TaNAM*-RNAi plants had delayed senescence relative to the WT and lower grain protein and mineral concentrations, N remobilization efficiency, and partitioning of N and most minerals to grain.

Conclusions — Nutritional quality of *TaNAM*-RNAi grain was consistently lower than WT. Delayed senescence of *TaNAM*-RNAi plants provided a yield advantage under

optimal temperatures but not under water or heat stress. Discovery of specific *NAM* protein targets may allow separation of the delayed senescence and nutrient partitioning traits, which could be used for improvement of wheat.

Keywords: Wheat, grain mineral concentration, senescence, stress

Introduction

Grain yield and grain protein concentration are two principal targets of most hard wheat breeding programs. In addition to protein, staple crops also supply minerals for the human diet. The World Health Organization (WHO) reports that over 2 billion people are anemic (de Benoist et al. 2008), largely due to iron deficiency in their diets. In developing countries, 40 % of preschool children are estimated to be anemic. Iron deficiency anemia leads to impaired cognitive and physical development and reduced work productivity, and is a contributing factor to 20 % of maternal deaths. The

WHO also estimates the global frequency of Zn deficiency to be 31 % and extrapolates that the loss of disability-adjusted life years from Zn deficiency, globally, is over 28 million. Zinc deficiency contributes significantly to the mortality of children worldwide, although the problem is most severe in Africa, the Eastern Mediterranean, and southeast Asia (Black et al. 2008). Wheat grain protein, iron and zinc concentrations generally are positively correlated (Cakmak et al. 2004; Gomez-Becerra et al. 2010a, b; Morgounov et al. 2007; Murphy et al. 2011), which will facilitate simultaneous improvement of multiple minerals in wheat (Karami et al. 2009; Welch and Graham 2004).

Nitrogen (N) use by cereal crops to produce grain protein involves multiple processes, including uptake, assimilation, translocation, and remobilization. Grain N is supplied from N taken up post-anthesis and also from pre-anthesis stored N that is remobilized (Bogard et al. 2010; Kichey et al. 2007). A substantial percentage of the protein in wheat grain is supplied by amino acids remobilized from vegetative tissue (Barneix 2007; Gregersen et al. 2008; Masclaux-Daubresse et al. 2008) during leaf senescence (Hopkins et al. 2007). The proportion of grain N derived from remobilization of stored N has ranged from 45 to 95 % (Bogard et al. 2010; Ercoli et al. 2008, 2010; Kichey et al. 2007; Masoni et al. 2007; Waters et al. 2009). Likewise, Fe and Zn are remobilized from vegetative tissues in several species (Drossopoulos et al. 1996; Hocking 1994; Hocking and Pate 1977; Miller et al. 1994; Waters and Grusak 2008).

During leaf senescence, photosynthetic capacity decreases, metabolism and gene expression change markedly, and the leaf transitions from a sink to source tissue. Chloroplasts and proteins are disassembled, and the resulting amino acids scavenged (Lim et al. 2007). A number of genes have decreased expression during senescence, while others are more highly or specifically expressed during leaf senescence (Gepstein et al. 2003), and are known as senescence associated genes (SAGs). Mineral element remobilization, which herein is defined as the net loss of stored mineral content from an organ over time, is directly or indirectly associated with expression of SAGs (Gregersen et al. 2008), although functionality, with respect to mineral remobilization, has been assigned to only a few SAGs (Masclaux-Daubresse et al. 2008, 2010). Leaf senescence can be initiated prematurely by plant stresses such as

heat stress, drought, and nutrient deficiency. Some SAGs are expressed under both natural and stress-induced senescence, while others are expressed exclusively during specific senescence conditions (Buchanan-Wollaston et al. 2005; Van der Graaff et al. 2006). Avoidance of leaf senescence during drought stress, a phenotype known as stay-green, is associated with grain yield in maize, wheat, and sorghum (Harris et al. 2007; Snape et al. 2007; Thomas and Howarth 2000; Zheng et al. 2009).

Correlations between onset of senescence and grain protein concentration (GPC) and between onset of senescence and grain yield vary depending on environmental conditions (Bogard et al. 2011). The relationship between senescence and grain protein concentration is particularly evident in the wheat *Grain Protein Content (Gpc)-B1* QTL on chromosome 6BS (Kade et al. 2005; Uauy et al. 2006a). Using map-based cloning in a *Triticum turgidum* ssp. *durum* × *T. turgidum* ssp. *diccoides* genotype, the *Gpc-B1* QTL was identified as a NAC transcription factor, *NAM-B1* (Uauy et al. 2006b). Members of the large family of NAC transcription factors have roles in plant development, abiotic stress, and biotic stress responses (reviewed by (Nakashima et al. 2012)). Overexpression of some NAC family genes have resulted in improved tolerance to abiotic stresses (Mao et al. 2012; Xue et al. 2011). Alleles containing a 1-bp insertion or a deletion of *NAM-B1* predominate in cultivated wheat varieties (Asplund et al. 2010; Uauy et al. 2006b). Introgression of a functional *Gpc-B1* allele increased grain protein concentration (Brevis et al. 2010) as well as grain Fe, Mn, and Zn concentration (Distelfeld et al. 2007). Conversely, decreased expression of the four endogenous wheat *NAM-B1* family genes *TaNAM-A1*, *D1*, *B2*, and *D2*, (which were upregulated in flag leaves rapidly after anthesis), using a *TaNAM*-RNA interference (RNAi) construct, resulted in delayed senescence (Uauy et al. 2006b). Therefore, these genes likely act early in the regulatory pathway of developmental senescence. *TaNAM*-RNAi also decreased N remobilization and N partitioning to developing grain (Waters et al. 2009). Although grain weight was not significantly affected by *TaNAM*-RNAi expression, protein, iron, and zinc concentrations were significantly lower in grain from *TaNAM*-RNAi plants (Uauy et al. 2006b; Waters et al. 2009).

The previous studies that characterized the senescence effects of the *TaNAM*-RNAi on wheat plants

were performed under optimal conditions of temperature, available moisture, and N-sufficiency. However, the effects of the *TaNAM*-RNAi trait have not been evaluated in plants grown under stress conditions. Additionally, since N supply substantially influences N efficiency (Górny et al. 2011), experiments evaluating the effects of the *TaNAM*-RNAi trait on N dynamics under sub-optimal N supply were needed. Moreover, mineral uptake is intimately connected with nutrient and moisture availability, as well as climatic conditions (Karami et al. 2009). For example, N availability has been shown to have effects on uptake and translocation of other grain minerals. Iron and zinc concentration of wheat grain, and Zn and Fe uptake and remobilization were substantially increased by improved N fertility (Aciksoz et al. 2011; Cakmak et al. 2010; Erenoglu et al. 2010; Kutman et al. 2010, 2011a; b; Shi et al. 2010).

Efforts to improve the nutrient use efficiency of crop plants are motivated by economic factors (e.g. increasing fertilizer costs), and by recognition of detrimental environmental effects of excess fertilizer use. Genetic variation for grain protein concentration appears to be particularly associated with post-anthesis N uptake (Bogard et al. 2010). Genetic variability for N remobilization efficiency has been observed in wheat within existing varieties (Bogard et al. 2010; Kichey et al. 2007; Van Sanford and MacKown 1987), and alterations in N remobilization efficiency in the *TaNAM*-RNAi line (Waters et al. 2009) indicate that there are metabolic processes that could potentially be targeted to improve this trait. Next Generation Sequencing of the GPC-RNAi and WT transcriptomes 12 days after anthesis, when *NAM* gene expression was decreased by 35 % relative to WT, revealed 691 genes that were differentially regulated by the *NAM* genes (Cantu et al. 2011), some of which may be involved with regulating senescence and N and mineral translocation to seeds.

Here, we address three primary objectives. The first objective was to determine whether expression level of *NAM* genes alters onset of senescence under stress conditions that induce premature senescence. A second objective was to determine whether delayed senescence in the *TaNAM*-RNAi line results in improved tolerance (in terms of yield) to post-anthesis abiotic stress. A third objective was to determine the effects of post-anthesis abiotic stress (heat, water, and low N stress) on N and mineral remobilization and partitioning to grain.

Materials and methods

Plant materials and growth

Seeds of WT hexaploid spring wheat cv. 'Bobwhite' and the *TaNAM*-RNAi line were generously provided by Jorge Dubcovsky (University of California-Davis, Davis, CA, USA). Six seeds were planted per pot (3 seeds *TaNAM*-RNAi, 3 seeds WT) in 2.2 kg 1:1 soil: sand in clay containers 15 cm deep, 16 cm diameter at the top and 10 cm diameter at the bottom. Seedlings were later thinned to 2 plants of each line, for a total of 4 plants per pot. Treatments were assigned randomly to pots. At least 6 pots (12 plants of each genotype) were assigned to each treatment. Plants were grown in a greenhouse with supplemental lighting to provide a 16 h photoperiod.

Two studies were conducted, a fertility response study and a moisture stress study. The fertility response study was a factorial arrangement of a completely randomized design where the factors were genotype (WT, *TaNAM*-RNAi) and fertility treatment (3 levels). Fertility treatment levels were selected to represent full fertility and two levels of reduced fertility (moderate and severe), with the severe deficiency meant to induce premature senescence. All treatments in the fertility response study were well-watered. The moisture stress study was conducted as a factorial arrangement of a completely randomized design where the factors were genotype (WT, *TaNAM*-RNAi) and post-anthesis applied irrigation (moisture stress, well-watered). Plants in the moisture stress studies were fertilized at the highest rate used in the fertility studies. The studies were conducted in two environments. Environment 1 was a heat stress regime: day temperatures were set for 27–31 °C, night temperatures at 21–24 °C. Environment 2 was an optimal temperature regime: day temperatures were set for 18–24 °C, night temperatures for 13–18 °C.

Fertilization was supplied in a nutrient solution containing 0.5 mM MgSO_4 , 1 mM CaCl_2 , 0.1 mM NaH_2PO_4 , with N supplied as KNO_3 at rates indicated below. Fertility treatments were initiated with a uniform base application 33 and 28 DAP in environments 1 and 2, respectively. Three differential fertility treatments were applied (0, 43.75, or 87.5 mg N per application) at 4–6, 11–13, and 16–20 d after the base application. Fertility treatments were completed 1 and 4 days prior to anthesis of

the first plants in environments 1 and 2, respectively. Total N applied to the three treatments was 35 mg, 166 mg, and 298 mg pot⁻¹. Soil test data indicated that potting soil provided an additional 35 mg of N to each pot, and as there were four plants per plot, available N was estimated as a combined (soil + applied) total of 17.5, 61.25, and 105 mg N per plant. Containers in the fertility studies received 250 ml water each 2 d from anthesis until the conclusion the experiment. For the moisture stress study, differential irrigation treatments were applied to plants beginning with the first irrigation during anthesis. Two treatments (moisture stress = 150 ml water, well-watered = 250 ml water) were applied each 2 d from anthesis until the conclusion the experiment.

Tissue sampling

Above ground tissue of 6 plants of each genotype and treatment were harvested at anthesis. At maturity, spikes and vegetative tissues were harvested separately from at least six plants of each genotype and treatment. Plant samples were dried in a drying oven at 60 °C for at least 48 h. Grains were removed from spikes, then spikes were pooled with vegetative tissues from the same plant. Spikes less grain are considered vegetative tissues for the purposes of this study. Tiller number, plant and seed dry weight and kernel number were recorded. Tillers were ground in a Wiley mill, while seeds were ground in a coffee grinder for analysis of mineral composition and N analysis. Mineral concentrations were determined by ICP-MS following digestion of tissues in nitric acid/hydrogen peroxide as described previously (Waters and Grusak 2008). N was determined by combustion analyses using a Costech Analytical ECS 4010 (Costech Analytical Technologies, Valencia, CA). Protein concentration was calculated as N × 5.7.

Senescence was monitored by the decrease in relative leaf chlorophyll of flag leaves using a handheld SPAD-502Plus chlorophyll meter (Konica Minolta, Japan). Means of 10 readings were recorded from 6 plants for each treatment and genotype through 42 days after anthesis (DAA) in the heat stress environment and through 38 DAA in optimal temperature environment in the fertility studies, and through 30 DAA in the irrigation studies.

Real-time RT-PCR

Total RNA was extracted from node, peduncle, lower leaf, stem, and spike tissues at anthesis, and from node, rachis, peduncle, senescing lower leaf, green lower leaf, flag leaf, florets + grain, and stem tissues at 14 DAA using the Aurum Total RNA Fatty and Fibrous Tissue kit (Bio-Rad, Hercules, CA, USA). RNA quality and concentration was determined by UV spectrophotometry. One µg of DNase-treated RNA (RNase-free DNase I, New England Biolabs, Ipswich, MA, USA) was used for cDNA synthesis, using the High Capacity cDNA Reverse Transcription kit (ABI, Foster City, CA, USA) with random hexamers at 2.5 µM final concentration. Real-time RT-PCR primers and conditions were based on those of a previous study (Uauy et al. 2006b). cDNA corresponding to 50 ng of total RNA was used in a 15 µL real-time PCR reaction performed in a MyIQ (Bio-Rad, Hercules, CA, USA) thermal cycler using SYBR GreenER qPCR Super-Mix (Invitrogen Technology, Carlsbad, CA, USA) and 0.2 µM of gene-specific primers (see below). The following standard thermal profile was used for all PCRs: 50 °C for 2 min, 95 °C for 8 min; 40 cycles of 95 °C for 15 s and 60 °C for 30 s. At the end of each reaction, a dissociation curve was performed to ensure primer specificity. The Ct values for all genes were calculated using BioRad IQ5 System Software version 2.0 (Bio-Rad, Hercules, CA, USA) and normalized to the Ct value of actin using the equation $Y = 2^{-\Delta Ct}$, where $\Delta Ct = Ct_{UBQ10} - Ct_{target\ gene}$. The PCR efficiency from the exponential phase (E) was calculated for each individual amplification plot using the equation $(1 + E) = 10^{\text{slope}}$ calculated by the LinRegPCR program (Ramakers et al. 2003). PCR efficiency ranged from 97 to 99.5 %, with correlation coefficient (r^2) ranging from 0.98 to 1.0. The calculated PCR efficiency was then used to normalize each obtained Ct value. The primer sequences used for actin, *NAM-A1*, *-D1*, *-B2*, and *-D2* were described previously (Uauy et al. 2006b).

Calculations

Vegetative N content (per plant): N concentration (mg g⁻¹) × Vegetative tissue dry weight (g plant⁻¹)
 Harvest index (HI): Seed yield (dw)/(seed yield + above ground vegetative tissue dry weight)
 Nutrient Harvest index: Seed nutrient content

Figure 1. Expression of *NAM* genes in wild-type wheat tissues, normalized to actin. Inset shows *NAM-A1*, *-D1*, *-D2*, and *-B2* quantification by real-time RT-PCR at anthesis. The main panel shows expression of the same genes at 14 DAA.

(mg plant⁻¹)/Whole plant nutrient content (mg plant⁻¹)

Net Remobilization: Plant mineral or N content at anthesis (mg spike⁻¹) – Vegetative content at maturity (mg spike⁻¹)

Remobilization Efficiency: 100 × Net remobilization/Plant mineral or N content at anthesis

Post-anthesis N Accumulation (PANA): Whole plant content at maturity (mg plant⁻¹) – Plant content at anthesis (mg plant⁻¹)

Normalized Partitioning Quotient (PQ): 100 × (Nutrient harvest index/Harvest index)/Experimental Mean

Statistical analysis

Data were analyzed by analysis of variance using PROC GLIMMIX in SAS. Growth in each environment was conducted for one season; therefore tests of environment effects could not be made. Residual variances in the two environments were highly heterogeneous ($p < 0.001$) in both the fertility and moisture stress studies for the key traits of plant DW, seed DW, and seed number. Therefore, environments could not be combined for statistical analysis and separate analyses were conducted of each study in each environment. Main effects of treatment and linear and quadratic effects of fertility rate, and their interactions with the main effect of genotype, were tested using Type I sums of squares. Least squares adjusted means are presented in tables, and mean

differences (net remobilization, PANA) were estimated using LSMESTIMATE in PROC GLM.

Results

Senescence effects of *NAM* expression

In a previous study, wheat *NAM* genes were up-regulated rapidly after anthesis, however, expression was quantified only in flag leaves (Uauy et al. 2006b). Since other tissues (i.e. spike, peduncle, lower leaves) also exhibit delayed monocarpic senescence in the *TaNAM-RNAi* line, we tested whether the *NAM* genes were expressed in other tissues in wild-type hexaploid spring wheat. By real-time RT-PCR, no or very little mRNA for *NAM-A1*, *-D1*, *-D2*, and *-B2* was detected in nodes, peduncles, green lower leaves, flag leaf, stem, or whole spike in pre-anthesis wheat plants (Figure 1, inset). In plants 14 d later, during grain fill, expression was still low in the rachis and florets + grain of the spike, the peduncle, stem, and green lower leaves. Expression of *NAM-A1*, *-D1*, and *-B2* were slightly higher in green lower leaves, stem, and nodes, and moderately expressed in senescing lower leaves. High levels of expression were observed for all four *NAM* genes in flag leaf during grain fill.

SPAD meter readings of flag leaf chlorophyll were used to monitor senescence (Figure 2), and demonstrated the previously described phenotypic difference in timing of senescence onset between

Figure 2. Flag leaf chlorophyll levels of *TaNAM*-RNAi and WT wheat plants determined by a Minolta SPAD meter. a–d: fertility study; 17.5, 61.25, and 105 refer to mg N supply per plant. a, *TaNAM*-RNAi plants grown in the heat stress environment; b, *TaNAM*-RNAi plants grown in the optimal temperature environment; c, WT plants grown in the heat stress environment; d, WT plants grown in the optimal temperature environment. e and f: moisture stress study; 150 and 250 refer to volumes of water supplied every 2 d. e, *TaNAM*-RNAi and WT plants grown under heat stress; f, *TaNAM*-RNAi and WT plants grown under optimal temperatures.

the *TaNAM*-RNAi and WT plants. In both environments, chlorophyll degradation was initiated earlier in WT plants than in *TaNAM*-RNAi plants. Senescence was earliest in plants of the low N treatments. Low N RNAi plants had 20 % lower chlorophyll at 37 d under heat stress (Figure 2a) and 38 d under optimal temperatures (Figure 2b), while WT low N plants had 20 % decreased chlorophyll at 24 d under the heat stress regime (Figure 2c) and 25 d under optimal temperatures (Figure 2d). *TaNAM*-RNAi plants of intermediate and high N treatments had lost less than 20 % of their chlorophyll concentration at the end of the experiment, while WT intermediate and high N lost 20 % of chlorophyll at 30 d in the heat stress environment and 28 d in the optimal temperature environment. Thus, heat stress did not accelerate senescence of either WT or *TaNAM*-RNAi at low or intermediate N supply. High N WT plants lost 20 % of chlorophyll at 30 d in the heat stress environment, while they senesced later under optimal temperatures, at 35 d. The moisture stress studies were imposed at the high N rate only, and because of more rapid senescence, were terminated sooner than the fertility studies. Under heat stress at 20 d after anthesis, flag leaf chlorophyll in moisture stress treated WT plants had decreased by 52 % (Figure 2e); under optimal temperatures, it had decreased by 14 % (Figure 2f). In contrast, flag leaf chlorophyll was unchanged in *TaNAM*-RNAi plants under both heat and optimal environments. At 30 d, WT plants had 86 % lower chlorophyll in the heat stress environment with moisture stress and 68 % lower under optimal temperatures with moisture stress. In comparison, chlorophyll in flag leaves of RNAi plants decreased 36 % in the heat stress environment with moisture stress and 27 % under moisture stress in the optimal temperature environment.

WT vs RNAi in heat stress and optimal environments

The fertility studies were conducted under well-watered conditions; therefore effects of environment were not confounded with effects of moisture limitation in these studies. The responses of genotypes to environments are discussed first, averaged across N treatments, and specific interactions with fertility will be discussed below. Heat stress and optimal temperature environments produced substantial differences in seed yield in the fertility studies

(Table 1). Plants grown in the optimal temperature environments had vegetative mass approximately 3 times larger (data not shown), and 5 times the seed yield of plants grown in the heat stress environment, averaged over N treatments (Table 1). The heat stress environment diminished tiller number: these plants produced only primary tillers, whereas under the optimal temperature environment in Exp 2, the RNAi and WT plants averaged 3.0 and 2.7 tillers, respectively. The diminished tillering in the heat stress environment did not fully explain the effects of the high temperature environment on grain yield. Grain yields per spike of *TaNAM*-RNAi and WT plants in the heat stress environment were 58 % and 56 %, respectively, of grain yield per spike in the optimal temperature environment, resulting from fewer seeds per spike under heat stress. The overall effect of genotype on seed yield was consistent and significant in the fertility studies (Table 1). In both experiments, *TaNAM*-RNAi plants produced significantly greater seed yield, with more seeds than WT, and with heavier seeds in heat stress environment (Table 1). The yield harvest index of WT, however, was greater than *TaNAM*-RNAi in both environments, because of greater vegetative biomass in the *TaNAM*-RNAi plants. Fertility treatments significantly affected kernel weight in both environments (Table 1). Under the heat stress environment, kernel weight was maximized by intermediate fertility (Figure 3); however, under optimal temperatures, kernel weight was lowest at intermediate fertility.

Seed protein concentration of *TaNAM*-RNAi was significantly lower than WT in both environments (Table 2, Figure 4). Despite increased seed yield of *TaNAM*-RNAi plants, WT plants had significantly greater total quantities of seed N (Table 2). Averaged across N treatments, N remobilization efficiencies were lower in *TaNAM*-RNAi than WT plants, at 19.4 % v. 42.3 %, respectively, under heat stress, and 59.1 % v. 74.9 %, respectively, under optimal temperatures (Table 2), although due to variability this difference was not statistically significant in the optimal temperature environment. Partitioning of N to seeds, measured as N harvest index, was greater in WT plants in both environments. Because treatments affected plant morphology and grain harvest index, the partitioning quotient (ratio of NHI/HI) was used to compare nitrogen use efficiency (NUE) among treatments. Partitioning quotients were normalized to the mean of

Table 1. Effect of genotype, *TaNAM*-RNAi or Wild-type (WT), on vegetative biomass and yield components in fertility studies conducted under heat stress and optimal temperature environments. Means are presented as the mean $\pm$ standard error

Genotype	Tiller no.	Plant dwt per tiller	Seed no. (seeds plant ⁻¹)	Seeds per spike	Kernel wt (mg kernel ⁻¹)	Seed yield (g plant ⁻¹)	Seed yield (g spike ⁻¹)	Harvest index
<i>Heat stress environment</i>								
<i>TaNAM</i> -RNAi	1.0	1.32 $\pm$ 0.04	21.0 $\pm$ 0.8	21.0 $\pm$ 0.8	36.4 $\pm$ 0.4	0.76 $\pm$ 0.03	0.76 $\pm$ 0.03	0.365 $\pm$ 0.007
WT	1.0	0.97 $\pm$ 0.04	18.6 $\pm$ 0.8	18.6 $\pm$ 0.8	35.1 $\pm$ 0.4	0.66 $\pm$ 0.03	0.66 $\pm$ 0.03	0.400 $\pm$ 0.007
F-value ^a								
Genotype	ns	38.5**	4.2*	4.2*	5.0*	5.5*	5.5*	11.1**
Fertility Trt	ns	7.6*(2)	ns	ns	12.2** (2)	4.4*(2)	4.4*(2)	ns
Geno $\times$ Fert	ns	ns	5.3*(2)	5.3*(2)	ns	4.2*(2)	4.2*(2)	ns
<i>Optimal temperature environment</i>								
<i>TaNAM</i> -RNAi	3.01 $\pm$ 0.12	1.58 $\pm$ 0.05	107 $\pm$ 5	36.0 $\pm$ 1.1	35.8 $\pm$ 0.5	3.78 $\pm$ 0.17	1.30 $\pm$ 0.04	0.452 $\pm$ 0.006
WT	2.74 $\pm$ 0.13	1.22 $\pm$ 0.05	88 $\pm$ 5	32.1 $\pm$ 1.1	35.0 $\pm$ 0.5	3.20 $\pm$ 0.18	1.16 $\pm$ 0.04	0.488 $\pm$ 0.007
F-value								
Genotype	ns	26.0**	8.9*	5.9*	ns	10.5**	5.4*	18.8**
Fertility Trt	4.9*(1)	ns	ns	ns	18.4** (2)	5.2*(1)	ns	ns
Geno $\times$ Fert	ns	ns	ns	ns	ns	ns	ns	4.6*(2)

a. The F-value was determined by analysis of variance. Values in parentheses after significant effects of fertility treatment indicate the highest order of the response (1 = linear; 2 = quadratic) to fertility treatment

* and ** indicate significance at $p < 0.05$ and 0.01 , respectively; ns = non-significant

the study to facilitate discussion of relative effects of treatments on nutrient use efficiency (as per (Masclaux-Daubresse and Chardon 2011)). The normalized partitioning quotient (NPQ) for N of the WT was >100 in under both heat stress and optimal temperature environments, while the NPQ of *TaNAM*-RNAi was <100 under both heat stress and optimal temperature environments (Table 2), indicating that WT plants had greater NUE than the *TaNAM*-RNAi line plants under both environments. Concentrations of minerals in vegetative tissues and contents for vegetative and seed tissues analyzed in the fertility studies are included in Online Resource 1. The mineral concentrations of seeds (Table 3) grown in the optimal temperature environment were similar to the concentration in seeds grown in the heat stress environment. The concentration of most minerals was greater in seed of WT plants than in seed of *TaNAM*-RNAi plants in both environments, with the exception of K, which was lower in WT. The *TaNAM*-RNAi genotype affected partitioning of minerals between reproductive and vegetative tissues. Similar to the N NPQ results, *TaNAM*-RNAi plants generally were less efficient in partitioning minerals into grain (NPQ <100), while WT plants generally had NPQs >100 (Figure 5). However, in both heat

stress and optimal temperature environments, WT partitioned less K to grain in proportion to dry matter than the *TaNAM*-RNAi line.

WT vs RNAi at differing N rates

WT seed yield and seed quantity had second-order responses to fertility when grown in the heat stress environment; however, seed yield and quantity of *TaNAM*-RNAi plants was not affected by fertility when grown under heat stress (Table 1, Figure 3). In the heat stress environment, fertility treatment did not affect seed protein concentration (Table 2, Figure 4). In contrast, in the optimal temperature environment, N concentration in seed increased linearly with increasing fertility. Seed protein concentration of the *TaNAM*-RNAi and WT genotypes responded similarly to fertility treatments in both environments, although the *TaNAM*-RNAi line had consistently much lower GPC (approximately 28 % lower) than WT.

N fertility treatment affected N harvest index (NHI) and NPQ similarly in WT and *TaNAM*-RNAi plants in both environments (Table 2). Effects of fertility treatments were small relative to effects of genotype in both environments. Therefore, despite

Figure 3. Grain yield and seed number and kernel mass of *TaNAM*-RNAi and WT wheat plants in the fertility study. a, grain yield; b, seed quantity per plant in the heat stress and optimal temperature environments; c, mass per kernel, averaged across genotypes. Asterisk indicates that F-value is significant between genotypes at $p < 0.05$

varying responses to fertility treatment in the two environments, in both environments, *TaNAM*-RNAi plants partitioned less N to grain than WT plants. *TaNAM*-RNAi plants consistently had lower net nitrogen remobilization efficiency than WT plants (Table 2). WT and *TaNAM*-RNAi plants had similar post-anthesis N accumulation (PANA) in each environment (Figure 5), which was affected by fertility treatment significantly in both environments. Similar to other rate responses in this experiment, PANA was at a maximum at the intermediate rate of N under heat stress. However, in optimal temperatures, PANA was relatively constant at the two lowest N rates (41 to 47 mg plant⁻¹), but was much greater (84 mg plant⁻¹) at the highest N rate. Again, despite the differences in effects of fertility treatments in the two environments, PANA of the two genotypes responded similarly to fertility treatment across environments.

Under the heat stress environment (Exp 1), fertility treatment interactions (across genotypes) were significant for partitioning of all minerals (except Cu) to seed (Online Resource 2), with NPQ increasing (mineral utilization efficiency increasing) as N rate increased. As exceptions, partitioning of Mn, Mo, and Cd decreased as N rate increased. The two genotypes responded similarly, except for S and Zn. Under the optimal temperature environment, partitioning of minerals to grain (across genotypes) increased with increasing N only for Mn, while partitioning of S was negatively correlated with N rate. There were significant quadratic effects of fertility on partitioning of K, Fe, and Cu. In the *TaNAM*-RNAi line, as N increased mineral partitioning of Mg, Ca, Mn, Fe, Cu, Zn, and Cd tended to remain constant or decrease, while in the WT mineral partitioning was lowest at low N and was highest at the intermediate N rate.

We observed decreased per tiller contents of Cu, Fe, P, and Zn at maturity, indicating remobilization, in certain conditions. In the fertility study, in the heat stress environment both *TaNAM*-RNAi and WT plants remobilized Cu only under high N supply (Figure 7a), Fe accumulated in low N *TaNAM*-RNAi plants and was remobilized in high and low N WT plants, and Zn was remobilized in high N WT plants. However, under optimal temperatures (Figs. 7b and d), both WT and RNAi plants remobilized Fe, P, and Zn in almost all conditions, however, the *TaNAM*-RNAi line plants did not remobilize Cu, and WT plants did not remobilize Cu under water limitation.

Table 2. Effect of genotype, *TaNAM-RNAi* or Wild-type (WT), on N related traits in fertility studies conducted under heat stress or optimal temperature environments. Means are presented as the mean $\pm$ standard error. The F-value was determined by analysis of variance

Genotype	Vegetative Tissue N content (mg · plant ⁻¹)		N Remobilization (mg · plant ⁻¹) ^a , Efficiency (%)	Seed protein concentration (g · 100 g ⁻¹)	Seed N content (mg · plant ⁻¹)	Whole plant N Content (mg · plant ⁻¹)	N Harvest Index	N Partition
	Anthesis	Maturity						
Heat stress environment								
TaNAM-RNAi	29.8±1.3	24.0±0.8	5.8±1.5, 19.4 %	14.2±0.3	18.9±0.9	42.9±1.4	0.440±0.012	89.6±1.7
WT	27.9±1.3	16.2±0.8	11.8±1.5, 42.3 %	19.6±0.3	22.4±0.9	38.7±1.4	0.580±0.012	108.7±1.6
F-value								
Genotype	ns	41.9**	7.9**	149.6**	8.3**	ns	67.8**	64.3**
Geno X Stage	7.9**							
Fertility Trt	ns	18.1**(2)	9.0**	ns	8.7**(2)	19.1**(2)	13.1**(1)	4.9*(2)
Geno X Fert	ns	ns	ns	ns	5.23*(2)	5.7*(2)	ns	ns
Optimal temperature environment								
TaNAM-RNAi	76.9±4.5	54.1±2.7	27.2±1.5, 59.1 %	11.5±0.3	77.8±4.3	132±6	0.598±0.013	92.2±1.5
WT	62.9±4.5	28.6±2.8	31.4±1.6, 74.9 %	16.2±0.3	91.0±4.6	120±6	0.763±0.014	109.6±1.7
F-value								
Genotype	4.7*	52.8**	ns	88.8**	ns	4.4*	67.5***	65.4**
Geno X Stage	4.5*							
Fertility Trt	7.9**(2)	29.7**(1)	ns	21.4**(1)	16.1**(1)	28.7**(1)	ns	7.6**(2)
Geno X Fert	ns	4.4*(1)	ns	ns	4.2*(1)	ns	ns	ns

The F-value was determined by analysis of variance. F-value of highest order response presented. Values in parentheses after significant effects of fertility treatment indicate the highest order of the response (1 = linear; 2 = quadratic) to fertility treatment

a. calculated on a per-tiller basis

* and ** indicate significance at $p < 0.05$ and 0.01, respectively; ns = non-significant at $\alpha = 0.05$

Figure 4. Grain protein concentration (GPC) of grain grown in heat stress and optimal temperature environments in fertility (a) and moisture stress (b) studies

Moisture stress

In general, *TaNAM*-RNAi and WT plants responded similarly to post-anthesis moisture stress. The RNAi line accumulated greater vegetative DW than WT, but grain yields were not different between the genotypes. Therefore yield harvest index was greater in WT plants than *TaNAM*-RNAi plants in the heat stress environment (Table 4). Across genotypes, moisture stress significantly reduced vegetative biomass in both temperature regimes. Total grain yield and seed number were decreased by moisture stress only under optimal temperatures, while seed size was reduced by moisture deficit only in the heat stress environment.

Total plant N content decreased with moisture deficit in both environments, and genotypes

responded similarly to moisture deficit (Table 5). Under optimal temperatures, seed protein concentration of *TaNAM*-RNAi grain under moisture stress and well-watered treatments was 12.5 % and 12.0 %, respectively, while seed protein concentration of WT grain under moisture stress and well-watered conditions was 15.2 % and 17.0 %, respectively (Table 5, Figure 4). Across genotypes, moisture deficit reduced NHI and normalized partitioning of N to grain (NPQ) in the heat stress environment, but did not affect NPQ or NHI under optimal temperatures.

Mineral concentrations of seeds generally were decreased by moisture deficit in both environments (Online Resource 3). Seed mineral concentrations of the two genotypes generally responded similarly to water limitation. Similar to the fertility studies, WT plants utilized minerals more efficiently than *TaNAM*-RNAi plants: WT plants had mineral NPQ > 100, while *TaNAM*-RNAi plants generally had NPQ < 100 (Online Resource 4). In both heat stress and optimal temperature environments K was an exception, and had higher relative partitioning to grain in *TaNAM*-RNAi than in WT. In the heat stress environment, WT and RNAi lines responded differently, in terms of NPQ, to moisture stress for Mg, P, S, K, Ca, Fe, and Cd, while under optimal temperatures genotypes responded differently only for S, K, and Cu.

Discussion

Senescence and yield under optimal and stress conditions

Field-grown wheat in temperate climates commonly experiences stressful conditions during the grain filling period, including temperature stress, moisture stress, and nutrient deficiency. Climate change is expected to increase the frequency and severity of heat stress and drought events (Ahuja et al. 2010; Long and Ort 2010). Avoidance of senescence under stress (stay-green) has been associated with maintained yield in grain crops (Harris et al. 2007; Snape et al. 2007; Thomas and Howarth 2000; Zheng et al. 2009). Because the *TaNAM*-RNAi line exhibited delayed senescence in previous studies, we hypothesized that this line may also have delayed senescence under stress conditions, which would potentially result in improved yield under stress. Senescence of

Table 3. Effect of genotype, *Ta*NAM-RNAi or Wild-type, on mineral concentrations in seeds of plants grown in greenhouse fertility studies conducted under heat stress and optimal temperature environments. Data are means over fertility treatments. ANOVA was used to test significance of genotype effects

	Concentration in Seed at Maturity			
	Heat stress environment		Optimal temperature environment	
	<i>Ta</i> NAM-RNAi	Wild-type	<i>Ta</i> NAM-RNAi	Wild-type
Mg	1260±30	1470±30**	1520±35	1586±37
P	3040±100	3880±100**	3791±107	4017±115
S	1520±40	1830±40**	2084±44	2291±48**
K	4900±80	4300±70**	7105±137	5647±147**
Ca	501±10	497±10	685±18	648±19
Mn	42.0±1.5	51.6±1.4**	50.8±1.9	54.8±2.0
Fe	32.5±1.7	48.5±1.6**	34.4±1.3	43.7±1.4**
Cu	3.80±0.14	4.84±0.14**	3.75±0.33	4.91±0.36*
Zn	45.6±2.2	56.2±2.1**	34.9±1.5	36.8±1.6
Mo	0.834±0.046	1.264±0.045**	0.825±0.035	1.123±0.037**
Cd	0.148±0.008	0.216±0.008**	0.201±0.013	0.257±0.014**

* and ** indicate that F-value for genotype effect is significant at $p < 0.05$ and 0.01 , respectively

*Ta*NAM-RNAi plants was significantly delayed relative to WT plants in all optimal and stress conditions, including low N, heat stress and water limitation (Figure 2), suggesting that triggering of stress induced senescence may rely at least partially on elements of the *NAM* gene regulatory network. Moisture deficit in the heat stress environment substantially accelerated senescence in WT plants, which had lost >50 % chlorophyll concentration when well-watered WT plants were just beginning to senesce. *Ta*NAM-RNAi plants also senesced earlier under the heat stress environment with moisture deficit compared to when well-watered, but the effect of moisture stress was modest relative to WT. However, the delayed senescence trait did not provide seed yield protection against post-anthesis moisture deficit (Tables 1 and 4). Under well-watered conditions, there

was a yield advantage for the *Ta*NAM-RNAi plants, as *Ta*NAM-RNAi plants produced more and heavier seeds than the WT. A previous study also observed increased seed numbers in *Ta*NAM-RNAi plants (Waters et al. 2009). The yield HI of WT, however, was greater than *Ta*NAM-RNAi in well-watered plants under both heat stress and optimal temperature environments (Exp 1 and 2, Table 1), and under moisture stress in the heat stress environment (Exp 1, Table 4).

How does stress affect N utilization and GPC?

Previous studies have reported higher wheat GPC under heat and/or moisture stress conditions (Correll et al. 1994; Ercoli et al. 2008, 2010; Gooding et al. 2003). In our experiments, the WT line

Figure 5. Normalized mineral partitioning quotients for *Ta*NAM-RNAi and WT wheat plants from heat stress (HS) and optimal temperature environments in the fertility study. *, ** indicate that F-value is significant between genotypes at $p < 0.05$ and 0.01 , respectively

Table 4. Effect of genotype, *TaNAM-RNAi* or wild-type (WT), and moisture availability on vegetative biomass and yield components in two greenhouse studies conducted under heat stress and optimal temperature environments. Means are presented as the mean $\pm$ standard error

Genotype	Vegetative dry weight (g plant ⁻¹)	Tiller number	Dwt per tiller (g plant ⁻¹)	Seed number	Seeds per spike	Kernel wt (mg kernel ⁻¹)	Seed weight (g plant ⁻¹)	Seed yield (g spike ⁻¹)	Harvest index
Heat stress environment									
<i>TaNAM-RNAi</i>	1.30 $\pm$ 0.04	1.0	1.30 $\pm$ 0.04	24.4 $\pm$ 1.3	24.4 $\pm$ 1.3	34.4 $\pm$ 0.6	0.84 $\pm$ 0.04	0.84 $\pm$ 0.04	0.393 $\pm$ 0.011
WT	0.94 $\pm$ 0.04	1.0	0.94 $\pm$ 0.04	23.1 $\pm$ 1.3	23.1 $\pm$ 1.3	33.8 $\pm$ 0.6	0.77 $\pm$ 0.04	0.77 $\pm$ 0.04	0.450 $\pm$ 0.011
F-value (genotype)	42.1***		42.1***	0.5 ns	0.5 ns	0.5 ns	1.3 ns	1.3 ns	14.6**
Moisture stress	1.00 $\pm$ 0.04	1.0	1.00 $\pm$ 0.04	23.9 $\pm$ 1.4	23.9 $\pm$ 1.4	32.9 $\pm$ 0.6	0.78 $\pm$ 0.04	0.78 $\pm$ 0.04	0.439 $\pm$ 0.011
Well watered	1.23 $\pm$ 0.04	1.0	1.23 $\pm$ 0.04	23.6 $\pm$ 1.3	23.6 $\pm$ 1.3	35.4 $\pm$ 0.6	0.83 $\pm$ 0.04	0.83 $\pm$ 0.04	0.414 $\pm$ 0.011
F-value (moisture)	16.8***		16.8***	0.0 ns	0.0 ns	8.3**	0.9 ns	0.9 ns	5.2*
F-value (geno x moisture)	0.0 ns		0.0 ns	0.1 ns	0.1 ns	2.7 ns	0.9 ns	0.9 ns	0.5 ns
Optimal temperature environment									
<i>TaNAM-RNAi</i>	3.77 $\pm$ 0.16	2.68 $\pm$ 0.14	1.44 $\pm$ 0.05	82.0 $\pm$ 4.5	31.2 $\pm$ 1.4	35.9 $\pm$ 0.7	3.01 $\pm$ 0.16	1.15 $\pm$ 0.06	0.443 $\pm$ 0.012
WT	2.90 $\pm$ 0.16	2.62 $\pm$ 0.14	1.18 $\pm$ 0.05	73.9 $\pm$ 4.4	29.3 $\pm$ 1.4	35.0 $\pm$ 0.7	2.65 $\pm$ 0.15	1.07 $\pm$ 0.06	0.470 $\pm$ 0.012
F-value (genotype)	14.4***	0.8 ns	12.2**	1.7 ns	0.9 ns	1.0 ns	2.7 ns	1.0 ns	2.7 ns
Moisture stress	2.81 $\pm$ 0.16	2.23 $\pm$ 0.13	1.33 $\pm$ 0.05	63.9 $\pm$ 4.3	29.9 $\pm$ 1.4	35.3 $\pm$ 0.7	2.28 $\pm$ 0.15	1.09 $\pm$ 0.06	0.444 $\pm$ 0.011
Well watered	3.86 $\pm$ 0.17	3.06 $\pm$ 0.14	1.29 $\pm$ 0.05	92.0 $\pm$ 4.6	30.6 $\pm$ 1.4	35.6 $\pm$ 0.7	3.38 $\pm$ 0.16	1.13 $\pm$ 0.06	0.468 $\pm$ 0.012
F-value (moisture)	21.0***	17.8**	0.3 ns	20.0***	0.1 ns	0.1 ns	25.1***	0.2 ns	2.2 ns
F-value (geno x moisture)	8.9**	0.9 ns	15.5**	0.9 ns	3.2 ns	3.4 ns	2.5 ns	5.2*	1.1 ns

* and ** indicate that F-value is significant at $p < 0.05$ and 0.01 , respectively

Table 5. Effect of genotype, *TaNAM*-RNAi or Wild-type (WT), and moisture availability on plant N related traits in two greenhouse studies conducted under heat stress (and optimal temperature environments. Means are presented as the mean $\pm$ standard error

Genotype	N Remobilization (mg $\cdot$ plant ⁻¹) ^a , Efficiency (%)	Seed protein conc (%)	Seed N content (mg plant ⁻¹)	Whole plant N content (mg plant ⁻¹)	N Harvest Index	N Partition
<i>Heat stress environment</i>						
<i>TaNAM</i> -RNAi	4.7 $\pm$ 1.9, 15.9 %	13.2 $\pm$ 0.6	19.4 $\pm$ 1.5	44.3 $\pm$ 1.9	0.440 $\pm$ 0.018	85.9 $\pm$ 2.3
WT	20.7 $\pm$ 1.9, 63.3 %	18.0 $\pm$ 0.6	24.2 $\pm$ 1.5	36.2 $\pm$ 2.0	0.673 $\pm$ 0.019	115.7 $\pm$ 2.4
F-value (genotype)	36.7**	31.3**	5.2*	8.8**	81.7***	82.4**
Moisture stress	16.9 $\pm$ 1.9, 54.3 %	15.3 $\pm$ 0.6	20.5 $\pm$ 1.5	34.7 $\pm$ 2.0	0.602 $\pm$ 0.019	105.0 $\pm$ 2.4
Well watered	8.5 $\pm$ 1.8, 27.3 %	15.9 $\pm$ 0.6	23.2 $\pm$ 1.5	45.9 $\pm$ 1.9	0.511 $\pm$ 0.017	96.6 $\pm$ 2.2
F-value (moisture)	10.4**	0.5 ns	1.7 ns	16.6***	12.6**	6.6*
F-value (geno x moisture)	0.4 ns	0.1 ns	0.6 ns	0.0 ns	0.1 ns	0.5 ns
<i>Optimal temperature environment</i>						
<i>TaNAM</i> -RNAi	29.6 $\pm$ 1.6, 64.2 %	12.2 $\pm$ 0.4	64.4 $\pm$ 4.1	108 $\pm$ 5.2	0.598 $\pm$ 0.019	91.4 $\pm$ 2.8
WT	33.3 $\pm$ 1.6, 74.2 %	16.0 $\pm$ 0.3	75.1 $\pm$ 4.0	103 $\pm$ 5.0	0.707 $\pm$ 0.018	108.1 $\pm$ 2.7
F-value (genotype)	2.7 ns	61.3**	2.9 ns	0.5 ns	17.9***	17.9***
Moisture stress	31.9 $\pm$ 1.6, 70.0 %	13.8 $\pm$ 0.3	55.0 $\pm$ 4.0	83.5 $\pm$ 5.0	0.647 $\pm$ 0.018	98.9 $\pm$ 2.7
Well watered	31.0 $\pm$ 1.6, 68.0 %	14.5 $\pm$ 0.4	83.6 $\pm$ 4.1	127.6 $\pm$ 5.3	0.659 $\pm$ 0.019	100.7 $\pm$ 2.9
F-value (moisture)	0.1 ns	2.0 ns	25.6**	36.8***	0.2 ns	0.2 ns
F-value (geno x moisture)	2.5 ns	5.3*	0.0 ns	1.6 ns	2.9 ns	2.9 ns

a. calculated on a per-tiller basis

* and ** indicate that F-value is significant at $p < 0.05$ and 0.01 , respectively

consistently had higher GPC than the *TaNAM*-RNAi line (Tables 2 and 5, Figure 4), consistent with previous studies (Uauy et al. 2006b; Waters et al. 2009). The exceptionally high seed protein concentrations ($>19\%$) of WT seeds from the heat stress environment (Table 2, Figure 4) suggests that the second-order effects observed may be a consequence of excess fertility application at the limited yield potential of the plants. The lower GPC in the *TaNAM*-RNAi line was not due to decreased pre-anthesis accumulation of N (Tables 2 and 5), which was greater in the *TaNAM*-RNAi line or equivalent between the two lines. For N, heat stress has been shown to increase (Corbellini et al. 1997) or reduce remobilization (Ercoli et al. 2010), while drought stress reduced remobilization (Ercoli et al. 2008). Here, we observed that plants grown in the temperature stress environment had reduced N remobilization efficiency in both fertility and moisture stress studies. Part of the decrease in GPC in *TaNAM*-RNAi grain relative to WT was attributed to decreased remobilization in prior work (Waters et al. 2009), and our results here are consistent with this hypothesis (Tables 2 and 5). While PANA was equivalent between WT and *TaNAM*-RNAi lines (Figure 6), NHI and NPQ were diminished in the *TaNAM*-RNAi line, indicating

that the *TaNAM*-RNAi line less effectively remobilized vegetative N and translocated new N from vegetative tissues to grain. Under the high temperature environment, plants that experienced moisture stress had greater NHI and NPQ than well-watered plants, but this relationship did not hold

Figure 6. Post-anthesis N accumulation (PANA) in above-ground tissues of *TaNAM*-RNAi and WT wheat plants from heat stress and optimal temperature environments in the fertility study. There were no significant differences between genotypes

Figure 7. Remobilization of Cu, Fe, P and Zn from above-ground tissues of *TaNAM*-RNAi and WT wheat plants. a and b, fertility study; c and d, irrigation study. a and c, plants grown under heat stress conditions; b and d, plants grown under optimal temperatures. Negative values indicate net remobilization from vegetative tissues between anthesis and maturity. Asterisk indicates that F-value for mineral content change is significant, positive indicates that F-value for mineral content change is significant between genotypes at $p < 0.05$

for plants in the optimal temperature environment (Table 5). At this point, it is not clear which specific gene products are needed to carry out partitioning, but it is clear that they are associated with senescence. Accompanying the delayed senescence in the *TaNAM*-RNAi line is differential regulation of several hundred genes (Cantu et al. 2011), which are not properly regulated in the *TaNAM*-RNAi line; it is likely that one or more of these genes are responsible for N partitioning to grain.

How does stress affect mineral partitioning?

Remobilization patterns of Cu, Fe, and Zn were altered in the heat stress environment (Figure 7). Optimal temperatures produced more consistency between N treatments. WT plants remobilized more Cu and Fe than the *TaNAM*-RNAi line in fertility studies under optimal temperatures, but P and Zn

remobilization was similar. The effects of moisture stress on mineral concentrations of wheat grain are not well defined. Water limitation significantly affected grain ash concentration, with less ash at more severe water stress, and Mg concentration also was affected by drought, but not in a linear fashion (Zhu et al. 2008). In a study of 22 accessions of *T. dicoccoides*, well-watered or water limited conditions produced differences in grain Fe and Zn, but some accessions had greater concentrations under water limitation while others had the opposite response to water limitation (Peleg et al. 2008). Collections of *T. spelta* and *T. dicoccoides* grown under different environments revealed that environmental effects were the greatest factor in grain mineral concentrations (Gomez-Becerra et al. 2010a, b). Here, moisture stress in the heat stress environment revealed differences in remobilization of not only Fe, but also P and Zn between genotypes (Figure 7). Similarly,

in the fertility study, remobilization of minerals affected the RNAi line greater than the WT in the heat stress environment. Thus, it appears likely that the lower remobilization capacity of the RNAi line is magnified by heat stress during the grain fill period. The greater remobilization in the WT certainly would have contributed to differences in partitioning between vegetative tissues and grain. Similar to N, partitioning quotients for most minerals were lower in the RNAi line than in wild-type (Online Resource 4), suggesting that similar processes are used to partition N and minerals to grain during grain fill, and that knocking down *NAM* genes by RNAi influences these processes. Although we cannot directly compare the samples from the two environments, normalized PQ appeared to be largely unchanged between the heat stress environment and optimal temperature environment (Figure 5). This is consistent with a previous study that showed decreased translocation of newly taken-up radiolabeled Zn to grain in the *TaNAM*-RNAi line (Waters et al. 2009). A notable exception to this relationship was partitioning of K to grain, which was significantly greater in the *TaNAM*-RNAi line than in the WT (Online Resource 4, Figure 5). Across genotypes, moisture stress altered NPQ of several minerals under both optimal and heat stress temperature environments, but there were few consistent patterns between the two environments (online Resource 4). Nutrient harvest indices of *TaNAM*-RNAi and WT plants generally responded similarly to heat and moisture stress. The heat stress environment tended to decrease mineral PQ in the fertility study (Online Resource 2, Figure 5).

TaNAM-RNAi plants were larger overall, but had smaller quantities of nutrients partitioned to the grain, with similar or higher grain yield. The end result of this is a more dilute grain mineral density, as reflected in grain mineral concentrations (Table 3, Online Resource 3). In the fertility study, in the heat stress environment all minerals except Ca were significantly different between genotypes, with all of these being lower in the *TaNAM*-RNAi line except for K, which was higher here and was higher in *TaNAM*-RNAi grain in all other conditions as well (Online Resource 3). Under the optimal temperature regime in both the fertility and moisture stress studies, most mineral concentrations were greater in WT grain. These results are mostly consistent with results of previous evaluations of *TaNAM*-RNAi

wheat (Uauy et al. 2006b; Waters et al. 2009) and extend the observations to stress conditions. However, in the previous studies, Zn concentration in *TaNAM*-RNAi seed was lower than in WT seed, while here the concentration of Zn in seed of *TaNAM*-RNAi plants was significantly lower only in the heat stress environment. When genotypes expressing the non-functional *Gpc-B1* allele were compared with genotypes expressing the functional allele (Distelfeld et al. 2007), only the concentrations of Fe, Mg, and Zn were significantly different. The remobilization of Zn from vegetative tissues was previously shown to be lower in the RNAi line (Waters et al. 2009), whereas here no differences were observed. The soil-sand mix employed in this study did not receive additional Zn fertilization, and soil Zn concentration may have been too low to allow expression of differences previously observed.

Several studies have shown that increasing N supply positively affected Fe and/or Zn uptake and grain Fe and/or Zn concentrations (Aciksoz et al. 2011; Cakmak et al. 2010; Erenoglu et al. 2010; Gooding et al. 2012; Kutman et al. 2010, 2011a, 2012; b; Shi et al. 2010). In our fertility study, grain Fe was consistently higher in WT than in RNAi, but Zn was higher only under the heat stress environment (Table 3), and we did not observe increasing grain Fe and Zn with increasing N supply. Our results did show, however, that across genotypes, increasing N from the lowest rate resulted in improved nutrient use efficiency (NPQ) for both Fe and Zn in the heat stress fertility study, and increased nutrient use efficiency of Fe under optimal temperatures. The *TaNAM*-RNAi and WT genotypes responded differently to N supply for Zn PQ in both heat stress and optimal temperature environment, and for Fe PQ under optimal temperatures (Online Resource 2). However, in regard to grain mineral concentrations, in general, both genotypes responded similarly to moisture stress, and relative mineral concentrations between the genotypes were maintained in both optimal temperature and heat stress environments (Online resource 3). Thus, the delayed senescence of the RNAi line did not provide an advantage in terms of grain mineral concentrations.

Conclusions and future directions

A previous study showed that *NAM* genes were upregulated rapidly after anthesis in flag leaves.

Here we show that the primary site of expression, at least in above-ground tissues, is the flag leaf, with some expression in senescing lower leaves (Figure 1). An unanswered question is how expression in flag leaf also controls senescence of peduncle and spike, which remained green in *TaNAM*-RNAi plants in previous studies and also this one. Nutritional quality, in terms of mineral and protein concentration, was decreased in the *TaNAM*-RNAi grain here and in the prior studies. Part of this decrease results from reduced net remobilization of minerals and part results from lower partitioning of minerals taken up during grain fill and directed to the grain as it is filling. Dry matter partitioning to grain was affected less than N and mineral partitioning to grain. Thus, another unanswered question is how decreased expression of *NAM* genes affects the translocation of N and minerals to grain. The pathways of N and minerals to grain are not fully defined (Waters and Sankaran 2011), and specific sites of phloem loading for transport to the grain are not known, although glumes, stem, roots, and leaves are all sources (Simpson et al. 1983). It is possible that N and minerals first move into the flag leaf to be directed into translocation pathways to the grain, which would make it easier to explain the relationship between the site of *NAM* expression and the effects of *NAM* knockdown. Since the *NAM* genes are transcription factors that potentially regulate a large number of direct target and downstream genes, in the future it may be possible to separate the effects on senescence onset from those on partitioning and grain nutritional quality. Transcriptome analysis identified 691 differentially regulated genes during senescence between the WT and *TaNAM*-RNAi (Cantu et al. 2011). It is unlikely that the *NAM* proteins directly regulate all of these genes, and some of them are probably part of secondary networks. Thus, increased knowledge of direct *NAM* targets and their downstream networks could enable targeting of specific components that may permit selection for the beneficial effects of delayed senescence while avoiding the detrimental effects on nutritional quality.

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Supplemental resource 1. Effect of genotype, *TaNAM-RNAi* or Wild-type (WT), on mineral concentrations in vegetative tissue and contents in vegetative tissue and seeds of plants grown in two greenhouse fertility studies. Data are means over fertility treatments. ANOVA was used to test significance of genotype effects.

	Concentration in Vegetative Tissue at Anthesis ($\mu\text{g g}^{-1}$)			
	Heat Stress Environment		Optimal Temperature Environment	
Mineral	<i>TaNAM-RNAi</i>	WT	<i>TaNAM-RNAi</i>	WT
Mg	2320 $\pm$ 60	2370 $\pm$ 60	2145 $\pm$ 56	2007 $\pm$ 56
P	1480 $\pm$ 34.3	1550 $\pm$ 34.3	2708 $\pm$ 84	2703 $\pm$ 84
S	2330 $\pm$ 110	2380 $\pm$ 110	3259 $\pm$ 99	3000 $\pm$ 99
K	31000 $\pm$ 520	31400 $\pm$ 520	41807 $\pm$ 882	39965 $\pm$ 882
Ca	5390 $\pm$ 140	5590 $\pm$ 140	5990 $\pm$ 205	5212 $\pm$ 205
Mn	63.7 $\pm$ 2.6	62.1 $\pm$ 2.6	87.2 $\pm$ 3.2	80.5 $\pm$ 3.2
Fe	53.5 $\pm$ 5.5	68.6 $\pm$ 5.5	56.3 $\pm$ 1.7	58.7 $\pm$ 1.7
Cu	9.05 $\pm$ 1.07	8.79 $\pm$ 1.07	5.44 $\pm$ 0.25	4.82 $\pm$ 0.25
Zn	23.0 $\pm$ 1.22	21.4 $\pm$ 1.22	27.2 $\pm$ 1.1	24.1 $\pm$ 1.1
Mo	1.96 $\pm$ 0.08	2.03 $\pm$ 0.08	0.772 $\pm$ 0.086	0.917 $\pm$ 0.086
Cd	0.310 $\pm$ 0.015	0.318 $\pm$ 0.015	0.371 $\pm$ 0.030	0.388 $\pm$ 0.030
	Concentration in Vegetative Tissue at Maturity ($\mu\text{g g}^{-1}$)			
	Heat Stress Environment		Optimal Temperature Environment	
	<i>TaNAM-RNAi</i>	WT	<i>TaNAM-RNAi</i>	WT
Mg	3130 $\pm$ 80	2970 $\pm$ 70	2217 $\pm$ 65	2038 $\pm$ 70
P	1240 $\pm$ 70	1330 $\pm$ 70	1148 $\pm$ 72	789 $\pm$ 78**
S	4760 $\pm$ 150	4140 $\pm$ 140**	3682 $\pm$ 108	3124 $\pm$ 116**
K	24000 $\pm$ 800	25000 $\pm$ 800	33438 $\pm$ 879	38860 $\pm$ 942**
Ca	8280 $\pm$ 260	7610 $\pm$ 250	6686 $\pm$ 250	6571 $\pm$ 270
Mn	90.4 $\pm$ 5.0	94.1 $\pm$ 4.8	111 $\pm$ 6	99.3 $\pm$ 6.0
Fe	37.6 $\pm$ 1.1	36.0 $\pm$ 1.0	54.7 $\pm$ 13.2	56.7 $\pm$ 14.2
Cu	6.91 $\pm$ 0.20	6.99 $\pm$ 0.19	6.54 $\pm$ 0.26	4.60 $\pm$ 0.28**
Zn	17.9 $\pm$ 1.0	14.8 $\pm$ 0.9	11.1 $\pm$ 0.8	7.1 $\pm$ 0.8**
Mo	2.82 $\pm$ 0.13	3.11 $\pm$ 0.12	1.95 $\pm$ 0.08	1.71 $\pm$ 0.09*
Cd	0.442 $\pm$ 0.022	0.481 $\pm$ 0.021	0.555 $\pm$ 0.032	0.480 $\pm$ 0.034

Supplemental resource 1, continued

Content in Vegetative Tissue at Anthesis ($\mu\text{g plant}^{-1}$)				
Heat Stress Environment		Optimal Temperature Environment		
Mineral	<i>TaNAM-RNAi</i>	WT	<i>TaNAM-RNAi</i>	WT
Mg	2190 $\pm$ 120	1990 $\pm$ 120	6182 $\pm$ 336	4651 $\pm$ 336 **
P	1400 $\pm$ 70	1290 $\pm$ 70	7925 $\pm$ 533	6314 $\pm$ 533 *
S	2210 $\pm$ 140	2000 $\pm$ 140	9383 $\pm$ 496	6970 $\pm$ 496**
K	29200 $\pm$ 1300	26200 $\pm$ 1300	120590 $\pm$ 4690	93030 $\pm$ 4390 **
Ca	5080 $\pm$ 280	4700 $\pm$ 280	17318 $\pm$ 934	11976 $\pm$ 934**
Mn	60.2 $\pm$ 3.6	52.0 $\pm$ 3.6	251 $\pm$ 13	185 $\pm$ 13**
Fe	50.1 $\pm$ 4.6	57.5 $\pm$ 4.6	162 $\pm$ 9	136 $\pm$ 9
Cu	8.31 $\pm$ 0.94	7.40 $\pm$ 0.94	16.0 $\pm$ 1.2	11.2 $\pm$ 1.2 **
Zn	21.6 $\pm$ 1.5	18.2 $\pm$ 1.5	78.8 $\pm$ 4.9	56.1 $\pm$ 4.9**
Mo	1.88 $\pm$ 0.12	1.68 $\pm$ 0.11	2.22 $\pm$ 0.26	2.09 $\pm$ 0.26
Cd	0.290 $\pm$ 0.020	0.269 $\pm$ 0.020	1.070 $\pm$ 0.075	0.875 $\pm$ 0.075
Content in Vegetative Tissue at Maturity ($\mu\text{g plant}^{-1}$)				
Heat Stress Environment		Optimal Temperature Environment		
	<i>TaNAM-RNAi</i>	WT	<i>TaNAM-RNAi</i>	WT
Mg	4150 $\pm$ 155	2880 $\pm$ 149**	10137 $\pm$ 349	6627 $\pm$ 374**
P	1640 $\pm$ 85	1260 $\pm$ 82**	5472 $\pm$ 390	2739 $\pm$ 419**
S	6300 $\pm$ 240	3950 $\pm$ 233**	17206 $\pm$ 676	10349 $\pm$ 726**
K	31400 $\pm$ 1240	24000 $\pm$ 1220**	154883 $\pm$ 4605	129110 $\pm$ 4605**
Ca	10900 $\pm$ 417	7330 $\pm$ 403**	30541 $\pm$ 1201	21432 $\pm$ 1277**
Mn	117 $\pm$ 7	92.5 $\pm$ 6.3**	504 $\pm$ 26	323 $\pm$ 28**
Fe	49.8 $\pm$ 1.8	34.6 $\pm$ 1.7**	257 $\pm$ 42	178 $\pm$ 45
Cu	9.21 $\pm$ 0.39	6.78 $\pm$ 0.38**	30.3 $\pm$ 1.4	15.1 $\pm$ 1.5**
Zn	20.4 $\pm$ 1.0	14.0 $\pm$ 0.9**	50.4 $\pm$ 3.5	24.5 $\pm$ 3.8**
Mo	3.73 $\pm$ 0.20	3.08 $\pm$ 0.20*	9.00 $\pm$ 0.43	5.59 $\pm$ 0.46**
Cd	0.621 $\pm$ 0.029	0.465 $\pm$ 0.028**	2.50 $\pm$ 0.14	1.57 $\pm$ 0.14**
Content in Seed at Maturity ($\mu\text{g plant}^{-1}$)				
Heat Stress Environment		Optimal Temperature Environment		
	<i>TaNAM-RNAi</i>	WT	<i>TaNAM-RNAi</i>	WT
Mg	955 $\pm$ 40	935 $\pm$ 38	5924 $\pm$ 278	4935 $\pm$ 298**
P	2300 $\pm$ 107	2450 $\pm$ 103	14878 $\pm$ 724	12363 $\pm$ 777**
S	1150 $\pm$ 52	1180 $\pm$ 51	8053 $\pm$ 381	7055 $\pm$ 409*
K	3720 $\pm$ 148	2790 $\pm$ 144**	27802 $\pm$ 1215	17637 $\pm$ 1292**
Ca	387 $\pm$ 18	322 $\pm$ 17*	2661 $\pm$ 131	2050 $\pm$ 140**
Mn	31.9 $\pm$ 1.5	32.4 $\pm$ 1.4	195 $\pm$ 10	172 $\pm$ 11
Fe	24.4 $\pm$ 1.5	30.7 $\pm$ 1.4**	134 $\pm$ 8	137 $\pm$ 8
Cu	2.87 $\pm$ 0.14	3.07 $\pm$ 0.14	14.4 $\pm$ 1.4	15.1 $\pm$ 1.5
Zn	34.2 $\pm$ 1.7	35.3 $\pm$ 1.7	131 $\pm$ 6	113 $\pm$ 7*
Mo	0.635 $\pm$ 0.042	0.802 $\pm$ 0.040**	3.19 $\pm$ 0.21	3.45 $\pm$ 0.22
Cd	0.114 $\pm$ 0.008	0.140 $\pm$ 0.008*	0.751 $\pm$ 0.053	0.792 $\pm$ 0.057

*, ** indicate that F-value for genotype effect is significant at $p < 0.05$ and 0.01 , respectively

Supplemental resource 2. Normalized partitioning coefficient for minerals in two greenhouse fertility studies.

Heat Stress Environment, Fertility		Normalized Partitioning Coefficient										
Genotype	Fertility (mg N plant ⁻¹)	Mg	P	S	K	Ca	Mn	Fe	Cu	Zn	Mo	Cd
	17.5	86.7	94.3	76.0	88.2	75.8	117.4	92.0	98.2	93.7	149.8	124.2
	61.3	96.3	97.4	88.7	98.5	91.0	102.9	95.7	99.7	100.0	96.3	100.4
	105	112.4	107.0	128.0	109.6	125.2	85.7	109.1	101.1	105.5	63.9	81.4
		** ⁽¹⁾ ^a	** ⁽¹⁾	* ⁽²⁾	** ⁽¹⁾	** ⁽¹⁾	** ⁽¹⁾	** ⁽¹⁾	ns	** ⁽¹⁾	** ⁽¹⁾	** ⁽¹⁾
RNAi	17.5	76.5	94.6	63.5	97.7	70.4	110.9	76.4	86.7	89.4	126.1	109.0
RNAi	61.3	88.2	98.4	81.7	99.5	88.2	104.2	85.2	91.1	103.4	85.4	85.7
RNAi	105	107.2	100.4	103.9	119.0	119.7	72.9	95.4	94.8	99.3	57.8	61.6
WT	17.5	96.8	94.0	88.5	78.7	81.3	123.9	107.7	109.7	98.1	173.6	139.4
WT	61.3	104.3	96.4	95.6	97.4	93.7	101.6	106.3	108.2	96.7	107.2	115.1
WT	105	117.5	113.6	152.1	100.2	130.7	98.6	122.8	107.3	111.6	70.0	101.2
		ns ^b	ns	* ⁽²⁾	ns	ns	ns	ns	ns	* ⁽²⁾	ns	ns

*, ** indicate that F-value for effect is significant at $p < 0.05$ and 0.01 , respectively; ns=non-significant

†Values in parentheses indicate order of response to fertilizer rate: 1=first-order (linear); 2=second-order (quadratic)

^aindicates response to N rate across genotypes

^bindicates whether response to N rate was different between genotypes

Supplemental Resource 2 continued

Optimal Temperature Environment, Fertility		Normalized Partitioning Coefficient										
Genotype	Fertility (mg N plant ⁻¹)	Mg	P	S	K	Ca	Mn	Fe	Cu	Zn	Mo	Cd
	17.5	97.1	101.0	108.8	101.6	96.5	90.7	92.5	98.7	97.2	101.7	99.2
	61.3	102.5	102.5	98.6	93.5	101.4	102.7	118.2	121.6	102.4	104.1	105.7
	105	102.5	97.8	93.5	103.1	104.7	110.4	100.6	89.5	101.7	98.6	100.0
		ns ^a	ns	** (1)†	* (2)	ns	** (1)	** (2)	** (2)	ns	ns	ns
RNAi	17.5	95.9	99.1	99.3	121.3	102.7	90.9	92.8	88.4	95.1	90.6	88.0
RNAi	61.3	92.5	97.2	85.3	101.7	90.7	87.5	86.4	89.8	95.5	83.6	80.5
RNAi	105	99.9	96.2	88.9	117.7	105.5	103.8	81.0	75.1	100.5	83.5	88.6
WT	17.5	98.3	102.8	118.4	81.9	90.4	90.5	92.1	109.0	99.3	112.7	110.3
WT	61.3	112.5	107.8	111.8	85.3	112.1	117.9	150.1	153.3	109.4	124.5	130.8
WT	105	105.2	99.4	98.1	88.4	103.8	116.9	120.3	103.8	103.0	113.8	111.4
		*(2) ^b	ns	ns	ns	** (2)	** (2)	** (2)	** (2)	*(2)	ns	*(2)

*, ** indicate that F-value for effect is significant at $p < 0.05$ and 0.01 , respectively; ns=non-significant

†Values in parentheses indicate order of response to fertilizer rate: 1=first-order (linear); 2=second-order (quadratic)

^aindicates response to N rate across genotypes

^bindicates whether response to N rate was different between genotypes

Supplemental resource 3. Effect of genotype, *TaNAM-RNAi* or Wild-type (WT), and water limitation on mineral concentrations seed in two greenhouse irrigation studies.

Heat Stress Environment, Irrigation		Concentration (ppm)											
		Mg	P	S	K	Ca	Mn	Fe	Cu	Zn	Se	Mo	Cd
RNAi		1311	3000	1809	5396	595	32.46	28.3	3.4	37.2	0.18	0.73	0.09
WT		1369	3521	2036	4434	505	37.49	42.3	4.1	43.1	0.28	1.07	0.14
		ns ^a	**	**	**	**	*	**	**	ns	**	**	**
	150	1191	2699	1792	4469	510	28.25	31.6	3.2	35.1	0.22	0.84	0.13
	250	1489	3822	2053	5362	590	41.70	39.1	4.3	45.2	0.24	0.96	0.11
		** ^b	**	**	**	**	**	**	**	**	ns	*	*
RNAi	150	1153	2454	1611	4664	548	26.05	25.8	2.9	33.0	0.17	0.71	0.11
RNAi	250	1469	3547	2007	6128	643	38.87	30.8	4.0	41.4	0.19	0.76	0.08
WT	150	1229	2945	1973	4273	473	30.45	37.3	3.5	37.1	0.27	0.98	0.16
WT	250	1509	4097	2099	4595	536	44.53	47.4	4.7	49.0	0.29	1.16	0.13
		ns ^c	ns	**	**	ns	ns	ns	ns	ns	ns	ns	ns

*, ** indicate that F-value for effect is significant at $p < 0.05$ and 0.01 , respectively; ns=non-significant

†Values in parentheses indicate order of response to fertilizer rate: 1=first-order (linear); 2=second-order (quadratic)

^aindicates whether concentration was different between genotypes across water treatments

^bindicates whether response to water treatment was significant across genotypes

^cindicates whether response to water treatment was different between genotypes

Supplemental resource 3, continued

Optimal Temperature Environment, Irrigation		Concentration (ppm)											
		Mg	P	S	K	Ca	Mn	Fe	Cu	Zn	Se	Mo	Cd
RNAi		1419	3143	1978	6659	680.3	40.09	30.08	2.705	29.43	0.193	0.722	0.137
WT		1544	3787	2345	5917	651.9	46.42	44.41	3.625	33.64	0.23	0.983	0.189
		* ^a	**	**	**	ns	**	**	**	ns	*	**	**
	150	1385	3258	2147	5985	605.4	38.05	39.37	3.525	34.19	0.218	0.842	0.155
	250	1578	3671	2175	6591	726.9	48.45	35.12	2.805	28.88	0.205	0.864	0.171
		** ^b	*	ns	**	**	**	ns	**	ns	ns	ns	ns
RNAi	150	1281	2757	1896	6057	614.7	34.94	30.8	7.892	31.61	0.195	0.712	0.137
RNAi	250	1557	3528	2059	7261	746	45.24	29.36	2.45	27.25	0.19	0.733	0.138
WT	150	1489	3759	2399	5913	596.1	41.17	47.94	4.088	36.76	0.24	0.972	0.173
WT	250	1599	3814	2290	5921	707.8	51.66	40.88	3.161	30.51	0.22	0.995	0.205
		ns ^c	ns	ns	**	ns	ns	ns	ns	ns	ns	ns	ns

*, ** indicate that F-value for effect is significant at $p < 0.05$ and 0.01 , respectively; ns=non-significant

†Values in parentheses indicate order of response to fertilizer rate: 1=first-order (linear); 2=second-order (quadratic)

^aindicates whether concentration was different between genotypes across water treatments

^bindicates whether response to water treatment was significant across genotypes

^cindicates whether response to water treatment was different between genotypes

Supplemental resource 4. Effect of genotype, *TaNAM-RNAi* or Wild-type (WT), and water limitation on seed mineral partitioning quotient in two greenhouse irrigation studies.

Heat Stress Environment, Irrigation	Normalized Partitioning Quotient											
	Mg	P	S	K	Ca	Mn	Fe	Cu	Zn	Se	Mo	Cd
RNAi	97.8	98.0	89.2	118.9	106.0	96.3	81.2	88.4	96.8	78.7	88.1	83.8
WT	103.7	102.8	115.3	80.4	95.6	105.8	120.9	113.8	103.8	124.2	115.8	120.4
	ns ^a	ns	**	**	ns	ns	**	*	ns	**	**	**
150	101.9	101.7	119.2	89.3	104.0	106.7	104.0	102.6	100.3	108.0	117.3	116.5
250	99.6	99.1	85.3	110.0	97.6	95.4	98.1	99.6	100.4	94.9	86.5	87.8
	ns ^b	ns	**	**	ns	ns	ns	ns	ns	ns	**	**
RNAi 150	91.5	96.0	97.9	96.7	100.3	95.5	80.3	81.6	93.6	80.4	98.3	91.2
RNAi 250	104.1	100.1	80.6	141.1	111.7	97.2	82.0	95.2	100.0	77.0	77.8	76.3
WT 150	112.4	107.5	140.6	81.9	107.7	117.8	127.6	123.6	107.0	135.5	136.3	141.7
WT 250	95.1	98.2	90.1	78.9	83.5	93.7	114.2	103.9	100.7	112.9	95.3	99.2
	** ^c	*	*	**	**	ns	*	ns	ns	ns	ns	*

*, ** indicate that F-value for effect is significant at $p < 0.05$ and 0.01 , respectively; ns=non-significant

^a indicates whether normalized partitioning quotient was different between genotypes across water treatments

^b indicates whether response to water treatment was significant across genotypes

^c indicates whether response to water treatment was different between genotypes

Supplemental resource 4, continued

Optimal Temperature Environment, Irrigation	Normalized Partitioning Quotient											
	Mg	P	S	K	Ca	Mn	Fe	Cu	Zn	Se	Mo	Cd
RNAi	94.0	96.8	88.2	106.4	97.9	92.1	78.2	73.5	97.8	85.6	87.4	88.3
WT	106.4	102.6	110.6	95.0	103.0	107.5	121.3	125.4	101.7	113.1	111.5	111.7
	** ^a	**	**	*	ns	**	**	**	ns	**	**	**
150	94.3	102.7	103.6	92.6	91.2	99.4	93.3	95.5	102.7	102.8	101.8	94.4
250	106.0	96.8	95.2	108.7	109.7	100.1	106.2	103.4	96.9	95.9	97.2	105.6
	** ^b	**	**	**	**	ns	*	ns	ns	ns	ns	**
RNAi	150	84.5	98.8	89.1	92.9	86.0	91.0	70.8	83.1	100.0	90.6	93.4
RNAi	250	103.4	94.9	87.4	119.8	109.8	93.2	85.6	63.8	95.5	80.7	81.5
WT	150	104.2	106.6	118.1	92.3	96.4	107.8	115.7	107.8	105.3	115.1	110.3
WT	250	108.6	98.7	103.0	97.7	109.6	107.1	126.8	143.0	98.2	111.1	112.8
	ns ^c	ns	*	*	ns	ns	ns	**	ns	ns	ns	ns

*, ** indicate that F-value for effect is significant at $p < 0.05$ and 0.01 , respectively; ns=non-significant

^aindicates whether normalized partitioning quotient was different between genotypes across water treatments

^bindicates whether response to water treatment was significant across genotypes

^cindicates whether response to water treatment was different between genotypes