

cis-Proline-mediated Ser(P)⁵ Dephosphorylation by the RNA Polymerase II C-terminal Domain Phosphatase Ssu72*[§]

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RNA polymerase II coordinates co-transcriptional events by recruiting distinct sets of nuclear factors to specific stages of transcription via changes of phosphorylation patterns along its C-terminal domain (CTD). Although it has become increasingly clear that proline isomerization also helps regulate CTD-associated processes, the molecular basis of its role is unknown. Here, we report the structure of the Ser(P)⁵ CTD phosphatase Ssu72 in complex with substrate, revealing a remarkable CTD conformation with the Ser(P)⁵–Pro⁶ motif in the *cis* configuration. We show that the *cis*-Ser(P)⁵–Pro⁶ isomer is the minor population in solution and that Ess1-catalyzed *cis*-*trans*-proline isomerization facilitates rapid dephosphorylation by Ssu72, providing an explanation for recently discovered *in vivo* connections between these enzymes and a revised model for CTD-mediated small nuclear RNA termination. This work presents the first structural evidence of a *cis*-proline-specific enzyme and an unexpected mechanism of isomer-based regulation of phosphorylation, with broad implications for CTD biology.

The C-terminal domain (CTD)² of the largest subunit of RNA polymerase II (RNAPII) consists of multiple tandem heptad repeats, with the consensus sequence Y¹S²P³T⁴S⁵P⁶S⁷, that serve as a flexible binding platform for nuclear factors (1). CTD-binding partners influence the initiation, elongation, and termination of transcription as well as a myriad of co-transcriptional processes (2). The recruitment of these activities is tied to the progress of the polymerase by cyclic phosphorylation and dephosphorylation of the CTD repeats. For example, phosphorylation at the Ser⁵ position (Ser(P)⁵) predominates at the 5'-end of genes, attracting CTD-binding partners that influence initiation complex formation, mRNA capping, and the transition into elongation (3). As the polymerase moves toward the 3'-end of genes, the level of Ser(P)⁵

declines, whereas phosphorylation at Ser² (Ser(P)²) increases, recruiting nuclear factors responsible for elongation, termination, and 3'-end formation (3). The variable phosphorylation patterns within each heptad repeat, and distributions of these patterns across the full domain create a "CTD code" with a staggering potential complexity (4).

In addition to phosphorylation, proline isomerization provides a second mechanism for regulating the association of CTD-binding partners (5). Due to its cyclic side chain, proline can adopt both *cis* and *trans* conformations about its peptide bond, creating distinct and interconvertible backbone structures with the *cis* isomer being energetically disfavored and therefore less populated (6). Each CTD heptad contains 2 proline residues, and both are preceded by serine residues that are critical targets of phosphorylation. Phosphorylation of Ser-Pro motifs in non-CTD peptides has been shown to modestly stabilize the *cis* form and decrease the rate of isomerization (7). A study with a Ser(P)² CTD peptide reported a *cis* population of <30% for the Ser(P)²–Pro³ motif with very slow interconversion of *cis* and *trans* isomers, on the order of s⁻¹ (8). Relative to the short timescale of transcriptional events, this slow intrinsic exchange presents two structurally distinct and kinetically isolated binding epitopes for each proline residue in the CTD, adding an additional layer of complexity to the CTD code (4).

The biological importance of CTD proline isomerization is unclear, but hints have been provided by studies of the yeast peptidyl prolyl isomerase Ess1 (Pin1 in humans). Peptidyl prolyl isomerases speed the interconversion of proline isomers by several orders of magnitude to restore *cis*-*trans* equilibria at a biologically relevant timescale (6). Regulation of the proline "conformational switch" has been proposed to give these enzymes control over the duration and amplitude of a variety of cellular processes (6). Ess1/Pin1 specifically targets phosphorylated Ser-Pro motifs and influences transcription by RNA polymerase II, likely by regulating the phosphorylation state of the CTD (9). Ess1/Pin1 acts on pCTD peptides *in vitro*, preferentially binding to the Ser(P)⁵–Pro⁶ site over the Ser(P)²–Pro³ site (10, 11). Numerous genetic links between Ess1 and CTD kinases and phosphatases have been reported (5), and abnormal levels of Pin1 activity cause the accumulation of aberrantly phosphorylated forms of the CTD (12). This latter phenotype may be due in part to the influence of Pin1 on the activity of Ser(P)⁵ CTD phosphatase Fcp1 (12–14). Despite these intriguing findings, there is no detailed understanding of how catalyzed proline isomerization modulates the CTD phosphorylation state.

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The atomic coordinates and structure factors (code 3P9Y) have been deposited in the Protein Data Bank, Research Collaboratory for Structural Bioinformatics, Rutgers University, New Brunswick, NJ (<http://www.rcsb.org/>).

[§] The on-line version of this article (available at <http://www.jbc.org/>) contains supplemental Experimental Procedures and Figs. S1–S3.

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² The abbreviations used are: CTD, C-terminal domain; pCTD, phosphorylated CTD; RNAPII, RNA polymerase II; LMW PTP, low molecular weight protein tyrosine phosphatase; snRNA, small nuclear RNA.

Structure of the Ssu72-pCTD Complex

One of the enzymes linked to Ess1 activity is Ssu72, a CTD phosphatase with specificity for the Ser(P)⁵ position (15). Despite its distinct substrate, Ssu72 shares many characteristics of the low molecular weight subfamily of protein tyrosine phosphatases (LMW PTPs) including the signature catalytic motif (CX₅R) and a similar predicted arrangement of secondary structure elements (16, 17). Biologically, Ssu72 influences all three stages of transcription. Ssu72 interacts genetically and physically with initiation factor TFIIB (18, 19) and has been implicated in gene looping, a proposed mechanism for transcription reinitiation that tethers the promoter and terminator regions of a gene (20–22). During elongation, impaired Ssu72 activity leads to increased RNAPII pausing (23). Finally, Ssu72 is a component of the cleavage and polyadenylation factor complex through its association with Pta1 and is essential for the proper termination of small nuclear RNA (snRNA) transcripts (16, 24–27).

Recently, two studies illuminated an *in vivo* connection between Ess1 and Ssu72 (28, 29). Impairing Ess1 catalytic activity in yeast cells resulted in a temperature-sensitive phenotype and the accumulation of Ser(P)⁵ CTD. Both defects were ameliorated by overexpression of Ssu72 (29). A genome-wide analysis of mRNA expression in these cells showed readthrough transcription for a set of genes, mainly snRNAs, with marked similarity to those previously identified in cells with impaired Ssu72 activity (28). The prevailing model explains these results in the following way (28, 29). Phosphorylation at the Ser⁵ position would cause the CTD to adopt predominantly the *cis* form of the Ser(P)⁵–Pro⁶ motif. Near the end of snRNA transcripts, dephosphorylation of Ser(P)⁵ would coordinate the exchange of Nrd1 and Pcf11, two factors involved in Nrd1-dependent transcription termination. However, Ssu72 would have isomeric specificity for the less populated *trans* form of the Ser(P)⁵–Pro⁶ motif and would require Ess1 activity to efficiently process the entire pool of available substrate. Consequently, reduced catalysis by Ess1 would lead to an increased level of Ser(P)⁵ CTD, causing an improper localization of Ser(P)⁵ CTD-bound Nrd1 and a concomitant blocking of Ser(P)² CTD-mediated Pcf11 recruitment and resulting in readthrough transcription. This model is supported by the fact that all reported enzymes that target Ser(P)–Pro motifs with isomeric specificity recognize the *trans* configuration. On the other hand, there is no evidence that the *cis*-Ser(P)⁵–Pro⁶ isomer of the CTD becomes the major population upon phosphorylation (7, 8).

To investigate the mechanism of proline isomer-based regulation of CTD phosphorylation states, we determined the structure of Ssu72 in complex with its Ser(P)⁵ CTD substrate. The structure reveals an enzyme fold that contains the conserved LMW PTP scaffold and unique additions and a surprising substrate conformation with the Ser(P)⁵–Pro⁶ motif of the CTD adopting the *cis* configuration. We show that the *cis* isomer is a minor population of substrate in solution and that Ess1 significantly enhances Ssu72 activity by catalyzing rate-limiting *cis-trans* interconversion. Together, this work presents structural and kinetic evidence for a unique *cis*-proline-specific enzymatic activity and a fascinating explanation for the *in vivo* relationship of Ess1 and Ssu72, with broad implications for the regulation of the CTD code and its role in coordinating co-transcriptional events.

EXPERIMENTAL PROCEDURES

Molecular Cloning and Protein Purification—The full-length WT *Drosophila melanogaster* Ssu72 gene was PCR-amplified from cDNA (Open Biosystems, clone ID: RE29729), digested, and ligated into a pET15b vector (EMD Biosciences, Inc.) between the NdeI and BamHI restriction sites. Point mutants of Ssu72 and Ess1 were prepared using the QuikChange site-directed mutagenesis kit (Stratagene). The correct sequences for all constructs were verified by DNA sequencing. Protein expression and purification procedures are described in the [supplemental material](#).

NMR—Isotopically enriched proteins for NMR studies were overexpressed in M9 minimal media with [¹⁵N]NH₄Cl and [¹³C]glucose (Cambridge Isotope Laboratories, Inc.) as the sole nitrogen and carbon sources. NMR experiments were conducted at 30 °C using Varian INOVA 600- or 800-MHz spectrometers. Backbone resonances for C13D/D144N *D. melanogaster* Ssu72 (dSsu72) were assigned using a ²H/¹³C/¹⁵N-labeled sample with standard three-dimensional triple-resonance experiments.

Crystallization and Structure Determination—The Ssu72-pCTD complex was prepared by incubating a 5:1 molar ratio of pCTD peptide (4.17 mM) to C13D/D144N dSsu72 (0.83 mM) for 30 min on ice. The complex was crystallized by hanging drop vapor diffusion; 1 μl of protein sample was mixed with 1 μl of 22% (w/v) PEG monomethyl ether 550, 100 mM imidazole, pH 6.5, and 150 mM DL-malic acid and suspended over a reservoir of the same solution. Equilibration at 4 °C for ~5 days produced 50 × 50 × 200-μm rhombic-shaped crystals, which were cryoprotected by dipping into a solution of mother liquor with 25% ethylene glycol before flash-cooling in liquid nitrogen. Methods for data collection, phasing, and model refinement are provided in the [supplemental material](#).

Enzyme Kinetics—Reactions with synthetic pCTD substrate (see Fig. 4, A–D) were followed by measuring phosphate release with a standard Malachite Green reagent (30). For experiments with WT Ess1, 350 μl-reactions were set up in 50 mM HEPES, 20% glycerol, 1 mM EDTA, 2 mM DTT buffer at pH 6.5 (kinetics buffer) with 60 μM pCTD peptide and varying amounts of Ess1. After incubating for 10–15 min, each reaction was started by the addition of 4 μM WT dSsu72 (or 20 μM for the faster timescale reactions in [supplemental Fig. S3](#)). 50-μl aliquots were taken each minute and mixed with 50 μl of Malachite Green reagent in a 96-well plate, which was placed in the dark for 30 min to allow for color development. Color was quantified by measuring A_{630 nm} with a SpectraMax microplate reader (Molecular Devices). Absorbance values were converted to phosphate content using a standard curve made with Na₃PO₄ solutions of known concentrations. The same protocol was followed for experiments with Ess1 mutants, with reactions containing 1 μM of each mutant. Preparation and characterization of the substrate peptide are described in the [supplemental material](#), along with a description of kinetic assays with full-length pCTD substrate (see Fig. 4, E and F).

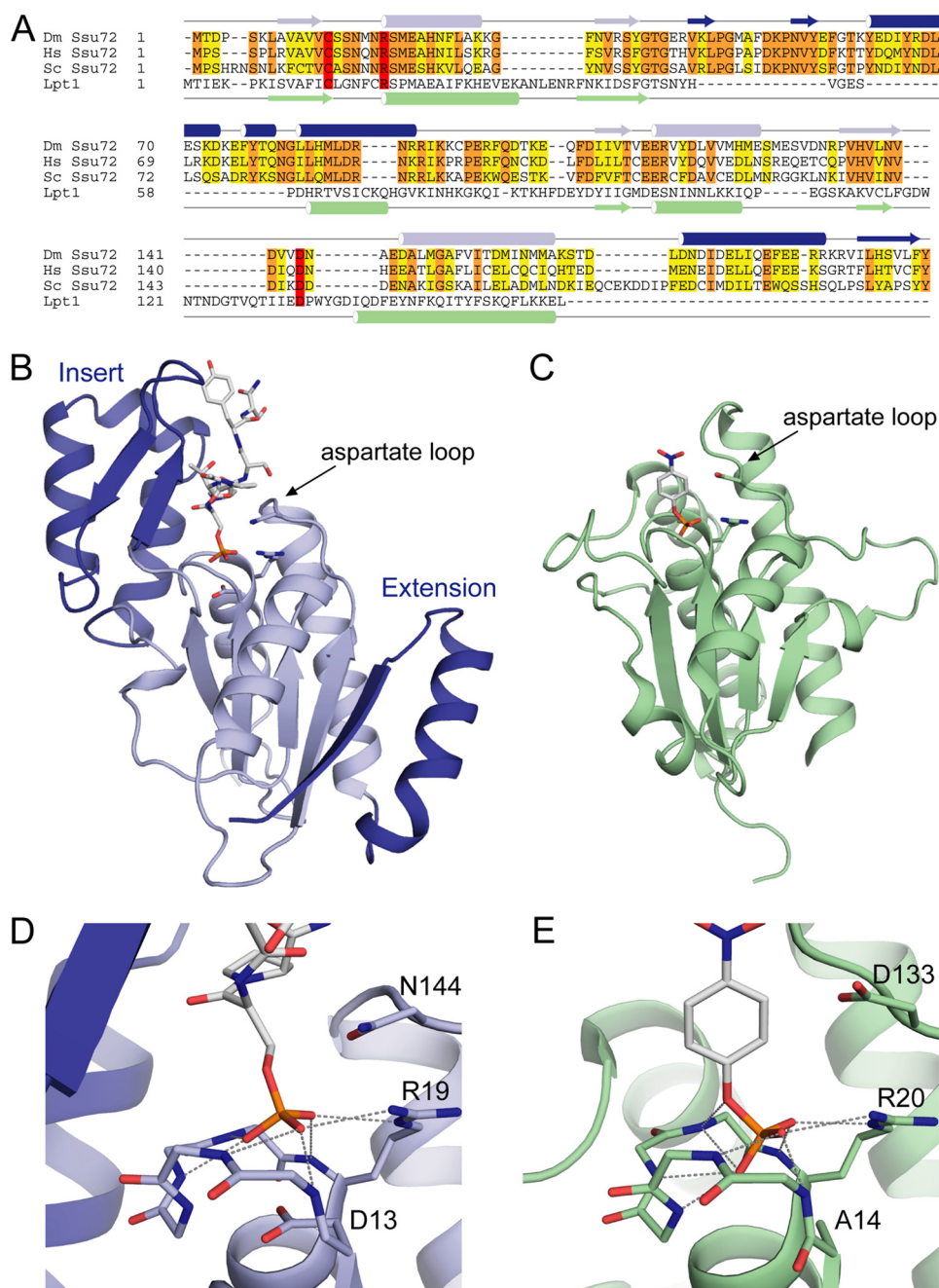


FIGURE 1. Structure of Ser(P)⁵ CTD phosphatase Ssu72. *A*, sequence alignment of Ssu72 orthologues and the LMW PTP Lpt1. Ssu72 proteins from *D. melanogaster* (*Dm Ssu72*), human (*Hs Ssu72*), and *S. cerevisiae* (*Sc Ssu72*) are shown with conserved and similar residues highlighted in orange and yellow, respectively. α -Helices and β -strands from the structures of dSsu72 and Lpt1 (PDB code 1D1Q) are denoted with cylinders and arrows, respectively. The secondary structure elements of Ssu72 that comprise the two additions to the LMW PTP scaffold are colored dark blue to match the structure in *B*. Catalytic residues are highlighted in red. *B*, ribbon diagram of the dSsu72-pCTD complex. Additions to the LMW PTP fold are colored dark blue. The substrate peptide and catalytic residues Cys¹³ (mutated to Asp¹³), Arg¹⁹, and Asp¹⁴⁴ (mutated to Asn¹⁴⁴) are shown as sticks. *C*, ribbon diagram of the LMW PTP Lpt1. The Lpt1 structure shows a catalytic mutant bound to the substrate *p*-nitrophenyl phosphate (PDB code 1D1Q). The substrate and catalytic residues Cys¹⁴ (mutated to Ala¹⁴), Arg²⁰, and Asp¹³³ are shown as sticks. *D* and *E*, active site close-ups of substrate-bound dSsu72 (*D*) and Lpt1 (*E*). The backbone of the phosphate-binding loops is shown as sticks along with important catalytic residues. Hydrogen bonds are represented by gray dashed lines.

RESULTS

Ssu72 Adopts the Conserved LMW PTP Fold with Unique Additions—Previous kinetic experiments suggest that Ssu72 shares the catalytic mechanism of protein tyrosine phosphatases (17, 31), a two-step reaction that requires: 1) the CX₅R catalytic motif with the cysteine side chain as a negatively charged thiolate group and 2) an aspartate residue on a flexible loop positioned near the active site (“the aspartate loop”).

In the first step, the thiolate of the catalytic cysteine attacks the substrate phosphorous atom to generate a phosphoenzyme intermediate, and in the second step, the aspartate protonates the leaving phosphate to regenerate the enzyme. By screening orthologs, we identified dSsu72 as a highly stable and soluble enzyme with significant sequence similarity to *Saccharomyces cerevisiae* Ssu72 (Fig. 1A). Mutation of both the catalytic cysteine (C13D) and the catalytic aspartate

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TABLE 1**Data collection and refinement statistics**

Values in parentheses are for the highest resolution shell.

Data collection	
Resolution range (Å)	50–2.10 (2.18–2.10)
Space group	R3
Unit cell dimensions (Å)	157.5, 157.5, 118.8
Completeness (%)	99.9 (99.8)
Unique reflections	63898 (6361)
Redundancy	3.9
R_{merge} (%)	6.1 (32.6)
$I/\sigma I$	23.7 (4.1)
Refinement	
$R_{\text{work}}/R_{\text{free}}$ (%)	20.90/24.11
Number of atoms ^b	
Protein	6360
Peptide	236
Water	658
Other molecules ^c	31
B -factors (Å ²) ^d	
Protein	33.38
Peptide	36.53
Water	34.55
Other molecules ^c	32.83
r.m.s.d. ^e from ideal geometry	
Bond lengths (Å)	0.003
Bond angles (°)	0.748
Ramachandran plot	
Favored (%)	97.2
Allowed (%)	99.9

^a R_{free} was calculated with the 5% of the data randomly omitted from refinement.^b Nonhydrogen atoms. Riding hydrogens were used in refinement and are included in the deposited structure.^c Imidazole and a PEG fragment.^d Only nonhydrogen atoms included.^e r.m.s.d., root mean square deviation.

(D144N) fully abolished the activity of dSsu72, and an NMR titration of ¹⁵N-labeled C13D/D144N dSsu72 with Ser(P)⁵ CTD peptide revealed tight binding (slow exchange on the NMR timescale) and significant perturbations of residues near the active site loop and the aspartate loop (supplemental Fig. S1).

To obtain a high resolution model of the Ssu72-pCTD interaction, we used x-ray crystallography to determine the structure of C13D/D144N dSsu72 in complex with a synthetic, 7-residue Ser(P)⁵ CTD peptide containing the minimal binding epitope (32) with an additional residue at each end (Ac-PtPSPSYS-NH₂, where the pS indicates Ser(P)). Phasing was carried out by molecular replacement using the apoWT dSsu72 structure deposited by the Northeast Structural Genomics Consortium (Protein Data Bank (PDB) code 3FDJ),³ and the model was refined at 2.1 Å resolution (Table 1).

The Ssu72 structure contains the LMW PTP fold and two sizable additions (Fig. 1, A–C, PDB code 3P9Y). The conserved scaffold consists of a central, four-stranded, parallel β -sheet flanked by two α -helices on one side and a third α -helix on the other and is represented in Fig. 1 by the structure of the LMW PTP Lpt1 in complex with its substrate *p*-nitrophenyl phosphate (33). The first addition to the Ssu72 fold is an ~60-residue insertion after the second β -strand that forms a subdomain of two β -strands packed against three α -helices. This insert structure sits immediately adjacent to

the active site loop and contributes to substrate binding (see below). The corresponding region in the LMW PTP fold contains ~25 residues and forms a single α -helix that packs against the back of the central β -sheet and is tethered by long loops that interact with substrate. The second addition is an ~30-residue C-terminal extension that forms a helix-turn-strand motif with the β -strand bonding in an anti-parallel fashion to the central β -sheet. Another distinguishing characteristic of Ssu72 is the length of its aspartate loop. The α -helix following the aspartate loop is one turn shorter in Ssu72, which allows a commensurate shortening of the aspartate loop and increases active site accessibility from the side opposite the insert subdomain.

The active site architecture of Ssu72 is very similar to LMW PTPs, providing structural support for a common mechanism of dephosphorylation. Fig. 1, D and E, show a close-up view of phosphate binding for the Ssu72-pCTD complex and the LMW PTP Lpt1-*p*-nitrophenyl phosphate complex (33). The conformations of active site loops in these structures are nearly identical, with the backbone amide groups forming a series of hydrogen bonds to the substrate phosphate group and the guanidinium group of the catalytic arginine extending this hydrogen bonding to form a complete circle. Despite the different length of aspartate loops, the positions of the catalytic aspartate side chains (D144N in the Ssu72 structure) are very similar. Although the carboxylate of the Lpt1 aspartate is turned away from the active site, presumably due to electrostatic repulsion with the substrate phosphate group, in the Ssu72 structure, the catalytic D144N side chain is pointed toward the phosphate. Another difference between the active sites is the position of the substrate phosphate. In the Lpt1 structure, the catalytic cysteine is mutated to alanine, and this loss of negative charge allows the phosphate to bind deeper in the active site, close to the position predicted for the phosphoenzyme intermediate (33), whereas the phosphate group in the Ssu72 structure is held ~0.7 Å higher by the negative charge of the C13D side chain.

Ssu72 Recognizes a cis-Proline Substrate—The electron density for the pCTD peptide is excellent and extends almost completely through both termini (Fig. 2A). While the substrate phosphate group is anchored in the active site with strong hydrogen bonds, the insert subdomain creates a deep and narrow recognition cleft that forces the C-terminal end of the CTD to extend away immediately from the phosphate in a nearly opposite direction. This constrained conformation is made possible by a *cis* isomer of the Ser(P)⁵-Pro⁶ peptide bond ($\omega = -2.42^\circ$) that provides an abrupt turn in the substrate backbone. The uniqueness of the *cis*-Ser(P)⁵-Pro⁶ isomer and its pronounced effect on the substrate conformation are readily apparent by comparison with the structure of Ser(P)⁵ CTD bound to the CTD phosphatase Scp1 (PDB code 2GHT) in Fig. 2, C and D (34). In the Scp1 complex, the *trans* isomer of the Ser(P)⁵-Pro⁶ bond provides an extended backbone configuration, in contrast to the sharp kink produced by the *cis*-Ser(P)⁵-Pro⁶ isomer.

To achieve its unusual isomeric specificity, Ssu72 must stabilize the high energy *cis*-proline substrate conformation (Fig. 2B). This is accomplished in part by facilitating the formation

³ A. P. Kuzin, Y. Chen, J. Seetharaman, F. Forouhar, Y. Chinag, Y. Fang, K. Cunningham, L.-C. Ma, R. Xiao, J. Liu, M. C. Baran, T. B. Acton, B. Rost, G. T. Montelione, J. F. Hunt, and L. Tong, unpublished results.

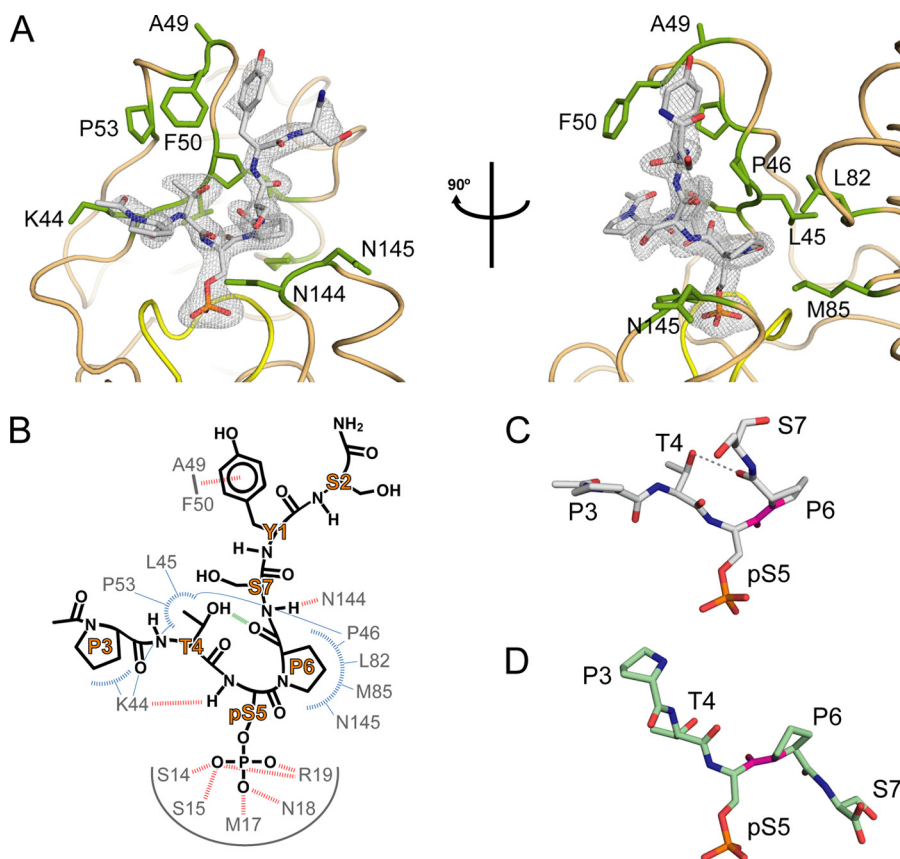


FIGURE 2. **Binding of *cis*-proline pCTD substrate by Ssu72.** *A*, orthogonal views of pCTD conformation and binding. The enzyme structure is shown in *tan* with *yellow* highlighting for the active site loop, and residues that form the substrate recognition surface are labeled and shown as *green* sticks. The *gray* mesh represents the $F_o - F_c$ omit map density (contoured at 3.0σ) surrounding the substrate. *B*, schematic of the Ssu72-pCTD interaction. Hydrogen bonds and van der Waals contacts are shown as *red* and *blue* dashed lines, respectively. The *green* line represents an intramolecular hydrogen bond in the pCTD substrate. *C* and *D*, examples of *cis* and *trans* substrate configurations. The conformations of Ser(P)⁵ CTD peptides in complex with dSsu72 (*C*) and phosphatase Scp1 (*D*) are shown with the Ser(P)⁵-Pro⁶ peptide bond highlighted in *magenta*. For clarity, only residues Pro³ to Ser⁷ are shown. The *cis* isomer is stabilized by an intramolecular hydrogen bond (*gray* dashed line) and causes a severe kink in the CTD backbone.

of an intramolecular hydrogen bond between the substrate Thr⁴ side-chain hydroxyl and the Pro⁶ backbone carbonyl. The Thr⁴ side chain is positioned by its methyl group binding in a hydrophobic pocket formed by the side chains of Lys⁴⁴, Pro⁴⁶, and Pro⁵³ and the backbone of Leu⁴⁵. Hydrogen bonding with the Thr⁴ hydroxyl forces the Pro⁶ residue to adopt the *cis* isomer, which is further stabilized by strong van der Waals interactions between the Pro⁶ ring and a hydrophobic patch of Ssu72 formed by the side chains of Pro⁴⁶, Leu⁸², and Met⁸⁵ and the backbone of Asn¹⁴⁵. Between the Thr⁴ and Pro⁶ residues, the Ser(P)⁵ backbone amide forms a hydrogen bond with the Lys⁴⁴ backbone carbonyl of Ssu72, while the Ser(P)⁵ side chain sits in the active site as described above. At the N-terminal end of the substrate peptide, the Pro³ residue makes minor van der Waals contacts with the Lys⁴⁴ side chain of Ssu72. Toward the C-terminal end, additional isomer-specific binding energy is provided by an intermolecular hydrogen bond between the substrate Ser⁷ main-chain amide group and the Asn¹⁴⁴ backbone carbonyl. While the side chains of the Ser⁷ and Ser² residues are largely solvent-exposed, the phenol ring of the intervening Tyr¹ side chain forms an aromatic-amide stacking interaction with the Ala⁴⁹-Phe⁵⁰ peptide bond of Ssu72 (35), causing a reorientation of the loop between the two β -strands of the insert subdomain in the

complex when compared with the apo structure (supplemental Fig. S2).

Ess1-catalyzed Proline Isomerization Stimulates Ssu72 Activity—A previous study of peptides with Ser(P)-Pro motifs showed that the *cis* isomer is less favored than the *trans* form, with populations of 12–20% depending on the adjacent residues (7). Similar work with a Ser(P)² CTD peptide reported a *cis* population under 30% for the Ser(P)²-Pro³ isomer (8). Although these results suggest that phosphorylation of the Ser⁵-Pro⁶ CTD motif is unlikely to change the minority status of the *cis* isomer, no direct measurements have been reported. To address this point, we measured the fractions of *cis*- and *trans*-Ser(P)⁵-Pro⁶ in our substrate, a synthetic peptide with the sequence Ac-PTpSPSYS-NH₂ (where the pS indicates Ser(P)), which is the same peptide used for crystallization. A natural abundance ¹³C-heteronuclear single quantum correlation spectrum shows two sets of signals for each proline C β and C γ atom, corresponding to the *cis* and *trans* forms (Fig. 3A). However, the substrate peptide contains 2 prolines, making it difficult to unambiguously assign peaks to the Pro³ and Pro⁶ residues. By collecting the same experiment with a shorter peptide containing only the Pro⁶ residue (Fig. 3B), we were able to assign the Pro⁶ resonances and readily map them to the longer peptide. Populations were calculated using the

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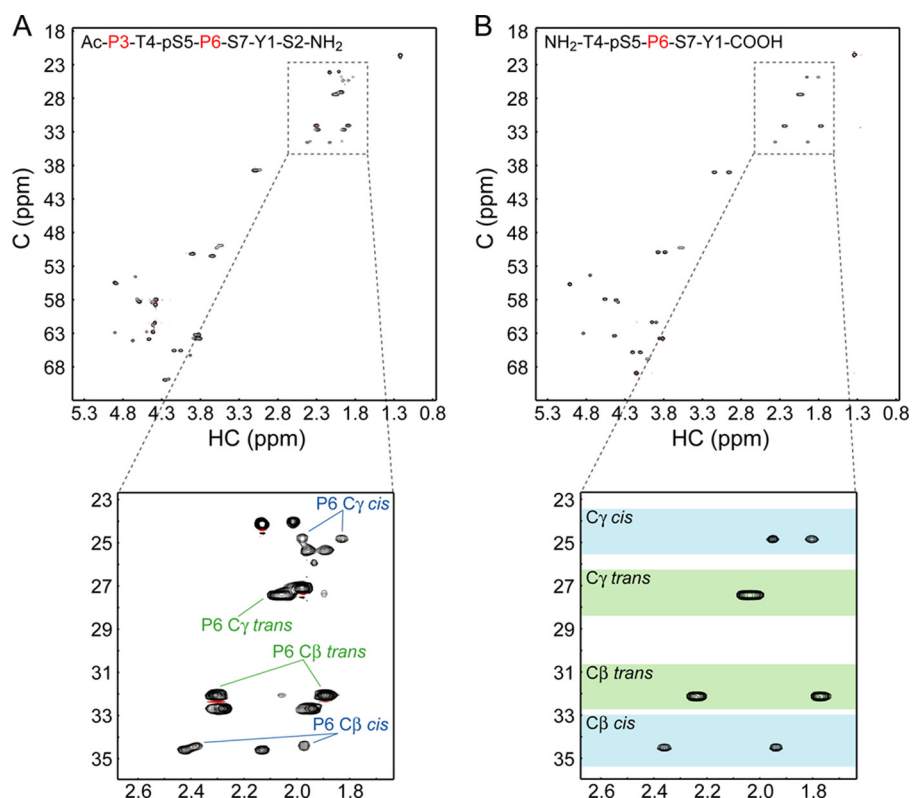


FIGURE 3. Measurement of proline isomer populations for Ser(P)⁵ CTD substrate. *A* and *B*, natural abundance ¹³C-heteronuclear single quantum correlation spectra were collected for the substrate peptide (*A*) and a shorter pCTD peptide with only one proline residue (*B*). Close-up views depict the proline β- and γ-region in each spectrum. Colored boxes indicate the range of chemical shifts (mean ± S.D.) for each proline atom in either the *cis* (blue) or the *trans* (green) form (43). pS5, Ser(P)⁵. C, carbon; HC, proton.

average ratio of *cis* and *trans* peak volumes for the Pro⁶ Cβ and Cγ atoms. This gave a fractional population of 12.4% for the *cis* isomer, verifying that the Ser(P)⁵–Pro⁶ motif is predominantly in the *trans* configuration.

Given that Ssu72 recognizes the minor population of *cis*-Ser(P)⁵–Pro⁶ isomers, the extremely slow interconversion of *cis*-*trans*-proline (s⁻¹) may present a rate-limiting step for dephosphorylation. To test this possibility, we examined the effect of catalyzed proline isomerization on Ssu72 activity. Reactions were monitored by measuring phosphate release with a Malachite Green-based assay. In a reaction with 60 μM pCTD and 4 μM WT dSsu72, less than half of the input substrate was dephosphorylated in 5 min (Fig. 4A). To speed the interconversion of proline isomers, we added increasing amounts of Ess1, a CTD proline isomerase that acts on Ser(P)⁵–Pro⁶ motifs. Ess1 significantly stimulated the activity of Ssu72, with a 2 μM Ess1 concentration causing an ~2-fold increase in reaction completion over 5 min. The increase in Ssu72 activity was saturable (Fig. 4B), and a concentration of 10 μM Ess1 was sufficient to maximize the completion percentage at all time points (data not shown). To ensure that Ssu72 was capable of completely dephosphorylating the pCTD substrate in the absence of Ess1, we followed the reaction without Ess1 over the course of 25 min (Fig. 4C), which showed that ~90% of substrate is dephosphorylated in 20 min, 4-fold slower than the reaction with 2 μM Ess1. This behavior implies a kinetic effect on dephosphorylation, with Ess1-catalyzed isomerization improving the availability of

cis-proline substrate for Ssu72 until the rate of isomer interconversion exceeds the rate of pCTD dephosphorylation.

To confirm that catalyzed isomerization is the cause of Ssu72 stimulation, we performed identical experiments with four catalytically impaired Ess1 mutants. Three of these mutations are in the catalytic domain of Ess1 (C120S, S122P, and H164R), whereas the fourth disrupts phosphate binding in the WW domain (K68A). All four mutations significantly decreased the enhancement of Ssu72 activity (Fig. 4D). To ensure that this result is not caused by reduced thermostability, we measured melting temperatures for each Ess1 mutant by circular dichroism (data not shown). Only the H164R mutation caused a large decrease in stability (melting temperature of 32 °C) that may contribute to its impaired stimulation. Together, these studies demonstrate that Ess1 enhances Ssu72 activity by catalyzing proline isomerization of the pCTD substrate.

Finally, to evaluate the effect of Ess1 on Ssu72 activity toward its natural substrate, we performed *in vitro* reactions using the full-length, 26-repeat CTD from *S. cerevisiae*. A GST-*S. cerevisiae* CTD-His fusion construct was hyperphosphorylated with the yeast CTD kinase CTDK-1 (36), and dephosphorylation by Ssu72 was monitored by measuring substrate depletion with a Ser(P)⁵-specific antibody in reactions with and without 100 μM Ess1 (Fig. 4, E and F). As in the reactions with synthetic pCTD substrate, Ess1 greatly facilitates dephosphorylation by Ssu72, decreasing the time required to reach ~90% completion by 20-fold (from 20 to 1 min).

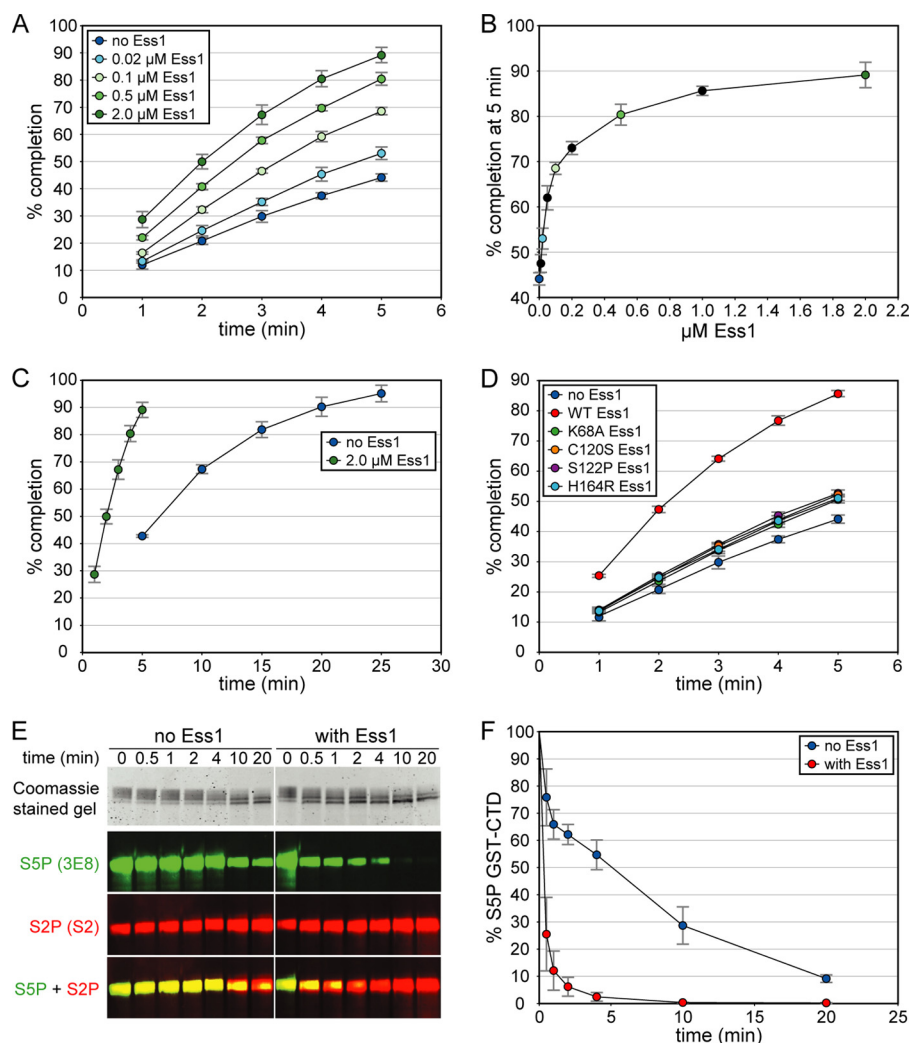


FIGURE 4. Ess1 stimulates pCTD dephosphorylation by Ssu72. *A*, the activity of Ssu72 was monitored with various concentrations of the proline isomerase Ess1. The maximum stimulation was ~2-fold under these conditions. In *A–D*, each point represents the average from three independent reactions, and *error bars* denote S.D. *B*, Ess1 enhancement of Ssu72 activity is saturable. The last points of the reactions in *A* are plotted versus Ess1 concentration, with *black circles* representing reactions omitted from *A* for clarity. *C*, pCTD dephosphorylation by Ssu72 reaches completion without Ess1. The *no Ess1* reaction in *A* was monitored over 25 min and reached ~90% completion in 20 min, 4-fold slower than the reaction with 2 μM Ess1. *D*, four catalytically impaired Ess1 mutants fail to significantly stimulate Ssu72. Reactions contained 1 μM of each Ess1 protein. Residual enhancement corresponds to a 50–100-fold reduction of WT Ess1 activity (compare with *A*). *E* and *F*, Ess1 stimulates Ssu72 dephosphorylation of full-length pCTD. The 26-repeat *S. cerevisiae* CTD was hyperphosphorylated *in vitro* and used as a substrate in reactions monitored by Western blotting with Ser(P)⁵ (S5P)-specific antibodies. As a control, Ser(P)² (S2P) levels were also measured; the apparent increase in the S2 signal is likely due to a higher affinity of the S2 antibody for singly phosphorylated Ser(P)² heptads over doubly phosphorylated Ser(P)²/Ser(P)⁵ heptads, as reported for other Ser(P)²-specific antibodies (44). Ser(P)⁵ CTD levels were quantified with infrared imaging, and the resulting substrate depletion curves are shown in *F*. Each point represents the average of three independent reactions, and *error bars* denote S.D.

DISCUSSION

Ssu72 as a New Family of LMW Phosphatases—The CTD phosphatase Ssu72 is an intriguing enzyme. Despite being built on the scaffold of a protein tyrosine phosphatase, its activity is directed at phosphorylated serine residues in the 5th position of the CTD heptad, with isomeric specificity for the *cis* configuration of the Ser(P)⁵–Pro⁶ peptide bond. Our complex structure also suggests a conserved catalytic mechanism, which agrees well with prior kinetic studies (17). Of the two additions to the LMW PTP scaffold, the insert subdomain plays a major role in the unique substrate specificity of Ssu72, contributing nearly all of the residues that form the non-phosphate substrate recognition surface and severely restricting access to the active site. Very recently, Xiang *et al.* (37)

reported a crystal structure of human Ssu72 in complex with pCTD substrate and symplekin (homologous to Pta1 in yeast), a scaffold protein in the cleavage and polyadenylation complex, which shows that the C-terminal extension of Ssu72 forms the symplekin-binding site. The Ssu72 fold, pCTD conformation, and effect of catalyzed proline isomerization on dephosphorylation are consistent with the data presented in this study.

Substrate Specificity of Ssu72—Prior kinetic work on Ssu72 has demonstrated a strict specificity for the Ser(P)⁵ position of the CTD heptad (15, 32), in contrast to CTD phosphatases Scp1 and Fcp1, which have only preferential activity for Ser(P)⁵ and Ser(P)², respectively. Interestingly, both potential sites are Ser(P)–Pro motifs, meaning that 2 of the 4

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residues recognized by Ssu72 are identical in the Ser(P)² and Ser(P)⁵ substrates. Therefore, discrimination must be based on the Thr⁴ and distal Tyr¹ positions. For a Ser(P)² substrate, the Tyr¹ position would be substituted by serine, causing a decrease in binding energy from the loss of the aromatic amide stacking interaction. However, the Thr⁴ position is likely the more important determinant of substrate specificity. The Thr⁴ residue would be replaced by tyrosine in a Ser(P)² substrate, which would disrupt substrate recognition in two major ways. First, the favorable hydrophobic interactions with the insert subdomain would be lost, and the accommodation of the large phenol group would require a significant reconfiguration of the substrate backbone to prevent steric clashes. Second, this substrate would be unable to replicate the intramolecular hydrogen bond we observe between the Thr⁴ hydroxyl and the Pro⁶ backbone carbonyl, making the already unfavorable *cis* conformation even less energetically stable.

Although the exact degree of isomeric specificity of Ssu72 is unclear, our structural and kinetic studies demonstrate a strong preference for the *cis*-Ser(P)⁵-Pro⁶ conformation. First, modeling the *trans*-Ser(P)⁵-Pro⁶ isomer in the Ssu72-pCTD complex structure leads to large steric clashes between the pCTD backbone and the side chains of the C-terminal α -helix of the insert subdomain. Also, if the activity toward *cis* and *trans* isomers were comparable, there would be no stimulation of dephosphorylation by Ess1. Finally, additional support for strong *cis*-proline specificity comes from a kinetic study that measured activity toward a set of Ser(P)⁵ CTD substrates with an alanine substitution at each heptad position (32). These experiments showed that CTD positions Thr⁴, Pro⁶, and Tyr¹ of the following heptad were critical for substrate recognition, a result that is consistent with the intermolecular contacts in our complex structure. Importantly, the largest decrease in Ssu72 activity was observed with the P6A mutant substrate, an \sim 30-fold reduction when compared with wild type. This likely reflects not only the loss of binding energy for the proline ring but also the reduced propensity of the Ser(P)⁵-Ala⁶ peptide bond to adopt a *cis* configuration.

Regulation of Isomer-specific Enzymes—To the best of our knowledge, Ssu72 is the first example of an enzyme with specificity for the *cis* isomer of proline. Isomeric specificity for *trans*-Ser(P)-Pro motifs has been previously established for two enzymes. The serine/threonine phosphatase PP2A dephosphorylates *trans*-Ser(P)-Pro motifs in its substrate Cdc25C and Tau proteins and shows increased activity in reactions that include Pin1 (40). Likewise, the serine/threonine kinase ERK2 phosphorylates *trans*-Ser-Pro motifs in the RNase T1 substrate and is also stimulated by proline isomerase activity (41). Importantly, catalyzed proline isomerization only increases the activity of these *trans*-specific enzymes toward Ser(P)-Pro motifs that adopt the *cis* conformation in the structure of their protein. For example, ERK2 phosphorylates two sites in RNase T1, *cis*-Ser⁵⁴-Pro⁵⁵ and *trans*-Ser⁷²-Pro⁷³; however, only phosphorylation at Ser⁵⁴-Pro⁵⁵ is enhanced by the addition of peptidyl prolyl isomerases. This illustrates a crucial point: catalyzed proline isomerization is an effective regulatory mechanism of phosphorylation/dephosphorylation only when the target motif exists predominantly in the oppo-

site isomeric configuration required for catalysis. In this case, the isomer-specific enzyme can only process a small fraction of the total substrate, and the extremely slow, uncatalyzed *cis-trans* isomerization prevents re-equilibration of the non-processed fraction. In the opposite scenario, the majority of substrate is already in the correct isomeric form and can be readily processed, with catalyzed proline isomerization having little influence.

The effect of proline isomerization on CTD-associated activities depends greatly on the CTD conformation during the transcription cycle. All structural studies of CTD-binding proteins have revealed binding that is, in fact, isomer-specific but preferential for *trans*-Ser(P)-Pro motifs. In the case of Pcf11, the CTD-interacting domain discriminates in favor of 3 consecutive *trans*-prolines, including one Ser(P)²-Pro³ motif (8). In the structure of CTD phosphatase Scp1 in complex with Ser(P)²/Ser(P)⁵ CTD, both the Ser(P)²-Pro³ and the Ser(P)⁵-Pro⁶ motifs adopt the *trans* configuration, with binding to either *cis* form seemingly prevented by steric clashes (34). For both of these proteins, the regulation of binding or catalysis by proline isomerization would demand a CTD conformation with a minority *trans* population. This is especially true for non-enzymatic proteins such as Pcf11, which may only require one correctly configured binding site out of dozens of CTD repeats. However, our measurements with a Ser(P)⁵ CTD peptide and prior work with a Ser(P)² CTD peptide (8) show that *cis*-proline is the less populated isomer. Given the lack of evidence for any stable full-length CTD structure, it seems likely that *cis*-prolines constitute the isomeric minority *in vivo* as well and that proteins with specific recognition of *cis*-proline are the major targets of regulation by isomerization.

The timescale of transcriptional events is another critical determinant of the effect of Ess1 on CTD-associated activities. We observed only an \sim 2-fold enhancement of Ssu72-catalyzed product accumulation in reactions with synthetic pCTD substrate as a significant amount of non-catalyzed proline isomerization occurs over the course of the 5-min reactions. A similar level of stimulation was reported for the PP2A phosphatase described above in reactions on the same timescale (40). However, at a faster timescale, the effect of Ess1 on Ssu72 activity is greater, as shown in [supplemental Fig. S3](#) in 1.25-min reactions monitored at 15-s increments. Although the exact *in vivo* timescale of Ssu72 activity is unknown, the effect of impaired Ess1 activity on Nrd1 localization and Pcf11 recruitment is largely confined to the 3'-end of genes (28), suggesting that Ser(P)⁵ CTD dephosphorylation by Ssu72 occurs mainly during termination, which may take only tens of seconds (42). This scenario is well illustrated by the kinetic assays with full-length pCTD in Fig. 4, *E* and *F*, where the reaction without Ess1 takes 20 min to dephosphorylate \sim 90% of the initial pCTD substrate. In contrast, the reaction with Ess1 reaches \sim 90% completion in 1 min, on par with the estimated timescale of transcription termination.

A Revised Model for Ess1-regulated Ssu72 Activity—Our kinetic experiments provide an excellent interpretation of recent *in vivo* studies of Ess1 (28, 29). We show that the stimulation of Ssu72 activity by Ess1-catalyzed *cis-trans* intercon-

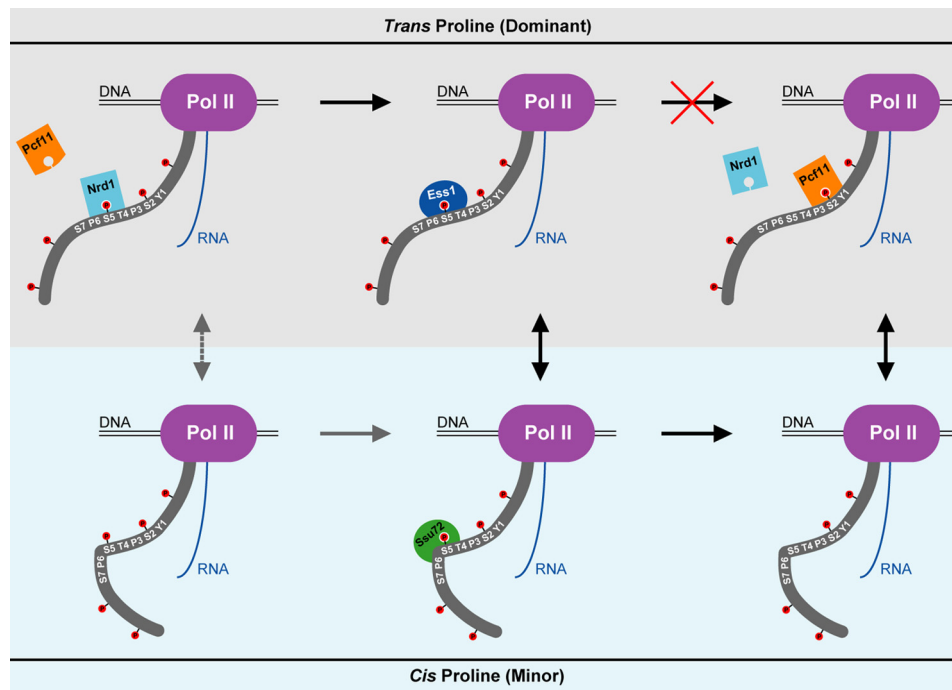


FIGURE 5. **A new model for Ess1-regulated Ssu72 activity.** RNAPII (*Pol II*) is shown near the termination of an snRNA transcript. For clarity, only one CTD heptad is labeled. The upper (gray) and lower (light blue) panels represent the *trans* and *cis* forms, respectively, of the Ser(P)⁵–Pro⁶ motif. Dephosphorylation cannot proceed through the *trans* form (red cross) due to the *cis*-proline substrate specificity of Ssu72. Instead, Ess1 catalyzes the interconversion of isomers, which is the rate-limiting step for Ssu72 activity. The minor pathway is indicated with gray arrows, with the dashed line representing non-catalyzed isomerization that occurs very slowly compared with transcriptional events. The change in CTD phosphorylation facilitates proper termination by promoting the release of Nrd1 and recruiting Pcf11.

version of Ser(P)⁵–Pro⁶ motifs in the CTD heptad is severely reduced with catalytically impaired Ess1 mutants, which explains the *in vivo* accumulation of Ser(P)⁵ CTD in *ess1* cells. Interestingly, overexpression of Ssu72 suppressed this phenotype in an A144T *ess1* strain, but not an H164R *ess1* strain (29). It is possible that different levels of residual activity or thermostability of the Ess1 mutants are responsible for this discrepancy. In contrast, overexpression of Ssu72 rescues the temperature sensitivity of both *ess1* strains (29), suggesting that improper regulation of CTD phosphorylation is the main cause of this phenotype. As described in the Introduction, the proposed proline isomer-based regulatory scheme that explains these results requires *trans* specific activity by Ssu72 (28, 29). Our results compel a simple but unexpected adjustment of this model: reversing the isomeric specificity of Ssu72 (Fig. 5). Interestingly, genetic evidence suggests that this model may be partially bypassed by overexpression of Ser(P)⁵ CTD phosphatase Fcp1 in yeast (16).

Implications of Proline Isomerization in CTD Biology—Ssu72 provides an exciting validation of proline isomers as a critical component of the CTD code but also raises many new questions about how this regulatory mechanism operates. For example, it is unclear whether the regulation of Ssu72 is a general property of RNAPIII transcription or a gene-specific one. Although impaired Ess1 and Ssu72 activity causes readthrough transcription of a small subset of genes consisting mainly of snRNA that undergo Nrd1-dependent termination (16, 28), it is possible this phenotype is only apparent for short transcripts that require a sudden drop in Ser(P)⁵ levels for proper termination. It will also be interesting to see what

other CTD-binding partners specifically recognize *cis*-proline and how their functions are regulated by catalyzed isomerization. Although these proteins may only regulate the CTD phosphorylation state, there may also be non-enzymatic CTD-binding factors with *cis*-proline recognition whose association with the transcribing polymerase is regulated by Ess1/Pin1 to fine-tune the recruitment of their co-transcriptional activities.

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