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# RNase Z<sup>S1</sup> processes *Ub<sub>L40</sub>* mRNAs and controls thermosensitive genic male sterility in rice

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Thermosensitive genic male-sterile (TGMS) lines, which are male-sterile at restrictive (high) temperatures but male-fertile at permissive (low) temperatures, have been widely used in breeding two-line hybrid rice (*Oryza sativa* L.). Here we find that mutation of *thermosensitive genic male sterile 5* (*tms5*) in rice causes the TGMS trait through a loss of RNase Z<sup>S1</sup> function. We show that RNase Z<sup>S1</sup> processes the mRNAs of three *ubiquitin fusion ribosomal protein L40* (*Ub<sub>L40</sub>*) genes into multiple fragments *in vitro* and *in vivo*. In *tms5* mutants, high temperature results in increased levels of *Ub<sub>L40</sub>* mRNAs. Overaccumulation of *Ub<sub>L40</sub>* mRNAs causes defective pollen production and male sterility. Our results uncover a novel mechanism of RNase Z<sup>S1</sup>-mediated *Ub<sub>L40</sub>* mRNA regulation and shows that loss of this regulation produces TGMS in rice, a finding with potential applications in hybrid crop breeding.

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Hybrid rice, which yields ~10–20% more grain than conventional rice, has been planted in ~60% (~17 million hectares) of the rice-growing area in China and is also cultivated in many other countries, thus providing a key component of the global food supply<sup>1–3</sup>. Rice is strictly self-pollinating; therefore, hybrid varieties are bred using male-sterile maternal lines that fail to produce viable pollen, thus preventing self-pollination. Hybrid rice breeding uses the well-developed three-line and two-line systems<sup>4,5</sup>. The three-line system uses a cytoplasmic male-sterile (CMS) line, a restorer line and a CMS maintainer line to produce F<sub>1</sub> hybrid seeds and maintain the CMS line<sup>6–8</sup>. Three-line hybrid rice has been grown since the 1970s and is a major type of hybrid rice<sup>1,4,9</sup>. However, the limited germplasm resources of restorer lines and the genetic diversity between CMS and restorer lines have limited further improvements in three-line breeding<sup>10</sup>. The two-line breeding system uses thermosensitive genic male sterility (TGMS) lines or photoperiod-sensitive genic male sterility (PGMS) lines as maternal parents to produce hybrid seeds. TGMS and PGMS lines are male-sterile under restrictive conditions (high temperatures for TGMS and long-day for PGMS) but convert to male-fertile under permissive conditions (low temperatures for TGMS and short-day for PGMS); thus, they can self-pollinate under permissive conditions<sup>10,11</sup>. The TGMS and PGMS traits are controlled by nuclear, recessive genes and most normal rice cultivars can restore male fertility, thus providing broader genetic resources for rice breeding to produce hybrids with strong hybrid vigour<sup>10–12</sup>. Two-line hybrid rice has been planted in China since 1993 and over 3.4 million hectares were grown in 2011 (Supplementary Fig. 1).

AnnongS-1 (AnS-1), the first *indica* rice (*Oryza sativa* ssp. *indica*) TGMS line, was found in 1987 as a spontaneous mutant<sup>13</sup>. AnS-1 and its derived TGMS lines have been widely used for two-line hybrid rice breeding, and their TGMS trait is controlled by a single recessive locus, *tms5* (ref. 13). A candidate region on the short arm of chromosome 2 has been reported<sup>14–17</sup>; however, the identity of the *TMS5* gene has not been confirmed and the molecular mechanism by which it produces male sterility remains unclear.

Here we report that *tms5* confers the TGMS trait through a loss-of-function mutation in the gene coding for RNase Z<sup>S1</sup>, which regulates *Ubl40* mRNA levels during male development. Our results uncover a novel mechanism of RNase Z<sup>S1</sup>-mediated *Ubl40* mRNA processing, which controls TGMS in rice and has potential applications in hybrid crop breeding.

## Results

### AnS-1 and Zhu1S are temperature-sensitive TGMS lines.

Zhu1S is one of the *indica* TGMS lines frequently used for commercial two-line hybrid rice breeding (Supplementary Fig. 1 and Supplementary Table 1). To examine the phenotype of AnS-1 and Zhu1S in detail, growth chambers and Phytotrons were used to grow plants under fixed photoperiod conditions and at different day average temperatures (DAT; See Methods and Supplementary Table 2a,b). AnS-1 and Zhu1S were grown in the field until the panicle length was ~1 cm and then the plants were transferred into growth chambers or Phytotrons for 2–3 weeks, respectively. At the permissive temperature (~23 °C DAT), AnS-1 is male-fertile, similar to the normal male-fertile line AnnongN (AnN, a parental line of AnS-1). At restrictive temperatures, AnS-1 becomes male-sterile, producing abortive pollen grains at ~26 °C DAT, and no pollen grains at 28 °C or higher DAT (Fig. 1a–h and Supplementary Fig. 2a). At 23 °C, AnS-1 developed normally (Fig. 1i–n), whereas at 28 °C few microspore mother cells (MMCs) undergo normal meiosis, and any

microspores that are produced eventually disintegrate (Fig. 1o–t). By contrast, the day length had little effect on male sterility of AnS-1 (Supplementary Fig. 2). We observed a similar phenotype in Zhu1S grown at the restrictive temperatures of >23 °C DAT (Supplementary Fig. 3).

### Cloning the gene associated with TGMS in AnS-1 and Zhu1S.

To clone *tms5* and the gene conferring the TGMS trait in Zhu1S, we developed and characterized two mapping populations (see Methods, Supplementary Fig. 4a,c), both of which narrowed the target locus to the same region on the short arm of chromosome 2 (Supplementary Fig. 4a,d). Sequencing of the mapped region in the male-sterile lines AnS-1 and Zhu1S revealed a C-to-A transition at position 71 of the gene *Os02g0214300* compared with that in AnN, creating a premature stop codon (Supplementary Fig. 4b,e). This change is identical to the premature stop codon mutation mapped to the same gene in the *ptgms2-1* locus<sup>18</sup>. The protein encoded by *Os02g0214300* was detected by immunoblotting in AnN, but not in AnS-1 (Fig. 1u). Therefore, we considered *Os02g0214300* to be a candidate gene for *TMS5*.

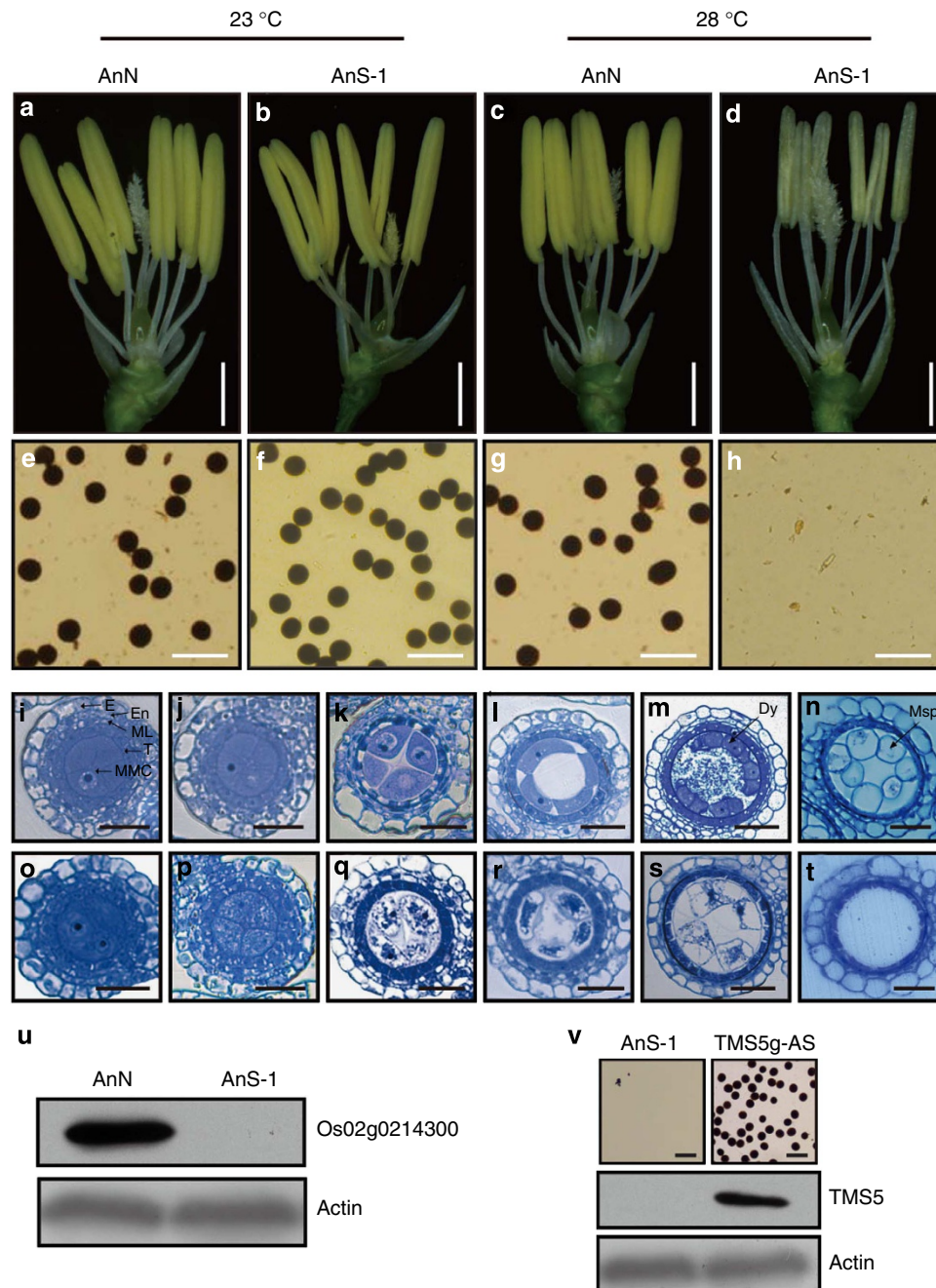
For genetic complementation to verify that *Os02g0214300* indeed corresponds to *TMS5*, genomic DNA fragments harbouring *Os02g0214300* from AnN and the rice cultivar Nipponbare (*O. sativa* ssp. *japonica*) were transformed into the *tms5* AnS-1 and Zhu1S lines, respectively. Male fertility was restored in transgenic plants of AnS-1 (designated TMS5g-AS) and Zhu1S (designated TMS5g-ZS) at the restrictive temperature (Fig. 1v, Supplementary Fig. 5). Moreover, knockdown of *TMS5* by RNA interference (RNAi) in the *japonica* lines Zhonghua11 (ZH11; designated TMS5i-ZH, Supplementary Fig. 6a–d) and Nipponbare (designated TMS5i-NB, Supplementary Fig. 6e–f) showed similar pollen abortion as in AnS-1 and Zhu1S at restrictive temperatures. These results demonstrate that the loss-of-function mutation of *TMS5* confers the TGMS phenotype in AnS-1 and Zhu1S.

### *tms5* is a major TGMS gene widely used in rice breeding.

Dozens of TGMS lines are used in two-line hybrid rice breeding<sup>19</sup>. To explore whether other TGMS lines also harbour a *tms5* mutation, we sequenced the *TMS5* locus in 25 other widely used TGMS lines and found that 24 of them carried the same *tms5* mutation (Fig. 2a). In 2011, 71 commercial two-line hybrid rice cultivars bred with *tms5*-containing TGMS lines accounted for at least 71% of all two-line hybrid rice cultivars and 83.8% (~2.9 million hectares) of all the land used to grow two-line hybrid rice in China (Fig. 2b,c and Supplementary Table 3). Accordingly, *tms5* serves as the major TGMS genetic resource for two-line hybrid rice breeding.

### *TMS5* encodes a conserved RNase Z protein.

*TMS5* encodes an evolutionarily conserved ribonuclease Z (RNase Z), which occurs in almost all kingdoms of life<sup>20–22</sup>. Two forms of RNase Z exist in eukaryotes, the short form (RNase Z<sup>S</sup>) and the long form (RNase Z<sup>L</sup>; Supplementary Fig. 7), while prokaryotes only have RNase Z<sup>S</sup> (refs 20,22). The *TMS5* protein, with 302 amino acids, belongs to the short-form group and we refer to it as RNase Z<sup>S1</sup>. In rice, RNase Z<sup>S1</sup> is constitutively expressed in various tissues (Supplementary Fig. 8a). At the protein level, RNase Z<sup>S1</sup> accumulated stably at both permissive and restrictive temperatures (Supplementary Fig. 8b). In anther, its mRNA expression was relatively enriched in MMCs and this did not change at low (23 °C) and high (28 °C) temperatures (Supplementary Fig. 8c,d). These results indicate that RNase Z<sup>S1</sup> itself is temperature-insensitive.

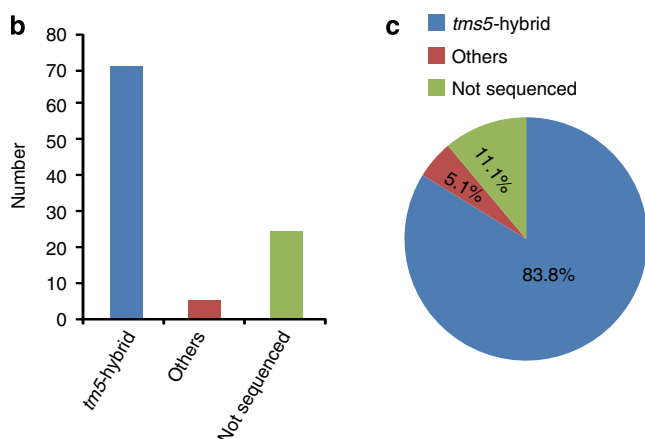


**Figure 1 | Phenotypes and complementation of AnS-1.** (a–h) Anther morphology and pollen fertility of AnN and AnS-1. Normal anthers (a,b) and pollen (e,f) in AnN and AnS-1 at the permissive temperature (23 °C). Normal anthers (c) and pollen (g) in AnN and abnormal anthers (d) and abortive pollen (h) in AnS-1 grown at the restrictive temperature (28 °C). (i–t) Comparison of transverse sections of anthers of AnS-1 grown at 23 °C (i–n) and 28 °C (o–t). At 23 °C, the MMCs underwent meiosis to produce microspores (immature pollen grains); at 28 °C, few or no microspores were generated. (i,o) Early MMC stage; (j,p) late MMC stage; (k,q) meiotic leptotene stage; (l,r) meiotic pachytene stage; (m,s) dyad stage; (n,t) young microspore stage. E, epidermis; En, endothecium; ML, middle layer; T, tapetum; MMC, microspore mother cell, Dy, dyad; Msp, microspore. Scale bars, 1 mm (a–d), 100  $\mu$ m (e–h) and 20  $\mu$ m (i–t). (u) The protein encoded by Os02g0214300 could not be detected in AnS-1 by immunoblot analysis. The constitutively expressed rice Actin protein was used as an internal control. (v) Pollen fertility was restored in the AnS-1 background containing *TMS5* genomic DNA (TMS5g-AS) grown at the restrictive temperature (28 °C; above, right). No pollen was produced in anthers of AnS-1 grown under the same conditions (above, left). *TMS5* protein was detected in TMS5g-AS line but not in AnS-1 (middle). The Actin protein was used as an internal control.

**RNase Z<sup>S1</sup> is cytoplasmic and processes tRNAs *in vitro*.** RNase Z proteins process the 3' ends of tRNAs<sup>21,23–25</sup>. The CCA sequence in 3' terminal of the tRNA is critical for tRNA maturation and function. RNase Z mainly processes CCA-less pre-tRNAs; however, it can process CCA-containing pre-tRNAs in the nucleus or mitochondria<sup>26,27</sup>. To investigate the enzymatic

activity of RNase Z<sup>S1</sup>, we tested whether RNase Z<sup>S1</sup> cleaved two pre-tRNAs, pre-tRNA<sup>9</sup>-AspATC (without CCA) and pre-tRNA<sup>35</sup>-MetCAT (with CCA). We expressed recombinant RNase Z<sup>S1</sup> in *Escherichia coli* (*E. coli*) and found that it processed the 3' ends of both pre-tRNAs, with or without CCA, *in vitro* (Fig. 3a), indicating that RNase Z<sup>S1</sup> possesses

	1	11	21	61	71
<b>Wild type</b>					↘
Nipponbare	ATG GCG AAC AGC GGC AAG TCA . . . G CCA CCG GGT CGG CCG AAG GCG AAG				
93-11	ATG GCG AAC AGC GGC AAG TCA . . . G CCA CCG GGT CGG CCG AAG GCG AAG				
AnN	ATG GCG AAC AGC GGC AAG TCA . . . G CCA CCG GGT CGG CCG AAG TCG AAG				
AnS-1	ATG GCG AAC AGC GGC AAG TCA . . . G CCA CCG GGT CGG CCG AAG TAG AAG				
Zhu1S	ATG GCG AAC AGC GGC AAG TCA . . . G CCA CCG GGT CGG CCG AAG TAG AAG				
Xiang125S	ATG GCG AAC AGC GGC AAG TCA . . . G CCA CCG GGT CGG CCG AAG TAG AAG				
Lu18S	ATG GCG AAC AGC GGC AAG TCA . . . G CCA CCG GGT CGG CCG AAG TAG AAG				
Guangzhan63S	ATG GCG AAC AGC GGC AAG TCA . . . G CCA CCG GGT CGG CCG AAG TAG AAG				
Y58S	ATG GCG AAC AGC GGC AAG TCA . . . G CCA CCG GGT CGG CCG AAG TAG AAG				
Xuan69S	ATG GCG AAC AGC GGC AAG TCA . . . G CCA CCG GGT CGG CCG AAG TAG AAG				
C815S	ATG GCG AAC AGC GGC AAG TCA . . . G CCA CCG GGT CGG CCG AAG TAG AAG				
Xin'anS	ATG GCG AAC AGC GGC AAG TCA . . . G CCA CCG GGT CGG CCG AAG TAG AAG				
ZhunS	ATG GCG AAC AGC GGC AAG TCA . . . G CCA CCG GGT CGG CCG AAG TAG AAG				
1892S	ATG GCG AAC AGC GGC AAG TCA . . . G CCA CCG GGT CGG CCG AAG TAG AAG				
Guangxiang24S	ATG GCG AAC AGC GGC AAG TCA . . . G CCA CCG GGT CGG CCG AAG TAG AAG				
Guangzhan63-4S	ATG GCG AAC AGC GGC AAG TCA . . . G CCA CCG GGT CGG CCG AAG TAG AAG				
GuangmoS	ATG GCG AAC AGC GGC AAG TCA . . . G CCA CCG GGT CGG CCG AAG TAG AAG				
Feng39S	ATG GCG AAC AGC GGC AAG TCA . . . G CCA CCG GGT CGG CCG AAG TAG AAG				
XinhuaS	ATG GCG AAC AGC GGC AAG TCA . . . G CCA CCG GGT CGG CCG AAG TAG AAG				
N422S	ATG GCG AAC AGC GGC AAG TCA . . . G CCA CCG GGT CGG CCG AAG TAG AAG				
HD9802S	ATG GCG AAC AGC GGC AAG TCA . . . G CCA CCG GGT CGG CCG AAG TAG AAG				
GD-5S	ATG GCG AAC AGC GGC AAG TCA . . . G CCA CCG GGT CGG CCG AAG TAG AAG				
GD-7S	ATG GCG AAC AGC GGC AAG TCA . . . G CCA CCG GGT CGG CCG AAG TAG AAG				
TianyuanS	ATG GCG AAC AGC GGC AAG TCA . . . G CCA CCG GGT CGG CCG AAG TAG AAG				
360S	ATG GCG AAC AGC GGC AAG TCA . . . G CCA CCG GGT CGG CCG AAG TAG AAG				
SE21S	ATG GCG AAC AGC GGC AAG TCA . . . G CCA CCG GGT CGG CCG AAG TAG AAG				
X07S	ATG GCG AAC AGC GGC AAG TCA . . . G CCA CCG GGT CGG CCG AAG TAG AAG				
AnxiangS	ATG GCG AAC AGC GGC AAG TCA . . . G CCA CCG GGT CGG CCG AAG TAG AAG				
Lubai76S	ATG GCG AAC AGC GGC AAG TCA . . . G CCA CCG GGT CGG CCG AAG TAG AAG				
AnlongS	ATG GCG AAC AGC GGC AAG TCA . . . G CCA CCG GGT CGG CCG AAG TAG AAG				
Pei'ai64S	ATG GCG AAC AGC GGC AAG TCA . . . G CCA CCG GGT CGG CCG AAG GCG AAG				



**Figure 2 | Application of *tms5* in two-line hybrid rice breeding.** (a) Left, names of wild-type and TGMS lines. Right, alignment of 75 base-pair DNA sequence of the coding region of *TMS5* in wild-type and different TGMS lines. Nucleotide numbers within the *TMS5*-coding sequence are shown at the top. The mutated nucleotide at position 71 (arrowed) in *tms5* lines is shown in red. (b) Number of different two-line hybrid rice cultivars bred using *tms5*-TGMS lines (*tms5*-hybrid) compared with other two-line hybrid rice cultivars grown in China in 2011 (left figure). Proportions of the two-line hybrid rice-growing area (83.4 million hectares) planted with *tms5*-hybrid cultivars and other hybrid cultivars bred with others and non-sequenced T/PGMS lines in China in 2011 (right figure). (c) Proportions of the two-line hybrid rice-growing area (~3.4 million hectares) planted with *tms5*-hybrid cultivars and other hybrid cultivars bred with others and non-sequenced T/PGMS lines in China in 2011.

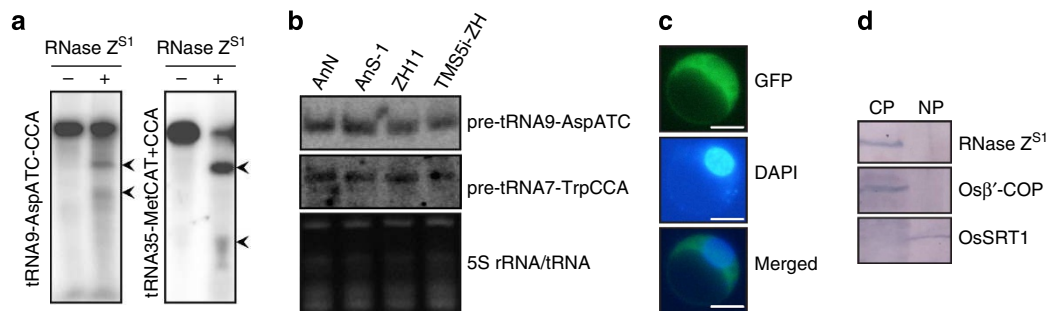
endonuclease activity. We then examined the *in vivo* accumulation of mature and precursor tRNA in AnN, AnS-1, ZH11 and TMS5i-ZH but found no obvious difference between wild-type and *tms5* plants (Fig. 3b and Supplementary Fig. 9a). The mature tRNA levels were unchanged at permissive or restrictive temperatures when analysed in NIL5 and NIL8 (near-isogenic lines 5 and 8 of Zhu1S in the background of the *indica* cultivar 93-11) and 93-11 (Supplementary Fig. 9b). These results indicate that RNase Z<sup>S1</sup> does not affect tRNA 3' end processing *in vivo*.

To explore where RNase Z<sup>S1</sup> localizes, we expressed RNase Z<sup>S1</sup>-GFP (green fluorescent protein) in rice protoplasts and found that it localizes in the cytoplasm but not in the nucleus (Fig. 3c).

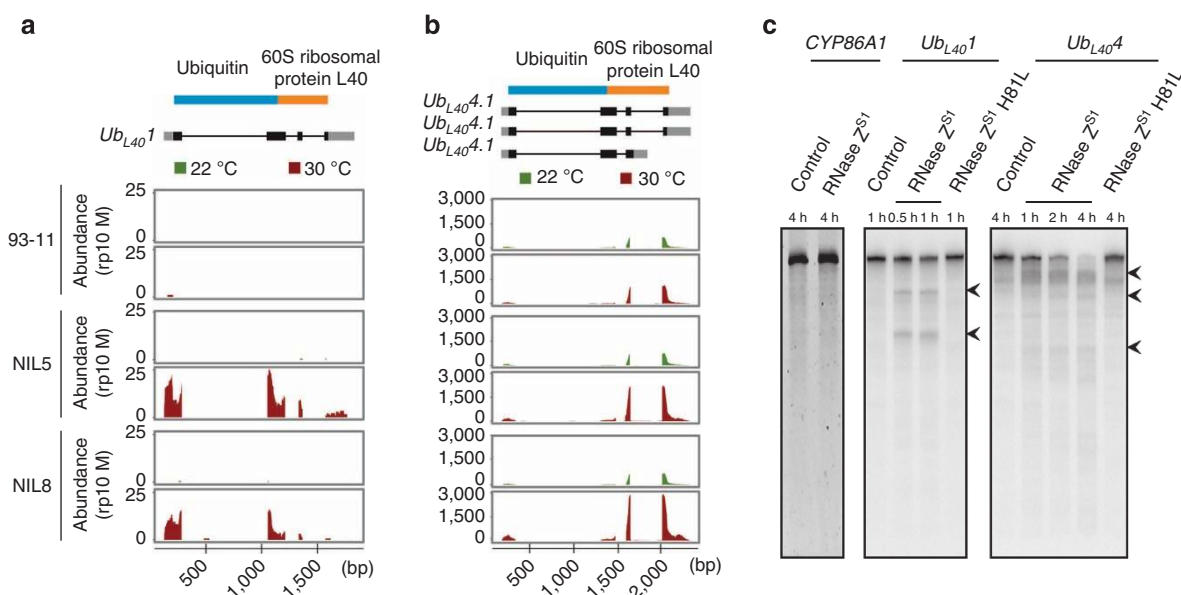
This observation was further verified by immunoblot analysis of partially purified nuclear and cytoplasmic proteins (Fig. 3d). Therefore, we propose that RNase Z<sup>S1</sup> functions in the cytoplasm but does not act in tRNA processing in the nucleus.

#### *Ubl40* mRNAs accumulate in *tms5* plants at high temperature.

To further address whether RNase Z<sup>S1</sup> has a role in mRNA metabolism in the cytoplasm, we performed independent whole-genome microarray and RNA-seq analyses, using RNA from young panicles (flower inflorescences) of wild-type lines (AnN, ZH11 and 93-11) and *tms5* or RNase Z<sup>S1</sup> knockdown lines (AnS-1, TMS5i-ZH, NIL5 and NIL8) grown at permissive and



**Figure 3 | RNase Z<sup>S1</sup> processes tRNA *in vitro* and localizes in the cytoplasm.** (a) Recombinant His-tagged RNase Z<sup>S1</sup> processed the 3' ends of both tRNA9-AspATC-CCA and tRNA35-MetCAT + CCA *in vitro*. Arrows indicate the processed product bands. (b) RNA blot analysis showed no obvious difference in tRNA precursor accumulation in wild-type (AnN and ZH11), AnS-1 and TMS5i-ZH plants. 5S rRNA/tRNA stained with ethidium bromide was used as a loading control. (c) RNase Z<sup>S1</sup>-GFP (green) was observed in the cytoplasm (top) but not in the nucleus (middle, stained with DAPI, blue). Scale bars, 10  $\mu$ m. (d) RNase Z<sup>S1</sup> was detected in cytoplasmic protein extracts using immunoblot analysis. Os $\beta$ '-COP and OsSRT1 were used as the indicators of cytoplasmic and nuclear protein controls, respectively. CP, cytoplasmic proteins; NP, nuclear proteins.



**Figure 4 | Ub<sub>L401</sub> and Ub<sub>L404</sub> mRNAs accumulate in *tms5* plants at high temperature.** (a, b) Overaccumulation of Ub<sub>L401</sub> and Ub<sub>L404</sub> mRNAs in *tms5* plants at the restrictive temperature. RNA-seq profile of Ub<sub>L401</sub> (a) and Ub<sub>L404</sub> (b) in 93-11, NIL5 and NIL8 (near-isogenic line 5 and line 8 of Zhu1S-derived *tms5* in 93-11 background) at 22 and 30 °C. The ubiquitin domain and 60S ribosomal protein L40 domain are shown as blue and orange boxes, respectively. Transcripts of Ub<sub>L401</sub> (one mRNA isoform) and Ub<sub>L404</sub> (three mRNA isoforms) are shown with thick boxes representing exons, thin lines representing introns and grey boxes representing untranslated regions (UTRs). Wiggle plots of RNA-seq data represent reads per 10 million (rp10M). (c) *In vitro* processing of CYP86A1 (left, a negative control), Ub<sub>L401</sub> (middle) and Ub<sub>L404</sub> (right) mRNAs by recombinant wild-type RNase Z<sup>S1</sup> and enzymatic null (RNase Z<sup>S1</sup>H81L), respectively. The assay time is marked above the lanes. Arrows indicate bands processed by RNase Z<sup>S1</sup>.

restrictive temperatures. These analyses identified three common mRNAs that accumulate at higher levels in *tms5* plants at the restrictive but not the permissive temperature (Fig. 4a,b; Supplementary Fig. 10a and Supplementary Table 4). These mRNAs are from genes of the conserved ubiquitin-60S ribosomal protein L40 family (Ub<sub>L40</sub>), namely Ub<sub>L401</sub> (Os09g0452700), Ub<sub>L402</sub> (Os03g0259500) and Ub<sub>L404</sub> (Os09g0483400; Supplementary Fig. 11). At the permissive temperature, Ub<sub>L401</sub>, Ub<sub>L402</sub> and Ub<sub>L404</sub> mRNA levels were similar in wild-type and *tms5* plants; at the restrictive temperature, the expression of these three genes, especially Ub<sub>L401</sub>, was dramatically elevated in *tms5* but not in wild-type plants (Fig. 4a,b; Supplementary Figs 10a, 12a and 13). Furthermore, Ub<sub>L401</sub>, Ub<sub>L402</sub> and Ub<sub>L404</sub> preferentially expressed in stamens and were largely induced by restrictive temperature in *tms5* stamens but not in other tissues

(Supplementary Fig. 12b–e). Thus, accumulation of Ub<sub>L401</sub>, Ub<sub>L402</sub> and Ub<sub>L404</sub> transcripts is temperature-sensitive in the *tms5* background, and RNase Z<sup>S1</sup>-dependent, suggesting that RNase Z<sup>S1</sup> might be involved in processing Ub<sub>L40</sub> mRNAs.

**RNase Z<sup>S1</sup> processes mRNAs of Ub<sub>L40</sub> genes.** To address this hypothesis, we transcribed Ub<sub>L401</sub>, Ub<sub>L402</sub> and Ub<sub>L404</sub> RNAs *in vitro* and incubated the RNAs with recombinant RNase Z<sup>S1</sup> proteins. All three Ub<sub>L40</sub> mRNAs were processed into multiple fragments by the wild-type but not the enzymatic null RNase Z<sup>S1</sup> proteins *in vitro* (Fig. 4c, Supplementary Fig. 10b). Furthermore, we employed RNA ligase-mediated 5' rapid amplification of cDNA ends (RNA ligase-mediated-5'RACE) and mapped multiple degradation sites in the Ub<sub>L401</sub> mRNA of wild-type plants

(ZH11, AnN and 93-11) but not in AnS-1 and NIL5 (Supplementary Fig. 14), suggesting that RNase Z<sup>S1</sup> is involved in the processing of *Ubl<sub>L401</sub>* mRNA *in vivo*. We hypothesize that RNase Z<sup>S1</sup> may recognize the secondary or tertiary structure of the target mRNAs and process them. Thus, we propose that at restrictive temperatures, the loss-of-function mutation of RNase Z<sup>S1</sup> in *tms5* plants results in unprocessed *Ubl<sub>L401</sub>*, *Ubl<sub>L402</sub>* and *Ubl<sub>L404</sub>* RNAs and the excess *Ubl<sub>L40</sub>* transcripts might lead to abnormal male development in the anther.

***Ubl<sub>L40</sub>* genes are expressed in MMCs.** To further test our hypothesis, we examined the spatial and temporal expression of *Ubl<sub>L40</sub>* genes by *in situ* mRNA hybridization. In wild-type anthers, compared with sense probe signals, antisense probes of *Ubl<sub>L401</sub>* and *Ubl<sub>L404</sub>* detected stronger signals in MMCs than in other cell layers (Fig. 5a,b). In addition, *Ubl<sub>L401</sub>*, *Ubl<sub>L402</sub>* and *Ubl<sub>L404</sub>* expression levels were similar between high (28 °C DAT) and low (23 °C DAT) temperatures in wild type (Fig. 5c and Supplementary Fig. 15). In contrast, all three *Ubl<sub>L40</sub>* genes tested showed stronger signals in MMCs at high temperature than in the RNase Z<sup>S1</sup> mutants at low temperature, indicating the interplay between RNase Z<sup>S1</sup> and *Ubl<sub>L40</sub>* mRNAs in male fertility (Fig. 5d and Supplementary Fig. 15). Among the *Ubl<sub>L40</sub>* family members, *Ubl<sub>L404</sub>* may have a major role in the regulation of thermosensitive genic male sterility since the expression level of *Ubl<sub>L404</sub>* was much higher than that of *Ubl<sub>L401</sub>* and *Ubl<sub>L402</sub>*, consistent with the RNA-seq data in NIL5 and NIL8, and RT-PCR data in AnS-1 (Figs 4a,b and 5d; Supplementary Figs 10, 12, 13 and 15).

**Overexpression of *Ubl<sub>L401</sub>* and 4 leads to male sterility.** To test whether the accumulation of *Ubl<sub>L401</sub>* and *Ubl<sub>L404</sub>* in *tms5* plants could affect male fertility, we generated *Ubl<sub>L401</sub>*- and *Ubl<sub>L404</sub>*-overexpressing transgenic plants in the ZH11 background (designated *Ubl<sub>L401</sub>*OE and *Ubl<sub>L404</sub>*OE). These plants produced higher levels of *Ubl<sub>L401</sub>* and *Ubl<sub>L404</sub>* mRNAs than ZH11 and exhibited partial pollen abortion (Supplementary Fig. 16), mimicking the phenotypes of *tms5* lines at restrictive temperatures. Furthermore, we created *Ubl<sub>L401</sub>* or *Ubl<sub>L404</sub>* knockdown plants in the *tms5* background, designated *Ubl<sub>L401</sub>*i and *Ubl<sub>L404</sub>*i. At restrictive temperatures, these plants exhibited reduced levels

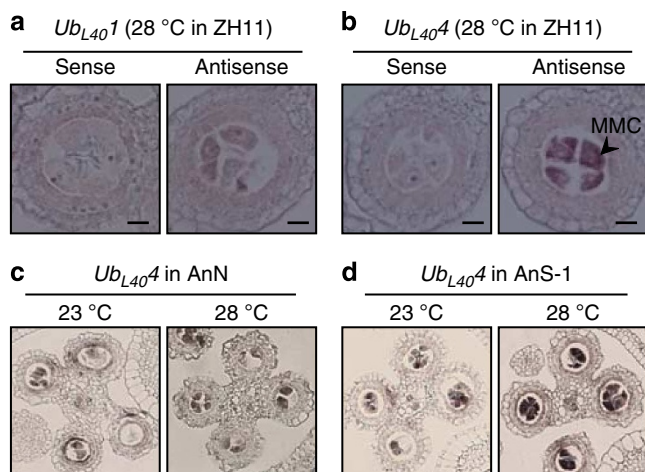
of *Ubl<sub>L401</sub>* and *Ubl<sub>L404</sub>* mRNAs and partially restored male fertility (Supplementary Fig. 17). Thus, we hypothesize that regulation of *Ubl<sub>L40</sub>* mRNA levels by RNase Z<sup>S1</sup> is critical for MMC development at high temperature in rice.

## Discussion

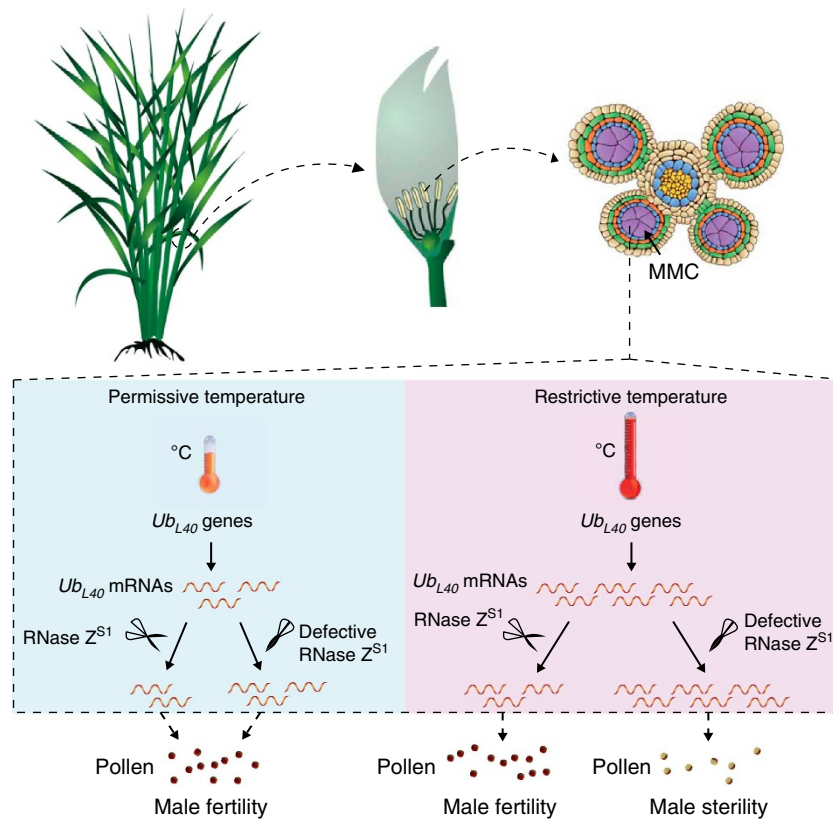
Male sterility has been extensively applied to the breeding of crops for hybrid vigour, and these hybrids have larger stature and higher yields than the parent inbred lines. In rice, CMS lines and TGMS/PGMS lines are widely used in three-line and two-line hybrid rice breeding, respectively. Here we uncovered the molecular mechanism of rice TMS5, which functions in RNase Z<sup>S1</sup>-mediated *Ubl<sub>L40</sub>* mRNA regulation during male development (Fig. 6). At the permissive temperature, the level of *Ubl<sub>L40</sub>* mRNAs remains low in the *tms5* mutant plants, allowing the production of normal pollen. At restrictive temperature, however, *Ubl<sub>L40</sub>* mRNAs are not processed by RNase Z<sup>S1</sup> and their high level cause male sterility. The preferential expression of *Ubl<sub>L401</sub>*, *Ubl<sub>L402</sub>* and *Ubl<sub>L404</sub>* in MMCs indicates that these mRNAs are crucial for MMC development. We notice that overexpression of *Ubl<sub>L401</sub>* and *Ubl<sub>L404</sub>* in wild type leads to male sterility without causing any other obvious defects. We hypothesize that additional factors that are restricted to the MMC might be required to assist *Ubl<sub>L40</sub>* genes in their functions (Fig. 6). Our results thus uncover a novel function for the RNase Z<sup>S1</sup> family, deepening our understanding of the molecular genetic basis of TGMS in rice. These findings have profound implications for hybrid breeding in crops.

Male reproduction is very sensitive to alterations in environmental conditions. A long non-coding RNA (lncRNA), the long-day-specific male-fertility-associated RNA, causes a PGMS trait and was identified in Nongken58S<sup>11</sup>. At the same locus in the *indica* line Pei'ai64S, a non-coding RNA producing small RNAs was shown to confer a P/TGMS trait<sup>10</sup>. In addition, a temperature-sensitive splicing defect of the transcript encoding UDPase leads to male sterility in rice at high temperature<sup>28,29</sup>. Our results show that, in *tms5* mutants, high temperature induces accumulation of *Ubl<sub>L40</sub>* mRNAs in MMCs and causes male sterility. It is possible that environmental conditions such as temperature and photoperiod may affect RNA metabolism at the post-transcriptional level. Defects in these processes might be harmful to the cells in anther development, leading to male sterility.

The endonuclease RNase Z exhibits two forms, RNase Z<sup>S</sup> and RNase Z<sup>L</sup>, which are evolutionarily conserved in almost all organisms<sup>21,22</sup>. RNase Z proteins have various RNA substrates, including tRNAs and mRNAs. Different RNase Z proteins might process different substrates and functions in different subcellular compartments<sup>21</sup>. In prokaryotes, RNase Z<sup>S</sup> is involved in tRNA 3' terminal processing and has a role in mRNA decay<sup>30</sup>. In *Arabidopsis*, four RNase Z proteins can process tRNA *in vitro*<sup>25</sup>: Trz1 and Trz2 (RNase Z<sup>L</sup>) localize in the nucleus and mitochondria and may process tRNAs; Trz3 and Trz4 (RNase Z<sup>S</sup>) localize in the chloroplasts and cytoplasm, respectively. Loss-of-function of *Trz4* leads to embryo lethality, and the function of *Trz3*, which localizes in the cytoplasm, is largely unknown<sup>25</sup>. The rice genome encodes three RNase Z proteins, including one long form and two short forms (RNase Z<sup>S1</sup> and RNase Z<sup>S2</sup>; Supplementary Fig. 7). RNase Z<sup>S2</sup> localizes in the chloroplast and is essential to chloroplast development<sup>31</sup>. In humans, ELAC2 (RNase Z<sup>L</sup>) localizes in the nucleus and mitochondria<sup>32,33</sup>. ELAC2 can process the lncRNA *MALAT1* and plays a role in the generation of a mammalian herpesvirus microRNA<sup>34,35</sup>. ELAC1 (RNase Z<sup>S</sup>) occurs in the cytoplasm but has unknown functions<sup>36,37</sup>. The enzymatic activity of ELAC2 is over 1,600-fold higher than ELAC1, suggesting that ELAC1 might



**Figure 5 | *Ubl<sub>L401</sub>* and *Ubl<sub>L404</sub>* mRNAs preferentially express in MMCs.** (a,b) *In situ* hybridization of *Ubl<sub>L401</sub>* (a) and *Ubl<sub>L404</sub>* (b) transcripts in anther. (c,d) At indicated temperatures, expression level of *Ubl<sub>L404</sub>* in MMCs under high (28 °C) and low (23 °C) temperature in AnN (c) and AnS-1 (d). Scale bars, 10 μm (a,b). MMC, microspore mother cells.



**Figure 6 | A functional model for RNase Z<sup>S1</sup> controlling TGMS in rice.** In rice anther, *Ub<sub>L40</sub>* genes preferentially express in MMCs (in purple). *Ub<sub>L40</sub>* mRNAs are induced by high (restrictive) temperature. In wild type, RNase Z<sup>S1</sup> processes mRNAs of *Ub<sub>L40</sub>* and maintains them at normal levels. In RNase Z<sup>S1</sup> defective mutant plants, excessive mRNAs of *Ub<sub>L40</sub>* accumulate in anthers and lead to male sterility. MMC, microspore mother cells.

have new functions other than tRNA processing in the cytoplasm<sup>38</sup>. Our data provide the first direct experimental evidence demonstrating the biochemical roles of a short form of RNase Z (RNase Z<sup>S1</sup>) in mRNA metabolism in the cytoplasm of an eukaryote. Our findings could provide clues to the functions of the RNase Z family not only in plants but also in animals.

## Methods

**Plant materials and growth conditions.** Rice accessions, including AnnonS-1 (*O. sativa* ssp. *indica*, AnS-1, *tms5*), AnnonN (*O. sativa* ssp. *indica*, AnN, WT siblings of AnS-1), Zhonghua11 (*O. sativa* ssp. *japonica*, ZH11, WT), *tms5*-ZH11 (the progeny of AnS-1 after two backcrosses to ZH11) and transgenic lines TMS5g-AS (RNase Z<sup>S1</sup> transgene in AnS-1 background), TMS5i-ZH (RNase Z<sup>S1</sup> RNAi in ZH11 background), *Ub<sub>L40</sub>*1OE (*Ub<sub>L40</sub>*1 overexpression in ZH11 background), *Ub<sub>L40</sub>*4OE (*Ub<sub>L40</sub>*4 overexpression in ZH11 background), *Ub<sub>L40</sub>*1i (*Ub<sub>L40</sub>*1 RNAi in ZH11 background) and *Ub<sub>L40</sub>*4i (*Ub<sub>L40</sub>*4 RNAi in ZH11 background) were grown in the field under normal conditions at the South China Agricultural University, Guangzhou, or grown in a growth chamber at indicated temperature when panicle length reaches ~1 cm (before the stages of early premeiosis) for ~2 weeks until flowering. All samples from AnS-1 and its controls were harvested between 1500 and 1700 hours in the growth chamber and Zhu1S and its controls were harvested between 0800 and 1100 hours in the Phytotron.

**Characterization of AnS-1 and Zhu1S phenotypes.** Rice florets were photographed with an OLYMPUS DP70 digital camera and an OLYMPUS SZX10 dissecting microscope. Mature pollen grains were stained with 1% I<sub>2</sub>-KI solution and

observed with an OLYMPUS BX51 or Leica DNRXA microscope. Panicles from various stages of development were fixed with a mixture of 4% paraformaldehyde and 1.5% glutaraldehyde in PBS solution at 4 °C. The samples were then fixed again for 1 h with 1% osmic acid. Fixed anthers were washed three times for 10 min each in PBS solution. After washing, the samples were dehydrated in five successive graded ethanol baths (from 50 to 95%) for 10 min each and then washed in 100% propylene oxide twice for 15 min each. The samples were infiltrated with resin/propylene oxide (1:1 mixture) for 2 h and resin/propylene oxide (3:1 mixture) for 3 h, embedded in pure resin overnight, and then placed in an oven at 60 °C for 48 h. After capsule-embedding, blocks were trimmed on an LKB pyramitome. Trimmed blocks were cut to 2-μm semithin sections on an LKB ultramicrotome with glass knives. The semithin sections were stained with 0.2% toluidine blue O (Chroma-Gesellschaft, Germany) and examined with the Leica DNRXA light microscope or OLYMPUS BX51 microscope.

**Fine mapping of *tms5* and the gene conferring TGMS in Zhu1S.** The F<sub>2</sub> mapping population was generated from crosses between Xiang125S (derived from AnS-1) and Jingxian 89 (*O. sativa* ssp. *indica*) and grown in the field during the summer in Guangzhou (DAT > 28 °C), Guangdong province in China. The introgressed mapping population (BC<sub>1</sub>F<sub>2</sub> and BC<sub>2</sub>F<sub>2</sub>) was generated according to the programme shown in Supplementary Fig. 4c and planted in the summer at Changsha (DAT > 28 °C), Hunan province in China. Details of the markers used for mapping are listed in Supplementary Table 5. The candidate genes were sequenced and compared in wild type (AnN and 93-11) and TGMS (AnS-1 and Zhu1S) using the primers shown in Supplementary Table 6.

**Vector construction, plant transformation and analysis.** For functional complementation tests, a 5.3-kb DNA fragment containing 2.3 kb of upstream sequence, the entire *TMS5*-coding sequence, and 0.9 kb of downstream sequence, was amplified from AnN genomic DNA using primers TMS5g-AS F/R (Supplementary Table 6), and then digested with BamHI and HindIII and ligated to the binary vector pCambia1300 for gene transformation. A 5.1-kb DNA fragment containing a 2.2-kb upstream sequence, the entire *TMS5* gene and a 1.0-kb downstream region from a BAC library (OSJNBb00118013, Arizona Genomics Institute), was digested with HindIII and ligated to the binary vector pCambia2300 to get the complementary construction, XF2108. For TMS5i-ZH and TMS5i-NB RNAi (construction of XF2114), two DNA fragments (415 and

360 bp, respectively) were cloned from *TMS5* cDNA using the primer pairs TMS5i-ZH F/R and TMS5i-NB F/R (Supplementary Table 6), and then inserted into pYL RNAi<sup>39</sup> and pUCC-RNAi<sup>40</sup>. For *Ubl<sub>40</sub>1* and *Ubl<sub>40</sub>4* RNAi, two DNA fragments (242 and 322 bp, respectively) were amplified from *Ubl<sub>40</sub>1* and *Ubl<sub>40</sub>4* using two pairs of primers, *Ubl<sub>40</sub>1* F/R and *Ubl<sub>40</sub>4* F/R (Supplementary Table 6), and inserted into pYL RNAi. For *Ubl<sub>40</sub>1* and *Ubl<sub>40</sub>4* overexpression, two DNA fragments (424 bp) from *Ubl<sub>40</sub>1* and *Ubl<sub>40</sub>4* were amplified using two pairs of primers, *Ubl<sub>40</sub>1*OE F/R and *Ubl<sub>40</sub>4*OE F/R, and inserted into the pYLox vector<sup>39</sup> driven by the ubiquitin promoter. Transgenic plants were generated via *Agrobacterium*-mediated transformation<sup>41,42</sup>. For each vector, 15 transgenic lines were selected and genotyped. Positive transgenic lines were selected for further analysis by expression level of target genes. Transgenic lines with moderate expression levels were next analysed and average phenotypes were photographed. The relevant PCR primers are listed in Supplementary Table 6.

**Subcellular localization.** The open reading frame of *TMS5* was amplified and fused with the N terminus of GFP<sup>43</sup> in the pUC18 vector under the control of the CaMV35S promoter. Protoplasts from leaf sheaths of rice plants were isolated as described previously<sup>44</sup>. Briefly, for protoplast transformation, 10 µl of plasmid carrying *TMS5* and GFP fragments, 100 µl of protoplasts and 110 µl of PEG solution (40% PEG4000, 0.3 M mannitol and 0.1 M CaCl<sub>2</sub>) were mixed gently and incubated for 15 min. After transformation, cells were washed with W5 solution and then resuspended in WI solution (4 mM MES, pH 5.7, 0.5 M mannitol and 20 mM KCl). Cells were incubated ~20–24 h after transformation. GFP signals were observed under the OLYMPUS BX51 fluorescence microscope.

**Expression analyses.** Total RNA was isolated from rice roots, stems, leaves, panicles, stamens and pistils at different temperatures using TRIzol (Invitrogen, USA) reagent. DNase I-treated total RNA (2.0 µg) was used for reverse transcription using the M-MLV-RT kit (TAKARA, Japan). RT-PCR and real-time quantitative PCR were performed as previously described<sup>39,40</sup>. Briefly, equal amounts of RT products were used to perform PCR. Quantitative PCR analyses were performed three repeats for each sample using the SsoFast EvaGreen Supermix kit (Bio-Rad, USA) with CFX96 Real-Time PCR Detection System (Bio-Rad). The cDNA levels of target genes were normalized to the internal standard genes *OsActin1* and *OsEF1-α*. RNA blotting was performed as previously described<sup>42</sup>. The hybridization signals were visualized using a Typhoon 9000 System (GE, USA). Nuclear and cytoplasmic proteins were purified using Plant Total Soluble Nuclear Protein High-Pure Kit (GENMED, USA). Total protein was extracted from rice panicles grown at different temperatures. Immunoblot analysis was performed using Li's methods<sup>42</sup>. The complementary sequences corresponding to tRNAs were used as probes labelled with γ-<sup>32</sup>P-ATP using T4 polynucleotide kinase. The hybridization was performed at 42 °C in a hybridizer. The protein extracts were separated by 12% SDS-polyacrylamide gel electrophoresis and transferred to a Pure Nitrocellulose Blotting Membrane (Pall Corporation). The membrane was successively incubated with the anti-RNase Z<sup>S1</sup> antiserum (1:1,000 dilution) and secondary antibody goat anti-rabbit IgG HRP (1:10,000 dilution). Actin (Code: M20009; Abmart, China) was selected as the internal standard protein. OsSRT1 (ref. 45) and Osβ'-COP<sup>46</sup> were selected as the nuclear and cytoplasmic protein controls. Anti-TMS5, anti-OsSRT1 and anti-Osβ'-COP were generated by immunizing rabbits. The relevant PCR primers are listed in Supplementary Table 6.

**Overproduction and purification of RNase Z<sup>S1</sup>.** The RNase Z<sup>S1</sup> and RNase Z<sup>S1</sup>H81L sequences were amplified (primers see Supplementary Table 6) and cloned into plasmid pET23d (Novagen, Germany). After transformation of the *E. coli* BL21 (DE3) strain with the resulting plasmid constructs in 2 l of LB culture medium at 37 °C, expression of histidine-tagged RNase Z<sup>S1</sup> was induced by the addition of 0.1 mM isopropyl-β-D-thiogalactopyranoside and grown at 20 °C overnight. The culture was harvested, pelleted, resuspended in 30 ml of buffer containing lysozyme and stirred for 3 h at 4 °C. The suspension was treated with an ultrasonic disintegrator (SONICS, USA) at 4 °C and then centrifuged at 15,000 g for 30 min to remove cell debris. The supernatant was loaded onto a Ni<sup>2+</sup>-NTA column (Qiagen, Germany). RNase Z<sup>S1</sup> was purified using the histidine tag according to the manufacturer's instructions (Novagen) and immediately dialysed. Protein purity was then verified using SDS-PAGE analysis (12%) and estimated at >95%.

**Synthesis of tRNA precursors and *Ubl<sub>40</sub>* RNAs *in vitro*.** Precursor tRNAs, *Ubl<sub>40</sub>* and *CYP86A1* were amplified using specific primers (Supplementary Table 6) and cloned into plasmid pEASY-T3 (Transgene, China). Precursor tRNAs, *Ubl<sub>40</sub>* and *CYP86A1* were transcribed *in vitro* using T7 RNA polymerase and labelled with [α-<sup>32</sup>P]-CTP, according to the manufacturer's instructions (Promega, USA). The template DNA was removed by the addition of DNase I (Fermentas, USA). The labelled RNAs were purified using phenol and ethanol.

**RNase Z<sup>S1</sup> enzymatic activity assays.** RNase Z<sup>S1</sup> activity was assayed as previously described<sup>24</sup>. Briefly, the labelled precursor tRNAs or *Ubl<sub>40</sub>* RNA were

incubated for 0.5, 1, 2 or 4 h at 37 °C with 200 ng purified RNase Z<sup>S1</sup> or enzymatic null RNase Z<sup>S1</sup>H81L, in which a substitution of histidine by leucine in position 81 in the conserved enzymatic domain of RNase Z proteins causes the loss of enzymatic activity<sup>47</sup>. The RNase Z<sup>S1</sup> cleavage products were purified using the RNA probe purification kit (OMEGA, USA). Equal volumes of reactant solutions and 2 × loading buffer (95% formamide, 20 mM EDTA, 0.05% bromophenol blue and 0.05% xylene cyanol) were mixed. Denaturing gel electrophoresis was performed by 8 or 15% PAGE containing 7 M urea and the signals were visualized with the Typhoon 9000 System (GE).

**RNA ligase-mediated 5' RACE.** A 5' RNA adaptor (5'-CUGACUGCACUCAG AGUACUACAGCCGAC-3') was ligated to ~2.0 µg of total RNA using T4 RNA ligase 1 (NEB, USA). The ligated mRNAs were then reverse transcribed using oligo(dT)<sub>15</sub> primer with AMV reverse transcriptase (TAKARA). Three rounds of 5' RACE reactions were performed with two nested primers (outer, CX1544: 5'-GCTGATGCGCATGAATGAACACTG-3'; inner, CX1545: 5'-CGCGGATCC GAACACTGCGTTTGCTGGCTTTGATG-3') and three gene-specific primers (Supplementary Table 6). The PCR products were gel-purified, cloned (pEASY-T1, Transgen, China) and sequenced.

**Microarray and RNA-seq analysis.** RNA samples used for microarray analysis were prepared from young panicles from stages of MMC to meiosis in wild-type (AnN and ZH11) and *tms5* (AnS-1 and TMS5i-ZH) plants grown at permissive and restrictive temperatures. RNA purification and Affymetrix microarray hybridization were performed by the Capital Bio Corporation (<http://www.capitalbio.com>, Beijing, China). The microarray data were analysed with the Gene Chip Operating software (GCOS 1.4). The different arrays were normalized using DNA-chip analyzer (dChip). The significantly differentially expressed genes between wild-type and *tms5* plants were analysed using the Significant Analysis of Microarray software<sup>48</sup>. Single-end 86-nucleotide RNA-seq reads were aligned to the TIGR6.1 genome sequences using TopHat<sup>49</sup>. Uniquely mapped reads were used for subsequent analysis<sup>50</sup>.

**In situ hybridization analyses.** Specific regions of *RNase Z<sup>S1</sup>*, *Ubl<sub>40</sub>1*- and *Ubl<sub>40</sub>4* were amplified using corresponding primers (Supplementary Table 6) and then transcribed *in vitro* as probes using the DIG RNA labeling kit (Roche, Switzerland). Fresh ZH11 young panicles from different developmental stages were fixed immediately, embedded in paraffin (Sigma-Aldrich, USA) and sectioned to 8-µm thickness. Hybridization and immunological detection were performed according to the previously described method<sup>51</sup>. Briefly, sections were over night incubated at 45 °C with coverslips in hybridization buffer (40 µl per slide) containing the probes. Immunological detection of the hybridized probes was performed using a DIG nucleic acid detection kit (Roche) according to the manual.

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## Author contributions

H.Z. and M.Z. performed most of the experiments; M.Z. organized the figures; Y.Y. bred Zhu1S, developed the NIL mapping population and performed the field experiments; J.L. performed cytological analysis; L.Zhu, D.J., M.F., P.Q., X.H. and X.S. performed the rough mapping and field experiments; Q.L. and L.G. performed the bioinformatics analysis; J.D. and L.Zho. participated in vector construction, plant transformation and transgenic plant analysis; E.N. performed *in situ* hybridization analyses; C.S., J.S., X.W., Q.D. and Z.L. contributed essential reagents; M.C., Y.L., X.C. and C.Z. conceived the experiments; X.C. and C.Z. designed the study and wrote the paper. All authors commented on the manuscript.

## Additional information

**Accession codes:** The microarray and RNA sequencing data have been deposited in the Gene Expression Omnibus (GEO) under the accession codes GSE42367 and GSE42314, respectively.

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