TOUSLED Kinase Activity Oscillates during the Cell Cycle and Interacts with Chromatin Regulators

Hashimul Ehsan2, Jean-Philippe Reichheld, Tim Durfee, and Judith L. Roe*

Program in Molecular, Cellular, and Developmental Biology, Division of Biology, Kansas State University, Manhattan, Kansas 66506–4901 (H.E., J.L.R.); Laboratoire Génome et Développement des Plantes, Université de Perpignan, 66860 Perpignan, France (J.-P.R.); and Department of Genetics, University of Wisconsin, Madison, Wisconsin 53706 (T.D.)

The TOUSLED (TSL)-like nuclear protein kinase family is highly conserved in plants and animals. tsl loss of function mutations cause pleiotropic defects in both leaf and flower development, and growth and initiation of floral organ primordia is abnormal, suggesting that basic cellular processes are affected. TSL is more highly expressed in exponentially growing Arabidopsis culture cells than in stationary, nondividing cells. While its expression remains constant throughout the cell cycle in dividing cells, TSL kinase activity is higher in enriched late G2/M-phase and G1-phase populations of Arabidopsis suspension culture cells, compared to those in S-phase. tsl mutants also display an aberrant pattern and increased expression levels of the mitotic cyclin gene CycB1;1, suggesting that TSL represses CycB1;1 expression at certain times during development or that cells are delayed in mitosis. TSL interacts with and phosphorylates one of two Arabidopsis homologs of the nucleosome assembly/silencing protein Asf1 and histone H3, as in humans, and a novel plant SANT/myb-domain protein, TKI1, suggesting that TSL plays a role in chromatin metabolism.

TOUSLED (TSL) is a widely expressed gene in Arabidopsis required for proper development (Roe et al., 1993, 1997b). TSL is a nuclear Ser/Thr protein kinase composed of a C-terminal catalytic domain and an N-terminal regulatory domain (Roe et al., 1997a). TOUSLED-like kinases (TLKs) are highly conserved and are found in both plants and animals. Humans and mice both have two highly related TLK genes, TLK1 and TLK2 (Yamakawa et al., 1997; Shalom and Don, 1999; Silije et al., 1999; Zhang et al., 1999; Li et al., 2001). Human TLK expression is constitutive throughout the cell cycle, while TLK protein kinase activity (both TLK1 and TLK2) oscillates during the cell cycle, with a peak of activity in S-phase (Silije et al., 1999). Inhibition of DNA polymerase α causes a decrease in TLK activity, suggesting that activation of TLKs is linked to DNA replication (Silije et al., 1999). In addition, recent evidence indicates that TLKs are targets of DNA damage checkpoints (Groth et al., 2003). Mutations in TLK genes cause severe defects in both plants and animals; tsl loss-of-function mutations in Arabidopsis cause flower and leaf abnormalities (Roe et al., 1993), whereas mutations in or suppression of TLK homologs in Drosophila melanogaster and Caenorhabditis elegans cause an early arrest in the embryo (Carrera et al., 2003; Han et al., 2003; T. Ratliff, M. Herman, and J. Roe, unpublished data). Overexpression of TLK1 in mouse cells conferred enhanced resistance to ionizing radiation (Li et al., 2001). A dominant negative form caused missegregation of chromosomes, whereas siRNA-mediated suppression of TLK1 blocked cell division in mouse cells (Sunavala-Dossabhoy et al., 2003).

The target(s) of TSL and the TLKs have remained elusive until recently. Silije and Nigg (2001) identified the human homolog of Asf1 (also known as CIA) as a candidate TLK target. Asf1 is a silencing protein first identified in yeast (Saccharomyces cerevisiae) that recruits histone H3 and H4 during nucleosome assembly (Le et al., 1997; Tyler et al., 1999; Munakata et al., 2000). It functions redundantly with chromatin assembly factor, CAF-1, and can interact directly with a CAF-1 subunit (Tyler et al., 2001; Mello et al., 2002). Asf1 also interacts with protein complexes involved in chromatin remodeling and transcription initiation, such as the SWI/SNF Brahma remodeling complex in D. melanogaster, and with transcription initiation factor IID (TFIID) in humans and yeast (Chimura et al., 2002; Mosshkin et al., 2002). Asf1 likely participates in several silencing pathways in yeast (Singer et al., 1998; Tyler et al., 1999; Sharp et al., 2001). It also acts during DNA repair, where it reassembles nucleosomes following repair (Emili et al., 2001; Hu et al., 2001). The two human Asf1 proteins, Asf1a and Asf1b, bind to and are substrates in vitro of both human TLK1 and TLK2 (Silije and Nigg, 2001). Phosphorylation of the Asf1s is cell cycle regulated and parallels the cell cycle-regulated activity of TLK1 and TLK2, suggesting that Asf1 is a TLK target in vivo (Silije and Nigg, 2001). TLKs have also recently been shown to be inhibited by...
the ATM (ataxia telangiectasia mutated) and the Chk1 protein kinase-DNA damage checkpoint pathways in response to double-strand breaks; possibly, transient inhibition of TLK-dependent activation of Asf1 allows chromatin restructuration prior to nucleosome reassembly during repair (Groth et al., 2003). Histone H3 is another potential TLK phosphorylation target; TLK1 phosphorylates histone H3 in vitro, and it complements a yeast mutation lacking the major histone H3 kinase, Ipl1, even though Ipl1 is not a TLK (Li et al., 2001). However, in C. elegans, it appears that RNA polymerase II large subunit, and not histone H3, may be a key TLK substrate (Han et al., 2003).

These results from animal systems suggest that TLKs participate in chromatin assembly, DNA repair, and in the regulation of gene expression. Although no mutations have been described in Asf1-like genes in Arabidopsis, several mutations in other chromatin assembly genes such as FAS1 and FAS2 (subunits of CAF-1) and cosuppressed AtMSI1 (another CAF-1 subunit) plants show pleiotropic defects in shoot, flower, and in some cases root development (Leyser and Furner, 1992; Kaya et al., 2001; Hennig et al., 2003). tsl loss-of-function mutations cause both leaf and flower defects; only approximately half of the normal complement of floral organs is formed due to a decrease in the number of organ primordia that are initiated, and the young floral meristems are abnormal in shape (Roe et al., 1993). To further examine the possible function of TSL in nuclear function in Arabidopsis, we tested whether TSL protein levels and kinase activity are cell cycle regulated, and we examined the expression pattern of the mitosis specific CycB1;1:GUS transgene (Colon-Carmona et al., 1999) in a tsl mutant. We demonstrate that TSL expression is fairly constant across the cell cycle in Arabidopsis cells, with higher activity seen in the G2/M-phase and G1-phase than S-phase, and that tsl mutants show ectopic expression of the cyclin B gene, CycB1;1. We also demonstrate the interaction of TSL with Asf1b/SGA1, one of two Arabidopsis Asf1 homologs, and with a novel SANT/myb-domain protein TKI1, which, together with other results, suggests that TSL may be involved in gene repression and/or chromatin regulation in Arabidopsis.

RESULTS

TSL and the Cell Cycle

To follow expression, protein levels, and kinase activity, we first prepared a TSL-specific antiserum. Anti-peptide antibody to the C-terminal TSL peptide (C)YNQEDRPDVLTMAQPDPYLA(S was generated and affinity purified as described in “Materials and Methods”. The peptide sequence is not found in kinases other than TLK family members (BLAST analysis, data not shown). The antibody specifically recognizes a GST-TSL fusion protein purified from yeast (Fig. 1A), and detects a single 80-kD protein in Arabidopsis flower protein extracts by western-blot analysis (Fig. 1B). A band at the same M, is not detected in extracts from tsl-1 flowers (Fig. 1B, lane 2), where no TSL protein is expected due to the T-DNA insertion in the gene, which inactivates gene expression. Equal amounts of protein extract were loaded in each lane, as determined by Coomassie Blue staining (data not shown). TSL protein is detected in tsl-2 mutants, as expected, where a single point mutation in the TSL gene affects only one amino acid in the predicted protein.

To determine if TSL is preferentially expressed in dividing cells, Arabidopsis suspension culture cells were harvested from cultures allowed to progress to stationary phase (10 d after subculture) and from cultures in the exponential phase of growth (3 d after subculture) and assessed for TSL expression. tsl-2 cultures in the exponential phase of growth (3 d after subculture) and assessed for TSL expression. TSL mRNA levels are slightly higher when the cells are rapidly dividing, and TSL protein is present in higher steady-state levels in exponentially growing cells than in stationary cells arrested in G0/G1 (Fig. 1C).

We next asked if the cell cycle expression of TSL is regulated in dividing cells. Arabidopsis suspension culture cells were synchronized with an aphidicolin block and release protocol which arrests cells in early S-phase, followed by synchronous progression through the cell cycle after release from the drug block (for review, see Planchais et al., 2000). Aliquots of cells were harvested at time points after the release. The culture contains cells in S-phase from time point 0 through hour 4, as shown by the northern blot detecting histone H4 mRNA levels, as expected (Fig. 2A). As cells progress out of S-phase during hours 6 to 8, the histone H4 mRNA levels had declined. Mitotic index was also determined by DNA staining and chromosome visualization, and mitotic cells were

Figure 1. TSL is more highly expressed in dividing cells. A, Western-blot analysis using α-TSL-peptide affinity-purified antisera of fusion proteins purified from yeast: lane 1, GST-TSL; lane 2, GST. B, Equal amounts of protein extracts from wild-type (lane 1, Wassilewskija ecotype, and lane 3, Columbia ecotype) and tsl (lane 2, tsl-1, and lane 4, tsl-2, in Wassilewskija and Columbia ecotype backgrounds, respectively) flowers were analyzed by western-blot analysis with α-TSL antibody. C, Northern- and western-blot analysis of TSL expression in stationary (st; 10 d after dilution) and exponentially (exp) growing (3 d) Arabidopsis suspension culture cells. Top, Northern-blot analysis of 20 μg total RNA hybridized with TSL cDNA probe (TSL). Hybridization of the same blot after stripping for the presence of 18S rRNA serves as a loading control (middle; 18S). Bottom, TSL protein was detected by western-blot analysis on equal amounts of protein extracts with α-TSL peptide antibody (TSL). Molecular weight markers (numbers) are in kD.
observed in samples from time points between 12 and 16 h, with a peak at 13 h (Fig. 2B). The percentage of cells in mitosis is underestimated with this method, as only the stages with visibly highly condensed chromosomes are counted; in fact, these numbers agree well with other published reports of synchronization in Arabidopsis cells that showed peak levels of 12% mitotic index (Menges et al., 2002; Menges and Murray, 2002). In contrast to histone H4 expression, TSL mRNA levels remain constant at all time points in the synchronized culture. Western-blot analysis on extracts across the time course with the TSL peptide antibody reveals that TSL protein is also present at fairly constant levels (Fig. 2C).

To assess when TSL is active across the cell cycle, we analyzed the protein kinase activity of TSL in immunoprecipitates from aphidicolin-treated and released cultures against the substrate myelin basic protein (MBP). Unfortunately, a highly active contaminating protein kinase showed high background activity in the preimmune controls. To avoid this contaminating kinase, in-gel kinase assays on immunoprecipitates were performed with gels containing MBP incorporated into the gel followed by addition of $[^{32}\text{P}]\text{ATP}$. We first tested purified TSL fusion proteins in the assay (Fig. 3A) and found that GST-TSL showed activity, whereas GST-K438E (contains a single point mutation that converts Lys-438 to Glu and inactivates kinase activity; Fig. 3A, lane 4) and GST alone (data not shown) did not. Also, we could detect TSL activity immunoprecipitated from exponentially growing cells (not shown). The assay was then performed on immunoprecipitates of extracts from aphidicolin-synchronized Arabidopsis suspension culture cells enriched in S-phase, G2/M-phase, and G1-phase (Fig. 3B). TSL could be identified specifically in this assay by its presence at 80 kD and absence in the nonrelated antibody control lanes. TSL is active in all the fractions but shows the highest level of activity in the enriched G2/M-phase sample, and more activity in the G1-enriched sample compared to the S-phase sample (Fig. 3B).

**Figure 2.** TSL expression levels are constant throughout the cell cycle in synchronized cells. A, Top, Suspension culture cells were incubated with aphidicolin for 24 h, and then washed extensively and fresh culture medium added. RNA was isolated from cells at time points (h) following release from aphidicolin block and subjected to northern-blot analysis. Total RNA (20 $\mu$g) was loaded per lane, and the blot was hybridized with a histone H4 cDNA probe. Bottom, The same blot was stripped and hybridized with 18S rRNA sequences as a control for RNA loading (after hybridizing with TSL cDNA in C). B, Samples from aphidicolin synchronized cell populations, as in A, were analyzed for mitotic index. C, Top, Northern-blot analysis of Arabidopsis TSL across the cell cycle. The same blot after stripping as in A was hybridized with a TSL cDNA probe. Bottom, TSL protein accumulation in synchronized Arabidopsis cell suspension culture. Total protein extract (300 $\mu$g) from samples of aphidicolin released cells collected at 2-h time intervals was run on a 10% SDS-polyacrylamide gel and then analyzed by western-blot analysis with $\alpha$-TSL peptide antibody.

**Mitotic Marker Gene Is Misexpressed in tsl Mutants**

 tsl mutants show abnormalities in meristem development (Roe et al., 1993). To assess potential cell cycle effects, the mitotic marker gene, CDG (Colon-Carmona et al., 1999), was introduced into tsl-2 mutants. This marker gene consists of the promoter and 5’ transcribed sequences from the Arabidopsis cyclin B gene CycB1;1 (Hemerly et al., 1992) cloned in frame with a uidA gene, which results in the production of a cyclin destruction box (CDB)-$\beta$-glucuronidase (GUS) fusion protein, CDG. The CDG marker gene allows for the visualization of cells in late G2 and mitosis, as the CycB1;1 promoter is activated in G2 just prior to mitosis, and the CDG enzyme that is synthesized is specifically degraded at the end of M via the CDB (Colon-Carmona et al., 1999; Menges and Murray, 2002). Thus, when plants carrying the marker gene are stained for GUS activity, only the cells in late G2 and M should stain. Inflorescences from tsl-2 plants carrying CDG were stained for GUS activity and compared to wild-type siblings carrying CDG (Fig. 4A).
In wild-type, the GUS activity was confined to a small population of cells in the youngest floral meristems where cell divisions are occurring, as expected (Fig. 4A). As the floral meristems develop and cells stop dividing, expression of the marker gene is lost, with the last expression seen in approximately stage 10 gynoecium on the carpel walls and ovule primordia. In contrast, tsl-2 meristems show greatly increased levels of GUS activity and the staining persists in older flowers, even after organ growth is near completion (Fig. 4A). Eventually, staining is lost in the outer organs. Cauline leaves showed only slightly increased GUS staining in mutant inflorescences compared to wild-type (not shown). This suggests that either more cells in G2 or M are present in the developing tsl flowers, or that deregulation of CycB1;1 expression or turnover is occurring.

To test if the increased expression of the CDG marker gene in tsl floral meristems reflects ectopic expression of the endogenous CycB1;1 promoter, we performed northern-blot analysis on mRNA from wild-type and tsl-1 mutant flowers using a CycB1;1 cDNA as a probe (Fig. 5B). 18S rRNA levels were monitored to control for loading consistency (Fig. 4B).

In comparing the level of transcripts in RNA isolated from wild-type and tsl flowers, a higher steady-state level of CycB1;1 mRNA is present in tsl flowers than in flowers of wild-type siblings. CycB1;1 mRNA levels were also slightly higher in tsl leaves than in wild-type (data not shown). Expression of a second cyclin B gene, CycB1;2, was also tested, as it is much more highly expressed in flowers than CycB1;1 (Day et al., 1996), and in contrast, the levels of CycB1;2 mRNA are comparable in the two samples (data not shown). Thus, CycB1;1 expression appears to be upregulated in tsl mutants. CycB1;1 protein levels could not be assessed, as an antibody that gave consistent results could not be obtained.

### Isolation of TKI1 as a TSL-Interacting Protein

To further investigate TSL function, a two-hybrid screen in yeast was performed to identify proteins that interact with TSL (Fields and Sternglanz, 1994). A TSL bait clone was constructed that encodes a near full-length TSL protein (amino acids 73–688) as a fusion protein with the DNA binding domain (DBD) of Gal4. The clone contains a centromere-based (CEN) origin of replication (Durfee et al., 1999) that was found to...
Leu, and Ura. A total of 1 were selected for by growth on medium lacking Trp, and clones encoding proteins that interact with TSL library in a Gal4-activation domain (ACT) plasmid, TSL bait vector pCD-NQ-TSL, followed by a cDNA thus the N-terminal deletion was used instead.

DBD weakly activates the reporter gene alone, and described previously, full-length TSL fused to Gal4-based plasmid vector in yeast (data not shown). As DBD fusions; right, TKI1-Gal4 ACT fusions) were lifted and assayed for -galactosidase activity. The top two panels were stained overnight and the bottom four for 5 h; blue color indicates activity. NQ, TSL N-terminal deletion containing amino acids 73-688; C-TKI1, C terminus of TKI1, amino acids 546-741; TKI1, full-length TKI1.

Figure 5. TKI1 interacts with active forms of TSL. Yeast transformants carrying two-hybrid plasmids expressing fusion proteins (left, TSL-Gal4 DBD fusions; right, TKI1-Gal4 ACT fusions) were lifted and assayed for -galactosidase activity. The top two panels were stained overnight and the bottom four for 5 h; blue color indicates activity. NQ, TSL N-terminal deletion containing amino acids 73-688; K438E, point mutation changing amino acid 438 from Lys to Glu; AN360, TSL N-terminal deletion containing amino acids 360-688; C-TKI1, C terminus of TKI1, amino acids 546-741; TKI1, full-length TKI1.

overcome the toxic effects of TSL expression from a 2-µ based plasmid vector in yeast (data not shown). As described previously, full-length TSL fused to Gal4-DBD weakly activates the reporter gene alone, and thus the N-terminal deletion was used instead.

The yeast strain YD116 was transformed with the TSL bait vector pCD-NQ-TSL, followed by a cDNA library in a Gal4-activation domain (ACT) plasmid, and clones encoding proteins that interact with TSL were selected for by growth on medium lacking Trp, Leu, and Ura. A total of 1 × 10^6 transformants were screened and putative positives were retested with the bait vector in a different strain, Y153, for activation of a UAS_Gal4::lacZ reporter gene. A number of clones were eliminated as false positives, and several were able to activate transcription alone (data not shown) and also were eliminated. One clone, however, showed strong reporter gene activation that was completely dependent on the presence of the TSL bait vector (Fig. 5). Furthermore, it was shown that the clone, pC-TKI1 (TSL-kinase interacting protein), did not show activation when cotransformed with other DBD-fusion protein vectors such as lamin (data not shown). So, it appears that the TKI1 gene encodes a polypeptide that specifically interacts with TSL.

DNA sequence analysis of the TKI1 cDNA insert revealed that the clone isolated in the two-hybrid screen is not a full-length cDNA and encodes the C-terminal region of the TKI1 polypeptide. PCR on a cDNA library was used to complete the full-length coding region (GenBank accession no. AF530160) corresponding to locus At2g36960, which differed only slightly from the predicted open reading frame (GenBank accession no. AC006922) and a complete cDNA sequence already present in the database (GenBank accession no. AY064068). The TKI1 protein sequence encoded by the cDNA is 741 amino acids (predicted Mr = 82 kD) in length. TKI1 contains several regions that share homology with other proteins; these were identified by BLAST analysis (Altschul et al., 1997) on the GenBank database and by Reverse Position Specific BLAST analysis on the Conserved Domain Database (Marchler-Bauer et al., 2002). The first region is a single SANT/myb domain shared with the human oncogene Myb and other related proteins (Fig. 6A; Aasland et al., 1996). The proteins most homologous to TKI1 include the putative Arabidopsis transcription factors LATE ELONGATED HYPOCOTYL (LHY) and CIRCADIAN CLOCK ASSOCIATED 1 (CCA1), but only the SANT domain is shared. These two factors play a role in circadian rhythm control (Schaffer et al., 1998; Wang and Tobin, 1998). LHY, CCA1, and TKI1 have only one myb repeat (the SANT/myb domain) and thus are not members of the large class of plant R2R3 myb domain proteins, which each have two myb domains (Romero et al., 1998). TKI1 also shares homology with the Polycomb-group D. melanogaster protein cramped (crn; GenBank accession no. Y13674; Yamamoto et al., 1997) in its SANT domain. The SANT domain is found in a number of proteins that are part of complexes that regulate transcription (Boyer et al., 2002, and see “Discussion and Conclusion”). We found that when swapped into the Gal4-DBD plasmid, the C terminus of TKI1 (amino acids 546-741), lacking the SANT/myb domain, acts as a strong transcriptional activator in yeast (Fig. 7).

Several proteins with weaker homology to other regions in TKI1 were found, including a predicted protein on chromosome 4 (At4g39380; GenBank accession no. NM_120098; Fig. 6B), the human protein mitofilin (not shown; GenBank accession no. NM_006839; Odgren et al., 1996), and a predicted protein on chromosome 1 (At1g78650; GenBank accession no. NM_106512; not shown). The importance of these similarities is unknown. TKI1 is expressed in wild-type Arabidopsis flowers, roots, and leaves (Fig. 6C), and a transcript of 3.2 kb is detected.

The interaction of TKI1 and TSL was investigated further using deletion and mutant versions of TSL (Fig. 5). TSL interacts strongly with both the original C-terminal portion of TKI (amino acids 546-741) and the full-length TKI protein in filter lift assays (Fig. 5). In quantitative assays (Fig. 7), TSL interacts approximately fourfold more strongly with the full-length protein. Furthermore, TKI1 interacts only with active kinase forms of TSL and does not bind inactive mutants such as an N-terminal deletion that removes the coiled-coil domain (AN360) or a point mutation
that changes a single amino acid in TSL and inactivates the ATP-binding site in the catalytic domain (Figs. 5 and 7). The K438E mutant is expressed in normal levels in the yeast cell, as has been shown previously by a strong interaction with both itself and wild-type TSL in two-hybrid assays (Roe et al., 1997a).

TSL Interacts with Asf1b/SGA1, an Arabidopsis Asf1 Silencing Factor Homolog

Sillje and Nigg (2001) described the interaction of human TLK1 and TLK2 with two human homologs of the yeast silencing factor Asf1 and their ability to phosphorylate the Asf1-like proteins. Also, TLK activity paralleled Asf1 phosphorylation status in synchronized human cells. Two highly similar homologs of Asf1 are present in the Arabidopsis genome: Asf1a/SGA2 and Asf1b/SGA1 (locus numbers At1g66740 and At5g38110, respectively, and see http://chromdb.biosci.arizona.edu). We tested the latter for interaction with TSL in the two-hybrid system, because a cDNA database entry confirmed its validity as an expressed locus. The coding region of Asf1b/SGA1 was cloned into a two-hybrid vector, creating a fusion protein of Asf1b/SGA1 with the Gal4-DBD (pCD1-Asf1b); this fusion did not show autoactivation (Fig. 7). pCD1-Asf1b was used for two hybrid interaction tests with different forms of TSL cloned as fusion proteins with the Gal4 ACT. o-Nitrophenyl β-D-galactopyranoside (ONPG) assays on cell extracts were used to quantify the interactions observed in colonies stained for β-galactosidase activity (Fig. 7, and data not shown). Asf1b/SGA1 interacted strongly with full-length TSL, and even more strongly with the N terminus of TSL alone (amino acids 1-406), including a clone that was missing the coiled-coil domain (Fig. 7). N-terminal deletions of TSL lacking the first 73, 171, and 360 showed little to no binding activity, as did the active site mutant K438E (Fig. 7). These results suggest that
Asf1b/SGA1 binds to the most N-terminal portion of the TSL protein, which may explain why it was not found in the library screen. The screen used an N-terminal deletion of TSL, lacking the putative Asf1b/SGA1 binding site. Thus, TSL binding to TKI1 and Asf1b/SGA1 appears to be dependent on different domains; Asf1b/SGA1 binds in the most N-terminal domain of TSL whereas TKI1 may bind to the active catalytic C-terminal domain.

To test if TKI1 and Asf1b/SGA1 are potential TSL substrates, in vitro kinase assays were performed using purified GST-TSL and purified His-tagged TKI1 and Asf1b fusion proteins (Fig. 8). Both Asf1b and the C terminus of TKI1 acted as efficient substrates of TSL in vitro, in contrast to purified His-GFP as a control (Fig. 8A, lanes 1–4). The C-TKI1 lane shows not only phosphorylation of the full C-terminal TKI1 protein but also a number of degradation products as well (Fig. 8A, lane 3). A kinase dead mutant of TSL showed no activity (Fig. 8A, lanes 5–8). All of the fusion proteins could be detected by western-blot analysis (Fig. 8B), except full-length TKI1 which did not give a high yield of fusion protein and thus is not shown. GST-TSL also phosphorylated histone H3 in vitro (Fig. 8C).

DISCUSSION AND CONCLUSION

**TSL Expression Is Increased in Dividing Cells and TSL Activity Oscillates during the Cell Cycle**

TSL is expressed in Arabidopsis suspension culture cells, and both TSL mRNA and protein expression levels increase when stationary cells are diluted and cells resume division. To follow expression of TSL during the cell cycle, Arabidopsis suspension culture cells were synchronized with aphidicolin. The results we obtained are highly similar to the results described in the recently reported synchronization of Arabidopsis culture cells (Menges et al., 2002; Menges and Murray, 2002). TSL mRNA and protein expression levels are stable throughout the cell cycle. These data are similar to results obtained in human cell cycle studies where TLK mRNA and protein levels remain...
constant, although we did not observe clear differentially mobile forms of TSL during the cell cycle as is seen in humans (Sillje et al., 1999). Also, TSL expression was not found to be cell cycle regulated in the microarray analysis of synchronized Arabidopsis cells by Menges et al. (2002), confirming these results. TSL is a unique gene in Arabidopsis, and thus its expression can be clearly determined in these experiments. In contrast, however, TSL protein kinase activity oscillates during the cell cycle in Arabidopsis cells, showing higher activity during G2/M-phase and G1-phase compared to S-phase. This contrasts with the data in humans, where TLK activity was found to be the highest during S-phase (Sillje et al., 1999). The reason for this difference is unknown. It is possible that the in-gel kinase assay does not reflect the cellular TSL activity, as the proteins are first denatured during SDS-PAGE and coactivators/inhibitors might have been removed. Posttranslational modifications should be intact, however, and the assay should reflect changes in activity due to such modifications.

We also did not find direct evidence for TSL acting specifically during DNA replication, as was found in humans (Sillje et al., 1999), although TSL is active during S-phase in Arabidopsis. In humans, aphidicolin inhibited TLK activation (Sillje et al., 1999), whereas no such inhibition was seen in the plant cells (H. Ehsan, unpublished data). Perhaps TSL activity in Arabidopsis is influenced by other factors or signals.

Cell Cycle Marker Gene Is Aberrantly Expressed in tsl Meristems

The CDG mitotic marker gene is highly overexpressed in tsl floral meristems. This result may reflect that cells are progressing more slowly through late G2- and/or M-phase when transcription of CycB1;1 is highest (Colon-Carmona et al., 1999; Menges and Murray, 2002), and thus positive cells accumulate in the tsl floral meristems. Alternatively, the greater abundance of the CDG-GUS fusion protein in the tsl meristems could be due to either increased transcription from the CycB1;1 promoter or to decreased turnover of the protein in the cells. Endogenous CycB1;1 mRNA levels are increased in tsl inflorescences, suggesting that the CycB1;1 promoter has been ectopically activated/derepressed. The effect is most pronounced in flowers, consistent with the more dramatic effect of tsl mutations on flower rather than leaf development (Roe et al., 1993). Transgenic plants overexpressing CycB1;1 do not show a tsl phenotype (Doerner et al., 1996), and thus its ectopic expression is probably not the primary defect in tsl mutants. In addition, expression of a nondegradable mutant version of CycB1;1 in cultured tobacco cells does not perturb progression through the cell cycle (Criqui et al., 2000).

Ectopic expression of CycB1;1 in tsl loss of function mutations suggests that TSL may participate in repression of this gene. The increase in TSL activity that we observe in late G2/M-phase (12 h) coincides with the time of peak expression of CycB1;1 in synchronized Arabidopsis cells, as measured by northern-blot analysis (Menges and Murray, 2002). This gene did not show up as one of a group of several mitosis-specific, oscillating mitotic cyclin genes in the microarray analysis of synchronized cells (Menges et al., 2002), probably because it is also expressed at a detectable level in S-phase, and its expression does not decline greatly after the peak in G2/M (Menges and Murray, 2002). In the earlier study, at approximately 14 h following aphidicolin release (end of M), CycB1;1 mRNA levels begin to decline (Menges and Murray, 2002), perhaps due to gene repression. The mechanism of cell cycle-regulated expression of CycB1;1 is not fully understood, although a set of Myb-like transcription factors have been identified in tobacco that participate in both activation and repression of tobacco CycB1;1 (Ito et al., 1998, 2001). Interestingly, TKI is a Myb-like factor, although other than in the single SANT/myb domain, the rest of the TKI1 sequence is not related to these tobacco Myb-like factors, however, which are R1R2R3 myb domain proteins.

Asf1b/SGA1, TKI1, and Histone H3 Are Potential TSL Substrates

Recent reports have shown that Asf1 and histone H3 are potential targets of TLKs in humans (Li et al., 2001; Sillje and Nigg, 2001) and that TLKs may therefore be
involved in chromatin regulation. Very little is known about the two Arabidopsis Asf1-like genes in Arabidopsis. Arabidopsis Asf1b/SGA1 does interact with TSL and is phosphorylated in vitro by TSL and could therefore be a substrate that leads to gene silencing via nucleosome assembly or chromatin regulation via interaction with specific protein complexes. It will be interesting to compare the tsl phenotype with a loss-of-function mutation in the two Asf1-like genes.

TSL also interacts with and can phosphorylate a novel SANT/myb domain protein, TKI1, that can act as a transcriptional activator as a fusion protein in yeast. While SANT/myb domains have been implicated in DNA binding as in Myb (Aasland et al., 1996), more evidence is accumulating that these domains play critical roles in the interaction of proteins in complexes involved in chromatin remodeling by interacting with both histone N-terminal tails and enzymes such as both histone acetyltransferase and histone deacetylase (Boyer et al., 2002; Sterner et al., 2002; Barbaric et al., 2003; Yu et al., 2003). Interestingly, Asf1 also interacts with a MYST acetyltransferase during chromatin silencing in yeast (Meising and Ehrenhofer-Murray, 2001; Osada et al., 2001), and these proteins are potentially in functionally similar complexes. The two interactors appear to bind differently to TSL. Whereas Asf1b/SGA1 binds strongly to the N-terminal regulatory domain, TKI1 binds only to active forms of the kinase and not to the regulatory domain alone. The SANT/myb domain of TKI1 is not required for TSL binding.

In addition to Asf1, histone H3 has been identified as a potential substrate of human TLK1 (Li et al., 2001). Phosphorylation of histone H3 modulates its interaction with chromatin, is likely involved during chromosome segregation, and a number of kinases have been identified that participate in this regulation (Jenuwein and Allis, 2001). TSL does phosphorylate histone H3 in vitro. However, histone H3 phosphorylation was not detectable in extracts of flower buds, either wild-type or mutant, by western-blot analysis using a Ser10-P specific antibody (J.L. Roe, unpublished data), and thus we could not confirm its importance in vivo. It will be interesting to follow the patterns of phosphorylation of all of these proteins in Arabidopsis, especially during nucleosome formation, transcription initiation, chromosome segregation, and DNA repair.

**Chromatin Assembly/Modification Requirement during Plant Development**

A number of genes that are likely involved in chromatin metabolism in Arabidopsis have been identified, and several mutations have been described which inactivate key regulatory genes. The effect is often a pleiotropic disruption of developmental events. Loss of either the FAS1 or FAS2 genes (CAF-1 subunits) results in a highly fasciated shoot meristem and abnormal flower and root development (Leyser and Furner, 1992; Kaya et al., 2001). Another mutation, splayed (syd), has a similar phenotype as tsl mutants, and the SYD gene encodes a SWI/SNF chromatin remodeling factor homolog (Wagner and Meyerowitz, 2002). The syd mutant has altered leaf shape, altered floral organ composition, and abnormal gynoecium and ovule development, and the mutation enhances weak leafy (lfy) and unusual floral organs (ufo) mutations, all features also caused by the tsl mutation (Roe et al., 1993; Wagner and Meyerowitz, 2002; J.L. Roe, unpublished data). tsl also shows a genetic interaction with leunig (Roe et al., 1997b); the LUG gene was identified as a putative transcriptional co-repressor (Conner and Liu, 2000). These as well as other mutations in chromatin regulatory genes, such as those in Polycomb group genes, have shown the importance of transcriptional silencing and chromatin remodeling in plant development (for review, see Wagner, 2003), including the control of flowering time (Noh and Amasino, 2003; Pineiro et al., 2003; for review, see Sung et al., 2003). It is interesting that one of the proteins with the most homology to TKI1 is the Polycomb group protein, cramped, which is likely involved in chromatin regulation during D. melanogaster development (Yamamoto et al., 1997), and further investigation of its function will show if it plays a similar role in Arabidopsis.

**Materials and Methods**

**Arabidopsis Suspension Culture Cells**

Arabidopsis suspension culture cells were grown at 22°C to 27°C, shaken at 120 rpm in the dark in Murashige and Skoog medium supplemented with Gamborg’s B5 vitamins (Sigma, St. Louis), kinetin (1 μg/mL), and α-naphthaleneacetic acid (1 μg/mL), and subcultured weekly.

**Cell Synchronization**

For cell cycle synchronization, 5 μg/mL of aphidicolin (Sigma) was added to a freshly diluted (1:5) stationary culture of Arabidopsis suspension culture cells, and cells were incubated with shaking for 24 h to block the cells in G1 phase. Cells were then washed by pouring through a mesh screen (150 μm), followed by three washing steps, all with sterile solutions. First, cells were washed with 3% sucrose (1 L); second with 1 L Murashige and Skoog medium; and finally with 500 mL Murashige and Skoog medium containing auxin and cytokinin as above. Cells were then suspended in 100 mL medium, still on the screen, and shaken gently but continuously 3 to 5 times for 2 min each. After resuspension in fresh medium, a t = 0 sample was removed, and cells were grown at 24°C with shaking in the dark. Aliquots of 5 to 10 mL were removed at time points following release, and cells were harvested by settling on ice. The medium was removed, and cells were quick-frozen in liquid nitrogen. Separate aliquots of cells were fixed with ethanol:acetic acid (3:1, v/v) and stained with 5 μM 4,6-diamidino-2-phenylindole (Sigma). Mitotic index was determined by counting the proportion of cells in late prophase to metaphase compared to cells in interphase observed by UV fluorescence on a Zeiss Axioskop microscope (Zeiss, Jena, Germany). For each sample, >300 cells were counted. Some samples were stained with propidium iodide (50 μg/mL) or Sytox (0.5 μM), Molecular Probes, Eugene, OR) and observed by confocal fluorescence microscopy.

**RNA and Protein Extraction**

RNA was purified from pelleted cells or quick-frozen (liquid nitrogen) ground tissue with Trizol, according to the manufacturer’s protocol (Life Sciences, St. Louis, MO). Protein extracts were prepared using the Mammalian Protein Extraction Kit (Pierce, Rockford, IL). Protein concentrations were determined by using a Bio-Rad Protein Assay (Bio-Rad, Hercules, CA).
Technologies, Rockville, MD). Total protein extracts were obtained by pelleting cells and resuspending in Lysis250 plus protease inhibitors (50 mM Tris, pH 7.4, 250 mM sodium chloride, 5 mM EDTA, 0.1% Igepal CA 630 [Nonidet P-40 substitute; Sigma], 50 mM sodium fluoride) supplemented with phenylmethylsulfonyl fluoride (100 μg/mL), aprotinin (2 μg/mL), leupeptin (2 μg/mL), pepstatin A (1 μg/mL), dithiothreitol (1 mM), and benzamidine (1 mM) and after freeze-thaw, vortexing with glass beads to solubilize the cells. After centrifugation, soluble protein was quantitated by A₂₈₀ measurement and by running aliquots on SDS-PAGE gels and staining with Coomassie Blue R-250. Flower extracts were obtained by grinding in liquid nitrogen and solubilizing in Laemmli sample buffer plus 6 μL urea.

Antibodies

Anti-TSL antibody was obtained as follows. The peptide (CYNQEDRPDVITMAQDPYLAY) was synthesized, anti-peptide antisera was generated in rabbits, and peptide-specific antibody was affinity purified on peptide-conjugated support by Zymed Laboratories (South San Francisco, CA).

Western-Blot Analysis

Western blot analysis was performed by standard techniques after transfer of SDS-PAGE gels to Immobilon-P membranes (Millipore, Bedford, MA). Blots were incubated overnight with anti-TSL, affinity-purified anti-peptide antiserum (12 μg IgG) or anti-His-horseradish peroxidase mouse monoclonal IgG (sc-8036, Santa Cruz Biotechnology, Santa Cruz, CA; 1:1000) or anti-GST (sc-459, Santa Cruz Biotechnology; 1:1000) rabbit polyclonal antiserum in washing buffer (50 mM Tris, pH 7.5, 150 mM sodium chloride, nonfat dry milk, and 0.05% Tween 20), washed three times for 30 min with washing buffer, incubated for 2 h at room temperature with 1:1000 secondary antibody (goat anti-rabbit IgG conjugated to horseradish peroxidase; Zymed or peroxidase-labeled anti-mouse IgG; Vector Laboratories, Burlingame, CA), washed three times for 30 min each with washing buffer, and rinsed with 50 mM Tris, pH 7.5, 150 mM sodium chloride. Presence of the secondary antibody was detected by chemiluminescence by incubation of blots with SuperSignal (Pierce, Rockford, IL) followed by autoradiography.

Immunoprecipitation

Suspension culture cells were lysed by freezing in liquid nitrogen, grinding on dry ice, followed by vortexing in Lysis250 plus protease inhibitors and glass beads. After centrifugation, total soluble protein extract was incubated with affinity-purified antibody (12 μg immune or preimmune) for 2 h at 4°C with shaking. Affi-Prep Protein A Matrix (Bio-Rad Laboratories, Hercules, CA) was added (30-μL beads), incubated for 30 min at 4°C, and beads were washed three times in extraction buffer. For SDS-PAGE, samples were boiled in Laemmli sample buffer.

In-Gel Kinase Assay

In-gel kinase assays were performed as described in Zhang and Liu (2001). Briefly, SDS-PAGE gels were formed containing MBP (Life Technologies) at 0.25 mg/mL in the resolving gel. After electrophoresis, SDS was removed, proteins were renatured, and the gel was incubated in [³²P-γ]ATP (NEN, Boston, 3,000 Ci/mmol). Following five washes in 5% (w/v) trichloroacetic acid/1% (w/v) sodium pyrophosphate, the gel was dried and autoradiographed.

Induction and Purification of Fusion Proteins and Kinase Assays

GST-fusion proteins were purified as described from yeast (Saccharomyces cerevisiae) after induction with galactose (Roe et al., 1997a). His-tagged fusion proteins were purified from IPTG-induced Erichichia coli strain BL21DE3-phyS using Talon resin (Clontech, Palo Alto, CA). Cells were sonicated in extraction buffer (50 mM HEPES, pH 7.0, 250 mM NaCl plus protease inhibitors), centrifuged 20 min at 12,000g to remove insoluble protein, incubated with resin, washed in extraction buffer, and His-tagged proteins were eluted in 150 mM imidazole. Kinase assays are as described with [³²P-γ]ATP (NEN, 3,000 Ci/mmol; Roe et al., 1997a). Purified histone H3 was obtained from Roche Diagnostics (Indianapolis).

Northern-Blot Analysis

Northern blot analysis was performed using standard techniques. RNA samples were run on formaldehyde-agarose gels, blotted on Hybond N membranes (Amer sham, Buckinghamshire, UK), and hybridized with probe labeled with [³²P]-dCTP (ICN Biomedicals, Costa Mesa, CA; 3,000 Ci/mmol) by random priming using Prime-it II kit (Stratagene, La Jolla, CA) and purified on NucTrap columns (Stratagene). After high stringency washing, blots were subjected to autoradiography. 18S rRNA and histone H4 were detected as described by Roe et al. (1993) and Elsahn et al. (1999). TSL was detected with a full-length cDNA probe, and TK1 was detected with the insert from the two-hybrid clone, p-TK11 (see below). CysB11 coding sequence was synthesized by PCR using the oligonucleotides 5'-GGCAGATC-GAGTTGCACTTCTCGTGCATTGATG-3' and 5'-GTCCATGGCACATGTCGCTTC-3' on the two-hybrid ACT library (see below), digested with BamHI and partially with NolI, and subcloned into a pBluescript-based vector which was sequenced for verification. An NolI fragment was used as a probe which contained the full coding sequence.

Plant Material and GS Activity Staining

tsl-1 and tsl-2 mutants are as described (Roe et al., 1993) and are in the Wassilewskija and Columbia ecotype, respectively. The CysC marker gene in Columbia ecotype was crossed into the tsl-2 background. Interferences from F2 individuals were stained for GS activity using X-glucaric acid as described by Sessions et al. (1999) overnight at 37°C, and destained in 70% ethanol.

Cloning of Asf1b/SGA1, TK1, Fusion Protein, and Two-Hybrid Clones

Oligonucleotides 5'-GGCGCATAGCTGCATCATGATCACC-3' and 5'-GGCGGATCCCTCGTGCAGATGATT-3' were used to clone the coding sequence of Asf1b/SGA1 locus (Asf1b/SGA1) and add an Ndel site and BamHI site by PCR on a cDNA two-hybrid library template (see below), followed by subcloning into pCD1 (Durfee et al., 1999) as a 0.65-kb Ndel-BamHI fragment to form pCD1-Asf1b, which encodes a fusion protein of the DBD of Gal4 with Asf1b/SGA1. The sequence was identical to that of the full-length cDNA sequence (GenBank accession no. AY072420). Some TSL two-hybrid and GST-fusion protein clones and the K438E mutation are as described (Roe et al., 1997). To generate the TSL bait vector, the fragment in p5S-ΔN73-TSL (Roe et al., 1997a) was subcloned into the CEN plasmid pCD1 (Durfee et al., 1999) to obtain the plasmid pCD-NQ-TSL which encodes amino acids 73-688 of TSL as a fusion protein with Gal4-DBD. Details of this cloning and other two-hybrid clones are available on request. pACTII (Bai and Elledge, 1996) was used for subcloning to obtain ACT plasmids. p-TK1 was isolated as a two-hybrid library plasmid in the vector pGAD424 (Ballas and Citovsky, 1997), which encoded amino acids 546-741 of TK1. Full-length TSL cDNA was assembled by PCR amplification of the N-terminal coding sequences using the two-hybrid cDNA library followed by subcloning into pCD-TKI (details available on request). The full-length TSL cDNA was then sequenced on both strands (GenBank accession no. AF350160). The TKI C-terminal coding region of p-TKI and the coding regions of Asf1b and GFP were subcloned into pRSSET (Invitrogen, Carlsbad, CA) to generate His-tagged fusion protein clones; details are available on request.

Two-Hybrid Library Screening and Two-Hybrid Assays

The TSL bait plasmid pCD-NQ-TSL was transformed into YD116 cells (Durfee et al., 1999; selected on Trp plates). A cDNA library in the pGAD424 vector constructed using mRNA from aerial parts of Arabidopsis including flowers (gift of Vitaly Citovsky; Ballas and Citovsky, 1997), which encoded amino acids 546-741 of TK1. Full-length TSL cDNA was assembled by PCR amplification of the N-terminal coding sequences using the two-hybrid cDNA library followed by subcloning into pCD-TKI (details available on request). The full-length TK1 cDNA was then sequenced on both strands (GenBank accession no. AF530160). The TKI C-terminal coding region of p-TKI and the coding regions of Asf1b and GFP were subcloned into pRSSET (Invitrogen, Carlsbad, CA) to generate His-tagged fusion protein clones; details are available on request.

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grown in liquid culture as described (Miller, 1972; Clontech, Yeast Protocols Handbook), and the data are reported in Miller units.

Sequence data from this article have been deposited with the EMBL/GenBank data libraries under accession number AF530160.

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