Research Article

Postrecruitment Function of Yeast Med6 Protein during the Transcriptional Activation by Mediator Complex

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Med6 protein (Med6p) is a hallmark component of evolutionarily conserved Mediator complexes, and the genuine role of Med6p in Mediator functions remains elusive. For the functional analysis of *Saccharomyces cerevisiae* Med6p (scMed6p), we generated a series of scMed6p mutants harboring a small internal deletion. Genetic analysis of these mutants revealed that three regions (amino acids 33–42 (Δ2), 125–134 (Δ5), and 157–166 (Δ6)) of scMed6p are required for cell viability and are located at highly conserved regions of Med6 homologs. Notably, the Med6p-Δ2 mutant was barely detectable in whole-cell extracts and purified Mediator, suggesting a loss of Mediator association and concurrent rapid degradation. Consistent with this, the recombinant forms of Med6p having these mutations partially (Δ2) restore or fail (Δ5 and Δ6) to restore in vitro transcriptional defects caused by temperature-sensitive med6 mutation. In an artificial recruitment assay, Mediator containing a LexA-fused wild-type Med6p or Med6p-Δ5 was recruited to the lexA operator region with TBP and activated reporter gene expression. However, the recruitment of Mediator containing LexA-Med6p-Δ6 to lexA operator region resulted in neither TBP recruitment nor reporter gene expression. This result demonstrates a pivotal role of Med6p in the postrecruitment function of Mediator, which is essential for transcriptional activation by Mediator.

1. Introduction

Promoter-specific mRNA synthesis requires a minimal set of proteins comprising Pol II (RNA polymerase II) and associated GTFs (general transcription factors), which is defined as in vitro “basal transcription” [1]. Regulated responses of this minimal transcription system by DNA-binding transcription factors or in vitro transcription from nucleosomal DNA template require additional sets of coregulator proteins involved in chromatin remodeling/modification as well as targeted recruitment of basal transcription machinery to the promoter [2, 3]. In vivo, these transcriptional cofactors indeed participate in the regulated expression of specific genes in response to environmental and physiological cues, which has been demonstrated in a wide range of eukaryotic model organisms.

Mediator complex was first identified from yeast *S. cerevisiae* [4], and Mediator homologs or its related complexes were subsequently found in other eukaryotic species, placing this evolutionarily conserved coregulator as an essential and general player in Pol II transcription [5, 6]. Yeast Mediator is required for diverse aspects of transcriptional regulation, such as activation, repression, stimulation of basal transcription, and TFIIH-dependent phosphorylation of the CTD (C-terminal repeat domain) of the Rpb1 subunit in vitro [2, 4]. The tight association of Mediator with the CTD of Pol II forms Pol II holoenzyme, which is generally required for the transcription of most class II genes in vivo [7]. Genetic and biochemical studies identified 21 polypeptides as bona fide subunits of the core Mediator, including transcriptional regulators previously identified by genetic studies, as well as the novel subunits collectively named Med proteins [8]. In addition, a subset of Srb/Ssn proteins was identified as the distinct kinase module whose association with core Mediator is known to suppress the positive function of Mediator [9, 10].

Molecular and genetic studies have revealed that some Mediator subunits are specifically required for the regulation
of a subset of genes, whereas others are necessary for general transcription of Pol II [11]. Consistent with these observations, our biochemical studies have demonstrated that functionally related Mediator subunits physically associate to form two stable subcomplexes, named Rgr1/Med14 and Srb4/Med17 [12]. The Med14 subcomplex was subsequently shown to contain several functional modules, including Gal11/Med15 module (Med2, -3, -15, and -16) and Med9/10 module (Med1, -4, -7, -9, -10, and -21), which suggested a modular structure of Mediator [12, 13]. While the Med14 subcomplex consists of several functional modules presumed to function in receiving diverse signals from activators or repressors, the Med17 subcomplex, composed of Med6p and genetically dominant Srb proteins (Med17, -18, -20, and -22), is thought to functionally interact with the Rpb1 CTD [12]. Based on electron microscopic (EM) images, the modular structure of core Mediator was represented as comprising distinct head, middle, and tail domains [14]. Recent X-ray structures of yeast Mediators suggest that the Med17 subcomplex corresponds to the head module while the Med14 subcomplex contains both tail and middle modules [15–18]. In holoenzyme and preinitiation complex (PIC) models based on refined EM and X-ray structures, both the middle and head modules make multiple contacts with diverse parts of Pol II and GTFs, explaining the molecular mechanism of Mediator-stimulated CTD phosphorylation by TFIIH and resultant enhancement of basal transcription [15, 16, 19].

Med6p is a hallmark component of evolutionarily conserved Mediator complexes and participates in transcriptional activation of many class II genes in S. cerevisiae [5, 20]. In metazoans, mutant flies deficient in dMED6 exhibit transcriptional defects in a wide range of developmentally regulated genes, validating the functional conservation and the requirement for Med6p in Pol II transcription in vivo [21]. Intriguingly, med17 mutation was isolated as a dominant suppressor of med6 temperature-sensitive (ts) mutation [12], and this genetic interaction was confirmed by an opposite genetic approach [22]. Consistent with functional interaction, a direct physical association between Med6p and Med17p was identified through successful in vitro reconstitution of the Med17 subcomplex [13, 23]. Recently, X-ray structural data have been obtained for the head modules of S. cerevisiae and S. pombe Mediators, which indicate that the structures of two head modules are well conserved despite the limited sequence similarities between these regions [17, 18]. The conserved Med6p structure revealed a core domain, which comprises a five-stranded antiparallel β-sheet, two pairs of α-helices flanking a conserved groove, and a flexible C-terminal α5 helix [18]. This domain is called “shoulder,” according to its position relative to the head module, and was shown to interact with other subunits within the head module (Med8 and Med17) and the middle module (Med4, Med10, and Med19) [18]. Recently, the C-terminal α6 helix of the Med6 subunit was shown to connect to the Med14 subunit of the middle module as one of the tethering molecules of the head module [15, 16]. These results strongly suggest that the Med6p may function as a critical interface between the head and middle modules and relay a regulatory signal from the tail/middle module to the head module/Pol II as proposed [12].

Although extensive structural studies have been made to reveal the action mechanisms of Mediator in Pol II transcription, analyses of each Mediator component to confirm the functions inferred from structural data remain to be completed. In order to enhance our understanding of the key role of S. cerevisiae Med6p (scMed6p) in Mediator functions, we generated a series of Med6p mutants harboring small internal deletions (10 amino acids) and analyzed the importance of each region of scMed6p in transcription activation by Mediator in vivo and in vitro. We identified three distinct domains of scMed6p required for its essential function: one N-terminal region (Δ2, amino acids 33–42) might be involved in Mediator association and/or protein stability, and the second (Δ5, amino acids 125–134) and third (Δ6, amino acids 157–166) regions have a role in post-recruitment function, such as signal-relay or conformational changes during transcriptional activation by Mediator.

2. Materials and Methods

2.1. Construction of med6 Deletions, Mutants, and Other Expression Plasmids. Serial deletion mutants of the scMED6 gene were constructed by an oligonucleotide-directed mutagenesis method. The MED6 gene harboring own promoter and terminator regions was obtained from pRS316-MED6 by EcoRI-XbaI digestion and subcloned into the pALTER-1 vector (Promega) for site-directed mutagenesis according to the manufacturer’s instructions. A series of mutagenic oligonucleotides were designed to delete 10 amino acids of Med6p spanning the whole MED6 ORF region (Figure 1). A Nar1 restriction site was introduced into deletion points within each mutagenic oligonucleotide, which was used for the construction of double deletion mutants (Δ3-4, Δ7-8, and Δ9-10). After the deletions were confirmed by sequencing, mutant med6 genes were subcloned into pRS313 plasmid.

For the copper-inducible expression of Med6p, wild-type and mutant MED6 genes were subcloned into pC (native form) or pCL (LexA-fused form) vectors using appropriate restriction sites [26]. pGEX-MED6 was made by subcloning the EcoRI-Sall fragment from pGBT-MED6 into pGEX-2TK (Pharmacia). The pGEX-med6 deletion constructs were made by replacing the indicated region of pGEX-MED6 with the corresponding fragments of mutant constructs in pRS313 by the use of restriction sites. In order to make the E. coli expression constructs for 6xHis-tagged Med6p and its deletion derivatives, the EcoRI-ClaI fragments from the pGEX-MED6 and deletion constructs were subcloned into pET vector series.

2.2. Growth Complementation Assay. To test the growth complementation by med6 deletion mutants, yeast strain YCL4 (MATa, ade2, his3, trp1, lys2, med6Δ::LEU2, and pRS316-MED6) was used as a host strain for plasmid shuffling. The med6 deletion mutants in pRS313 were introduced into YCL4 strain and the resulting transformants
were tested for their growth on synthetic complete media containing 5-fluoroorotic acid via the plasmid shuffling method.

2.3. Protein Purification. Gal4-VP16, GTFs, wild-type, and med6\textsuperscript{ts} mutant Pol II holoenzymes were prepared using several chromatographic steps as described previously [20].
Recombinant Med6p and mutant derivatives were expressed in E. coli strain BL21(DE) (Novagen) and purified through Ni²⁺-NTA columns (Qiagen) under denaturing conditions. Purified proteins were renatured at 4°C through stepwise dialysis according to the protocol described by Thompson et al. [27] and stored at −80°C before use.

2.4. In Vitro Transcription. Reconstituted in vitro transcription was performed as described previously [20]. Either wild-type or med6 ts mutant Pol II holoenzymes were preincubated for 10 min at 25°C or 37°C with other supplements containing two DNA templates (pGAL4::G- and pGCN4::G-), GTFs, 0.5 mM ATP, and Gal4-VP16, with or without recombinant Med6p derivatives. After initiation complex formation, the transcription reaction (30 min, 25°C) was initiated by the addition of [α-³²P]-UTP and CTP. The purified reaction products were analyzed on the 7% denaturing PAGE and autoradiography.

2.5. LacZ Reporter Assay, GST Pull-Down Analysis, Western Analysis, and Chromatin Immunoprecipitation. For the analysis of reporter gene expression in yeast, yeast strains were grown to the mid-log phase in selective synthetic complete media and β-galactosidase activity of cultured cells was measured in triplicate by the permeabilized-cell method [28]. To investigate the expression levels of mutant Med6ps, yeast strain YCL4 was transformed with plasmids encoding LexA-Med6p derivatives (in pCL313) and/or native forms of Med6p derivatives (in pC314). The resulting transformants were cultured in synthetic glucose media containing 0.5 mM CuSO₄ for 2 h to induce the expression of Med6 proteins. Preparation of yeast whole-cell extracts (WCEs), western analysis, and immunoprecipitation of Mediator were performed according to the protocols described previously [12]. For chromatin immunoprecipitation, L40 yeast strains expressing LexA-Med6p derivatives (in pCL313) were grown in synthetic media containing glucose (Glc) or galactose plus raffinose (Gal + Raf) for 6 h in the presence of 0.5 mM CuSO₄. Cells were harvested, fixed in 1% formaldehyde solution, and sonicated for the fragmentation of chromatin according to the protocol described by Kuo and Allis (1999). WCEs containing chromatin fragments were immunoprecipitated with the use of appropriate antibodies and subjected to a decrosslinking reaction. Purified DNAs were subjected to PCR (25 cycles) with the use of oligomers specific to promoter regions of LacZ reporter or GAL1-10 genes, respectively.

3. Results

3.1. Identification of yMed6p Regions Required for Cell Viability. For the unbiased and systematic analysis of yMed6p, a series of Med6p mutants harboring a small internal deletion was made. We designed 10 amino acid deletions throughout the entire Med6p for approximately every 30 amino acid interval and introduced these into MED6 genes harboring own promoter and terminator regions (Figures 1 and 2). The mutant med6 constructs were then transformed into host strain YCL4 for plasmid shuffling and tested for supporting cell viability upon the removal of wild-type MED6 gene in pRS316 by 5-fluoroorotic acid treatment (Figure 3(a)). Despite the considerable size for deletions (10 amino acids), seven out of ten deletion mutations (Δ1, Δ3, Δ4, Δ7, Δ8, Δ9, and Δ10) had no obvious effect on the viability of yeast cells (Figures 2 and 3(b)). Notably, the med6 deletion mutants containing Δ2 (residues 33–42), Δ5 (residues 125–134), or Δ6 (residues 157–166) were unable to support the cell growth.

The multiple-sequence alignment of Med6p homologs using the Expresso program indicated a high degree of conservation between yeast and metazoan Med6 proteins (Figure 1). However, scMed6p has distinct features from other Med6 homologs in four regions: two internal and one C-terminal spacer regions (50, 8, and 30 amino acids long, resp.) that are exclusively found in scMed6p and one variable region among Med6 homologs (amino acids 230–240 in scMed6p). According to alignment data, three regions (Δ2, Δ5, and Δ6), identified as essential for MED6 function(s), closely matched the highly conserved regions of Med6 homologs (Figure 1).

To confirm the functional regions of scMed6p, larger deletion mutants were made and used in a complementation test. First, we constructed a Med6p mutant having the combined deletion of Δ3 and Δ4 (Δ3-4), resulting in the removal of the first region specific for scMed6p. Interestingly, this mutant could fully complement the MED6 function, indicating that this region is dispensable for essential function (Figures 2 and 3(b)). In addition, one-third of the C-terminal domain of scMed6p, corresponding to the most divergent region of Med6p, also proved to be unnecessary for MED6 function, since two mutants (Δ7-8 and Δ9-10) spanning this region had no obvious effect on cell survival (Figures 2 and 3(b)). This assumption was confirmed by the fact that scMed6p comprising only residues 1–210 could support cell viability (see Supplementary Figure 1).

Taken together, the functional domains of yMed6p essential for cell viability were mapped to the highly conserved regions of Med6p, whereas the scMed6p-specific region and the divergent region of Med6ps were not required for essential function(s) of scMed6p.

3.2. Phenotypes or Transcriptional Defects of the Viable Med6 Mutants. Although the deletion analysis indicated that substantial parts of scMed6p were dispensable for cell viability, a significant level of sequence conservation is apparent at these regions, including Δ1 and Δ7 regions. Thus, we investigated whether the viable med6 deletion mutants showed any growth defects or transcriptional defects in vivo, such as ts lethality or limited carbon source utilization, as observed in med6 ts mutants [12]. All of the viable mutant strains had no observable ts phenotype and utilized galactose or raffinose at 30°C as efficiently as the wild-type did (Supplementary Figure 2A).

Based on the fact that MED6 is essential for galactose-induced GAL1 gene activation [20], β-galactosidase activity of the GAL1₆₆₆₆-TATA-LacZ reporter was measured for the
viable mutants under induction conditions. None of the viable med6 deletion mutants were defective in transcriptional activation of the GAL1 promoter (Supplementary Figure 2B). These results strongly suggest that the pivotal function(s) of scMed6p resides in the evolutionarily conserved regions.

3.3. In Vitro Transcriptional Activity of Mutant Med6 Proteins. Next, we investigated whether scMed6p regions essential for viability are also required for Med6p-dependent transcriptional activation in vitro. For this, we purified recombinant Med6p having ts-2 mutation or internal deletion mutations and examined their capabilities in restoring the transcriptional defects of Pol II holoenzyme caused by med6 ts-2 mutation [20]. When transcriptional PIC was formed at 25°C, both wild-type and med6-ts2 Pol II holoenzymes showed 23- and 21-fold activation of GAL4 enhancer-containing template (GAL4:G-) by Gal4-VP16 over basal transcription from GCN4:G-template, respectively (Figure 4, lanes 1 and 2). However, PIC formation at 37°C specifically impaired the transcription activity of med6-ts2 holoenzyme in that the mutant holoenzyme gave only a 2.5-fold activation in comparison to the 21-fold activation by wild-type holoenzyme (Figure 4, lanes 3 and 4). The temperature-dependent transcriptional defect of mutant holoenzyme was recovered by the addition of the recombinant form of wild-type Med6p, but not Med6-ts2 protein, prior to the PIC formation at 37°C (lanes 5–7). Next, we examined whether each deletion mutant of Med6p (-Δ2, -Δ5, -Δ6, and -Δ7) had the ability to restore the transcriptional defect of med6-ts2 holoenzyme. The Med6p-Δ7, which showed no apparent defect in vivo, did rescue the activation defect of the heat-inactivated med6-ts2 holoenzyme (lane II). However, Med6p-Δ5 and -Δ6 mutants were not able to complement this defect (lanes 9 and 10), in good agreement with their in vivo phenotypes. Notably, Med6p-Δ2, which did not support cell viability, could partially restore the transcriptional defect of med6-ts2 holoenzyme to half that of the wild-type (lanes 5, 6, and 8; see Discussion).

**Figure 2**: Schematic presentations of Med6p derivatives and summary of their functional defects in vivo and in vitro. Viability1 was tested for the ability of deletion mutants to complement the med6 null mutation via plasmid shuffling as described in Materials and Methods. In vitro activity2 represents the capability of each recombinant form of Med6 proteins to rescue the in vitro transcriptional defects of med6-ts2 holoenzyme based on the quantitative data shown in Figure 4. β-galactosidase activity3 indicates the expression levels of the reporter gene in the artificial recruitment assay of the indicated Med6p derivatives as shown in Figure 6(a).

**Figure 3**: Identification of the regions in Med6p required for cell viability. (a) Schematic depiction of plasmid shuffle technique for the complementation test of mutant MED6 genes (med6Δ). (b) The growth complementation test of the med6 deletion mutants. Wild-type (WT) MED6 and indicated med6 deletion mutants in the pRS313 vector were introduced into the host strain YCL4 (med6Δ::LEU2 and pRS316-MED6) for plasmid shuffling. The resulting transformants were grown for 4 days at 30°C on synthetic complete media containing 5-fluoorotic acid to remove the pRS316-MED6. (−): pRS313 vector only.
These results indicate that the Med6p regions which are essential for in vivo function (cell viability) are also required for activation function of Med6p in vitro.

3.4. Med6p-Δ2 Has a Defect in Association with Mediator. In our previous reports, the Pol II holoenzyme prepared from med6-ts mutant (Ts) Pol II holoenzyme (700 ng each) was preincubated for 10 min at 25°C or 37°C with other supplements containing two DNA templates, GTFs, 0.5 mM ATP, and Gal4-VP16 (30 ng), with or without the indicated recombinant Med6p (80 ng each). After PIC formation, the transcription reaction (30 min, 25°C) was initiated by the addition of [α-32P]-UTP and CTP. The mRNA products were purified and resolved on 7% denaturing polyacrylamide gel, followed by autoradiogram. Transcription (%) was set to 100% as to the fold activation (signal from GAL4 template divided by signal from GCN4 template) shown by wild-type holoenzyme upon PIC formation at 37°C. The data quantitation was performed with the use of Personal Molecular Imager FX system and associated program (Bio-Rad).

3.5. Artificial Recruitment Assay with the Functionally Defective Med6p Mutants. The artificial recruitment of Mediator to a promoter region via Mediator subunit fused with DNA-binding domain resulted in activator-bypass activation of target promoter. We successfully performed an artificial recruitment assay by using Mediator containing LexA-Med6p [29]. Thus, we used LexA-fused Med6p mutants for the artificial recruitment assay to activate the chromosomal 8xLexA-op-LacZ reporter gene. As expected from the expression data, LexA-Med6p-Δ2 was completely inactive for reporter gene activation (Figure 6(a)).

![Figure 4](image4.png)

**Figure 4**: Three lethal Med6p mutants also show transcriptional defects in a reconstituted in vitro system. Either wild-type (WT) or med6 ts-2 mutant (Ts) Pol II holoenzyme (700 ng each) was preincubated for 10 min at 25°C or 37°C with other supplements containing two DNA templates, GTFs, 0.5 mM ATP, and Gal4-VP16 (30 ng), with or without the indicated recombinant Med6p (80 ng each). After PIC formation, the transcription reaction (30 min, 25°C) was initiated by the addition of [α-32P]-UTP and CTP. The mRNA products were purified and resolved on 7% denaturing polyacrylamide gel, followed by autoradiogram. Transcription (%) was set to 100% as to the fold activation (signal from GAL4 template divided by signal from GCN4 template) shown by wild-type holoenzyme upon PIC formation at 37°C. The data quantitation was performed with the use of Personal Molecular Imager FX system and associated program (Bio-Rad).

![Figure 5](image5.png)

**Figure 5**: The deficiency of Med6p-Δ2 in WCEs and purified Mediator fraction. (lanes 1–12) WCEs were prepared from YCPL4 yeast strains expressing the indicated LexA-Med6p derivatives (pCL-MED6) along with (+) or without (−) coexpression of the native forms of Med6p derivatives (pC-MED6) by copper induction for 2 h. Samples were resolved on an 8% denaturing polyacrylamide gel, and expression levels of LexA-fused and native forms of Med6p derivatives were examined by western analysis using anti-LexA antibody and anti-Med6 antibody, respectively. Western analysis for TBP was done for loading control. The degradation product of LexA-Med6p-Δ2 is indicated by an asterisk. (lanes 13–16) WCEs were prepared from the indicated yeast strains and subjected to DEAE column fractionation to enrich the Pol II holoenzyme. Pol II holoenzyme in each sample was immunopurified with beads-coupled anti-Med14 antibody and subjected to western analysis to measure the amounts of LexA-Med6p derivatives (α-LexA) contained in the immunopurified Mediator fraction (α-Med14).
Intriguingly, LexA-Med6p-Δ5 had comparable transcriptional activity to that of wild-type Med6p, suggesting its transcriptional defect can be reversed by artificial recruitment via an unknown mechanism (see Discussion). In the case of the Med6p-Δ6 mutant, its artificial recruitment had no effect on reporter gene expression (Figure 6(a)).

Next, we performed chromatin immunoprecipitation analysis to examine in vivo association of LexA-Med6p derivatives, Med14 (for Mediator), and TBP (α-TBP) to promoter regions of chromosomal reporter (8xlexAop) and GAL1-10 (GAL1-10p) genes. L40 strains expressing indicated pCL313-Med6p derivatives were grown in synthetic media containing glucose (Glc) or galactose plus raffinose (Gal+Raf) for 6 h in the presence of copper. Chromatin fragments were prepared from WCEs and immunoprecipitated with the use of the indicated antibodies. DNAs were purified from immunoprecipitates and subjected to PCR (25 cycles) with the use of oligomers specific to promoter regions of genes. L40 harboring pCL313-Med6p derivatives were cultured in synthetic glucose media containing 0.5 mM copper. The β-galactosidase activity of chromosomal 8xlexAop-LacZ reporter gene in cultured cells was measured in triplicate.

Figure 6: Artificial recruitment assay and chromatin immunoprecipitation analysis of Med6p derivatives. Yeast strain L40 harboring pCL313-Med6p derivatives were cultured in synthetic glucose media containing 0.5 mM copper. The β-galactosidase activity of chromosomal 8xlexAop-LacZ reporter gene in cultured cells was measured in triplicate. (b) Chromatin immunoprecipitation for recruitment of Mediator (α-Med14), LexA-Med6p (α-LexA), and TBP (α-TBP) to promoter regions of chromosomal reporter (8xlexAop) and GAL1-10 (GAL1-10p) genes. L40 strains expressing indicated pCL313-Med6p derivatives were grown in synthetic media containing glucose (Glc) or galactose plus raffinose (Gal + Raf) for 6 h in the presence of copper. Chromatin fragments were prepared from WCEs and immunoprecipitated with the use of the indicated antibodies.

4. Discussion

In this report, we dissected the functional domains of yeast Med6p on the basis of its requirement for cell viability and transcriptional activation via molecular genetics and biochemical approaches. The functional domains of scMed6p (Δ2, Δ5, and Δ6 regions) identified by complementation assays were mapped to the highly conserved regions of Med6ps, whereas yeast-specific (spacer region) and mostly divergent regions (one third of C-terminus) of scMed6p were not required for its essential function(s) (Figures 1 and 2). Three lethal deletion mutants had defects in transcriptional activation in a reconstituted in vitro system although only the Gal4-VP16 activator was tested in our in vitro transcription (Figure 4). These results again clearly demonstrate that the major and essential function of Med6p is focused on transcriptional activation.

One interesting point is that some residual activity to support in vitro transcription was retained in Med6p-Δ2, showing a discrepancy with its in vivo defect. Although we cannot explain clearly this phenomenon, we surmise that this partial activity might have resulted from an in vitro dosage effect. If Med6p-Δ2 has a normal activation function but has a defect in Mediator association, Med6p-Δ2 could provide in vitro activation function to some extent via
a dosage effect when an excess amount of Med6p\(\Delta 2\) is added (which is not applicable in vivo).

In contrast to Med6p\(\Delta 5\) and \(-\Delta 6\) mutants, the Med6p\(\Delta 2\) protein was barely detectable in WCEs and was even absent in enriched Pol II-Mediator fraction. As mentioned above, the comparable proportions of \(med6-ts\) proteins detected in WCEs seem to be associated with Mediator in vivo. The observed deficiency and the degradation pattern of Med6p\(\Delta 2\) in WCEs suggest that the \(\Delta 2\) region of scMed6p might be involved in the association with other Mediator subunit(s) (Figure 5). Based on recent X-ray structure of \(S.\ pombe\) Mediator in PIC, the C-terminal residues of \(a3\) helix of spMed6p are shown to directly interact with the Med10/Nut2 subunit to form a head-middle module interface [16]. Since the \(\Delta 2\) region of scMed6p contains N-terminal residues of \(a3\) helix, its deletion might disrupt Med6-Med10 interaction by affecting \(a3\) helix structure (Figure 1). In this scenario, the dissociated form of the \(\Delta 2\) mutant seems to be rapidly removed via specific degradation pathway probably due to its instability. Notably, one (Q49L) of three critical mutations in the \(med6-ts2\) allele is found in the \(a3\) helix of scMed6p, suggesting the molecular basis of the deficiency of \(med6-ts2\) protein in WCEs and purified Mediator [20]. The other two mutations (I68L and L94P) were mapped to the \(\Delta 3\) and \(\Delta 4\) regions, respectively, which correspond to the nonessential spacer region of scMed6p (Figure 1).

Previous biochemical analyses and artificial recruitment experiments using mutant Pol II holoenzymes suggested that facilitated recruitment of a holoenzyme to a promoter is necessary but not sufficient for transcriptional activation and that an unknown biochemical activity is required for transcriptional activation (postrecruitment step) [29]. Consistent with this, the structural data suggested that Med6p may function as a critical interface between the head and middle modules, and relay a regulatory signal from the tail/middle module to the head module with this, the structural data suggested that Med6p may

required for its essential function: one N-terminal region (\(\Delta 5\), amino acids 33–42) and second (\(\Delta 6\), amino acids 157–166) might be involved in Mediator association and/or protein stability, and the second (\(\Delta 5\), amino acids 125–134) and third (\(\Delta 6\), amino acids 157–166) regions have a role in postrecruitment activity of Mediator complex, such as signal-relay or conformational changes. This study proved the previous notion that Med6p is involved in the postrecruitment function of Mediator complex, which is required for gene activation by Mediator.

### 5. Conclusion

Through the systematic dissection of functional domains of scMed6p, we identified three distinct domains of scMed6p required for its essential function: one N-terminal region (\(\Delta 2\), amino acids 33–42) might be involved in Mediator association and/or protein stability, and the second (\(\Delta 5\), amino acids 125–134) and third (\(\Delta 6\), amino acids 157–166) regions have a role in postrecruitment activity of Mediator complex, such as signal-relay or conformational changes. This study proved the previous notion that Med6p is involved in the postrecruitment function of Mediator complex, which is required for gene activation by Mediator.

### Abbreviations

- CTD: C-terminal repeat domain
- GTFs: General transcription factors
- Pol II: RNA polymerase II
- ts: Temperature-sensitive
- PIC: Preinitiation complex
- TBP: TATA-binding protein
- WCEs: Whole-cell extracts

### Conflicts of Interest

The authors declare that there are no conflicts of interest regarding the publication of this article.

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Supplementary Materials

Supplementary Figure 1: The growth complementation test of Med6p comprising residues 1–210. Supplementary Figure 2: The viable Med6 mutants showed no detectable phenotypes or transcriptional defects. (Supplementary Materials)

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