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**Iwr1 Protein Is Important for Preinitiation Complex Formation by All Three Nuclear RNA Polymerases in Saccharomyces cerevisiae**

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**Abstract**

**Background:** Iwr1, a protein conserved throughout eukaryotes, was originally identified by its physical interaction with RNA polymerase (Pol) II.

**Principal Findings:** Here, we identify Iwr1 in a genetic screen designed to uncover proteins involved in Pol III transcription in *S. cerevisiae*. Iwr1 is important for Pol III transcription, because an *iwr1* mutant strain shows reduced association of TBP and Pol III at Pol III promoters, a decreased rate of Pol III transcription, and lower steady-state levels of Pol III transcripts. Interestingly, an *iwr1* mutant strain also displays reduced association of TBP to Pol I-transcribed genes and of both TBP and Pol II to Pol II-transcribed promoters. Despite this, rRNA and mRNA levels are virtually unaffected, suggesting a post-transcriptional mechanism compensating for the occupancy defect.

**Conclusions:** Thus, Iwr1 plays an important role in preinitiation complex formation by all three nuclear RNA polymerases.

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Eukaryotic cells contain three multi-subunit RNA polymerases (Pol) that transcribe the nuclear genome and are responsible for the production of selected classes of RNAs [1–5]. Pol I is responsible for synthesis of the tandem repeated ribosomal RNA genes, Pol II synthesizes mRNA and many non-coding RNAs, and Pol III synthesizes tRNA, 5S rRNA, and few other small untranslated RNAs. These RNA polymerases share 5 subunits, and their catalytic cores are similar to each other and to *E.coli* RNA polymerase [6]. Unlike bacterial and bacteriophage RNA polymerases that bind specifically to promoter sequences, the eukaryotic enzymes work in conjunction with transcription factors that directly bind promoters and recruit the appropriate RNA polymerase to initiate transcription [7]. The TATA-binding protein (TBP) is required for transcription by all three RNA polymerases [8], and it is a component of multi-protein complexes that function specifically with a particular RNA polymerase machinery [9].

Described by the basal transcription machinery that contains TBP, Pol II, and general transcription factors [10]. Upon initiation, Pol II dissociates from these general factors and associates with “elongation factors” that travel down the mRNA coding region [11]. In vivo, efficient transcription requires activator proteins that bind specifically to regulatory DNA sequences and, via co-activators, stimulate the basal transcription machinery [7]. Some Pol II-transcribed genes are regulated by repressors that bind to specific DNA sequences. The identity, quality, and location of regulatory sequences are gene-specific, with the consequence that every gene has a distinct pattern of expression.

For the vast majority of Pol III-transcribed genes, promoter recognition elements are located internally within the RNA coding region, and Pol III transcription involves a multi-step assembly of general initiation factors [1,4,12]. In general, the six-subunit TFIIC binds to the A- and B-boxes, and it acts as an assembly factor directing binding of the TBP complex, TFIIB, to a position just upstream of the initiation site. Transcription of 5S rRNA genes requires an additional factor, TFIH, that binds to the A-box, G-box, and IE-element. Once TFIIB is assembled, the RNA polymerase is recruited and directs multiple rounds of transcription. Although Pol III genes are not individually regulated in the manner of Pol II genes, they are collectively subject to the negative

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**Introduction**

Eukaryotic cells contain three multi-subunit RNA polymerases (Pol) that transcribe the nuclear genome and are responsible for the production of selected classes of RNAs [1–5]. Pol I is responsible for synthesis of the tandem repeated ribosomal RNA genes, Pol II synthesizes mRNA and many non-coding RNAs, and Pol III synthesizes tRNA, 5S rRNA, and few other small untranslated RNAs. These RNA polymerases share 5 subunits, and their catalytic cores are similar to each other and to *E.coli* RNA polymerase [6]. Unlike bacterial and bacteriophage RNA polymerases that bind specifically to promoter sequences, the eukaryotic enzymes work in conjunction with transcription factors that directly bind promoters and recruit the appropriate RNA polymerase to initiate transcription [7]. The TATA-binding protein (TBP) is required for transcription by all three RNA polymerases [8], and it is a component of multi-protein complexes that function specifically with a particular RNA polymerase machinery [9].
regulator Ma1, which inhibits transcription in response to stress signals such as oxidative stress, cell wall stress, DNA damage, or nutrient limitation [13].

After transcription, specific nucleosides in primary rRNA transcripts become modified to yield a mature functional rRNA [14]. In S. cernua, the initiator methionine tRNA (tRNA\(^{\text{Met}}\)) contains a unique 2'-O-riboyl phosphate modification (Ar(p)) at position 64 [15] that is important for the discrimination between translational initiation and elongation [16–18]. Rtx1, the initiator 2'-O-riboyl phosphate transferase, is not required for normal cell growth, but a synergistic growth defect is observed in a rtx1 deletion strain also containing a reduced number of initiator methionine tRNA (IMT) genes [19].

Using a genetic screen based on synthetic lethality in a rtx1 mutant background to identify genes important for Pol III transcription, we have isolated mutations in the IVR1 gene. This was unexpected, because Iwr1 was originally identified by its physical association with Pol II [20,21], and it affects the basal and regulated expression of specific Pol II-transcribed genes [22], presumably through a direct effect on importing Pol II into the nucleus [23]. We show that Iwr1 is important for Pol III transcription, as an iwr1 mutant strain shows reduced association of TBP and Pol III at Pol III promoters, a decreased rate of Pol III transcription, and lower steady-state levels of Pol III transcripts. In addition, we show that Iwr1 is important for association of TBP to the Pol I-transcribed rDNA locus and for recruitment of TBP and Pol II to Pol II-transcribed loci. These data suggest that Iwr1 plays an important role in preinitiation complex formation by all three nuclear RNA polymerases in yeast.

Materials and Methods

Screen for mutants that require the RIT1 gene for growth

The genetic screen utilized to identify mutants requiring RIT1 for growth was based on a colony sectoring assay as described previously [24]. Candidate synthetic-lethal strains were crossed to UMY2395 and investigated for dominance/recessiveness and for 2:2 segregation of the non-sectoring phenotype. A YCp50 genomic library was used to transform iwr1-2 (UMY2299), rpc160-101 (UMY2304), and spb3-101 (UMY2309) mutants, and transfor-

Strains and DNAs

The source and genotypes of yeast strains used in this study are listed in table 1. Yeast strains and DNA molecules were constructed by standard methods, and the details are provided in Methods S1. Sequence analyses on chromosome IV revealed that iwr1-1 has an insertion of an adenine at position 255085, the iwr1-2 mutant allele carries an insertion of a thymine at position 254821, and in the iwr1-3 mutant there is a substitution from a guanine to a thymine at position 254368.

Immunofluorescence

To localize Iwr1, cells were grown in 5 ml YEPD at 30°C to an OD\(_{600}\) of 0.3, 670 µl formaldehyde (37%) was added and the cells were incubated for 40 min at RT. Cells were collected and washed once with 1× PBS, pH 7.4. The primary antibody, mouse anti-HA (12CA5), was diluted 1:2000, and the secondary antibody, goat anti-mouse linked to Cy3 (PA43002, Amersham Biosciences), was diluted 1:200. Cells were viewed in a Zeiss Axioscope 50 microscope using a 100 x objective. Images were acquired using a Hamamatsu-digital camera (C4742-95).

Polysome profiles

Cells were grown in 200 ml at 30°C in selective medium to an OD\(_{600}\) 0.4. Cycloheximide was added (100 µg/ml) 5 min before transferring the culture to an ice water bath for 15 min. Cells were collected at 4°C, washed twice in ice-cold breaking buffer (Bb; 20 mM Tris-HCl pH 7.4, 10 mM MgCl\(_2\), 100 mM KCl, 0.5 mM DTT, 100 µg/ml cycloheximide). The cells were resuspended in 1 volume of Bb, followed by addition of 1 volume of glass beads, and cells were disrupted by 6 x 20 sec on a vortex mixer, and the insoluble material was pelleted by centrifugation at 10,000 x g for 5 min at 4°C. The supernatant was transferred to a microfuge tube and subjected to a second centrifugation at 10,000 x g for 20 min at 4°C. The supernatant was applied to a 12 ml linear 10 to 45% sucrose gradient prepared in Bb lacking cycloheximide and centrifuged for 2.5 hrs at 40,000 rpm in a SW41 rotor at 4°C. The gradients were collected from the top, and A\(_{254}\) absorbances were monitored with the ISCO detection system.

RNA analysis

Northern blots were performed by standard methods using the following oligonucleotides for detection: GGGACATCAGGGT-TATGAGGC- (tRNA\(^{\text{Met}}\)); TGGCTCCAGGGGAGGCCGA-AC (tRNA\(^{\text{Met}}\))gCGGTTGTTCATCGAT (5.8S rRNA), and the levels of RNA quantified by phosphorimager analysis. For direct measurements, gels were stained with ethidium bromide, and RNA were quantified by using a BioRad Fluor-S TM Multimager and the QuantityOne-4.2.3 software. To analyze RNA levels by quantitative reverse-transcriptase PCR, total RNA was treated with DNase I on Qiagen RNaseasy columns, and first-strand cDNA synthesis was performed with random hexamers and Superscript III reverse transcriptase (Invitrogen) on 1 µg of total RNA. The relative representation of specific loci in this material was assayed by quantitative PCR in real-time on an Applied Biosystems 7500 machine.

In vivo labeling of RNA

In labeling experiments, cells were grown to an OD\(_{600}\) of 0.8 in 120 ml SC-uracil medium at 30°C before 125 µCi of 'H-Uridine was added (33 Ci/mmol, Amersham Biosciences). Samples (20 ml) were collected after 5, 10, 20, and 40 min. Total RNA was prepared and separated on an 8% polyacrylamide 8 M urea gel. The gel was stained with ethidium bromide, quantified, soaked in NAMP100 Amplifier for 15 min (Amersham Biosciences), dried onto 3 MM Whatman paper, and exposed to film. Signals were quantified using QuantityOne-4.2.3 software (BioRad), and the rate of synthesis for each RNA type was calculated. For the pulse-chase experiment, cells were grown to an OD\(_{600}\) of 0.8 in 50 ml SC-uracil medium at 30°C and pulse-labeled using 125 µCi of 'H-Uridine (33 Ci/mmol Amersham Biosciences) for 45 min. Cells were collected and resuspended in 250 ml pre-warmed SC medium containing excess (2 mM) uracil to begin the chase. Samples (40 ml) were collected after 0, 1, 2, 3, and 4 hrs. RNA was prepared, separated, and quantified as described above.

Immunoprecipitation

50 µg total protein in buffer 1 (0.15 M Tris-HCl, pH 7.8, 50 mM KAc, 20% glycerol, 1 mM EDTA, 1 × Protease inhibitors, 0.5 mM DTT) was incubated for 2 hrs with agarose beads
Table 1. Yeast strains used in this study.

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(Sepharose 4 Fast Flow, Amersham Biosciences) in a rotating chamber. Beads were recovered and the supernatant was transferred to tubes containing agarose beads linked to either anti-HA or anti-MYC antibody (Sigma A2095 or M5546), and incubated 2 hrs at 4°C in a rotating chamber. Beads were recovered and washed six times using 1 ml of buffer 2 (0.15 M Tris-HCl, pH 7.8, 50 mM KAc, 20% glycerol, 1 mM EDTA, 1× Protease inhibitors, 0.5 mM DTT, 0.5% Triton-X100), and proteins bound to the beads were recovered by incubating samples at 100°C for 3 min in 1× loading buffer. Recovered tagged proteins were detected using standard western blot techniques.

Chromatin immunoprecipitation

Yeast strains BY4741 and the isogenic strain containing an iwr1 null allele RN3812 were obtained from Invitrogen. Cells were crosslinked with 1% formaldehyde and total chromatin was sonicated to an average size between 300–500 bp as described previously [23]. Chromatin immunoprecipitation was performed using antibodies against Tfc4, Rpc34 (both antibodies kindly supplied by Steve Hahn), Bdpl (kindly provided by Ian Willis), and Rpb1 (8WG16 antibody from Covance). Immunoprecipitated DNA and total input control DNA were assayed by real-time quantitative PCR using the Applied Biosystems 7500 Real-time PCR System. Immunoprecipitation efficiency was determined for each locus by dividing the yield of PCR product in the immunoprecipitation sample by the amount of product obtained from the input control. Relative occupancy values were determined by dividing the immunoprecipitation efficiency at each locus by the immunoprecipitation efficiency at a negative control locus (either the middle of the POL1 ORF or an ORF-free region of chromosome V). The occupancy value of the negative control, 1.0, was subtracted from all values to yield a baseline of 0. All occupancy values were normalized to set the occupancy of each factor in the wild-type strain at 1.0, and the occupancy at 1.0 is the occupancy of each factor in the negative control strain.
Results

Strains with a mutation in *IWR1* require the *RIT1* gene for growth

We reasoned that screening for mutations that are synthetic-lethal in combination with a *rit1Δ* mutant might identify genes required for efficient Pol III transcription. A *rit1Δ* mutant is viable with no apparent growth defects unless the steady-state level of tRNA\text{Met}\_i is reduced by deleting *IMT* genes, whereupon severe/modest growth defects are observed [19]. The genetic screen that was employed here to identify mutants requiring Rit1 for growth has been described in detail [24,26]. Briefly, a *rit1Δ* null strain carrying a plasmid with the *RIT1* gene was mutagenized, and colonies were screened for the inability to lose the *RIT1* plasmid. Thirty-two strains were identified that required the *RIT1* gene for survival. Here we describe one complementation group consisting of three slow growing and temperature sensitive (ts) mutants.

To identify the gene mutated, a genomic library was introduced into one of the mutants, and plasmids complementing the *RIT1* dependence, ts, and slow growth phenotypes were identified. DNA sequencing and sub-cloning showed that the *IWR1* gene was responsible for the complementation. Genetic linkage between *IWR1* and the three original *iwr1* mutations was confirmed by targeted integration and tetrad analysis. To identify and characterize the *iwr1* mutations, we cloned and sequenced the gene from the three different mutant alleles. All three *iwr1* mutations generated truncated Iwr1 proteins of various lengths (Fig. 1A) (see Materials and Methods). When yeast Iwr1 was compared to translated ORFs from multi-cellular eukaryotes, potential homologues were identified. All identified proteins shared three distinct regions with high similarity (Fig. 1B). Thus, the *IWR1* gene encodes a conserved protein that is required for viability in a *rit1Δ* null strain.

The *iwr1*-2 mutant has reduced Pol III transcripts and a defect in initiation of translation

A *rit1Δ* null strain with two of the four *IMT* genes deleted displays a synergistic growth defect that is overcome by increased gene dosage of *IMT* [19]. Introduction of a high-copy number plasmid carrying *IMT* into an *iwr1* mutant suppressed the *RIT1* dependence for growth (data not shown). Suppression was specific for *IMT* genes, as *iwr1* mutants with increased dosage of the *EMT5* gene, encoding elongator methionine tRNA (tRNA\text{Met}\_m), required *RIT1* for growth (data not shown). This suggests that the *RIT1* dependence of the *iwr1* mutant might be caused by reduced levels of tRNA\text{Met}. To investigate the level of tRNA\text{Met} in the *iwr1*-2 mutant, northern blot were conducted using total RNA isolated from wild-type and *iwr1*-2 strains. The blots were probed for tRNA\text{Met}\_i, tRNA\text{Met}\_m, and the Pol I transcript 5.8S rRNA. The *iwr1*-2 mutant showed a ten-fold reduction in the level of tRNA\text{Met}\_i and a three-fold reduction of tRNA\text{Met}\_m as compared with the isogenic wild-type (Fig. 2A).

As the levels of both tRNA\text{Met}\_i and tRNA\text{Met}\_m decreased in the *iwr1*-2 mutant, we considered that all Pol III transcripts could be affected. Total RNA isolated from wild-type and *iwr1*-2 strains was separated on a denaturing polyacrylamide gel, stained with ethidium bromide, and the total tRNA and 5S rRNA signals were quantified after normalization to 5.8S rRNA. The *iwr1*-2 mutant showed a three-fold reduction in total tRNA levels and a 40% reduction in 5S rRNA compared to wild-type (Fig. 2B). We also investigated the levels of total tRNA and 5S rRNA in the *iwr1*-1 mutant.
and iwr1-3 mutants and found that total tRNA was 64% and 22% reduced, and 5S rRNA was 35% and 31% reduced, respectively (data not shown). As reduced levels of tRNA<sub>Met</sub> affect initiation of translation, we expected that the iwr1-2 mutant should have a reduction in initiation of translation. Consistent with this idea, the iwr1-2 mutant showed a six-fold reduction of polysome to monosome ratio when compared to wild-type (Fig. 2C). We conclude that the iwr1 mutant has decreased levels of Pol III-transcribed RNAs, and it is the low level of tRNA<sub>Met</sub> that likely causes the RIT1 dependence and defect in translation initiation.

**Iwr1 is important for Pol III transcription**

The reduction in tRNA and 5S rRNA levels in the iwr1 mutants could be caused by defects in stability, transcription, or a combination thereof. To test whether Iwr1 is needed for stability of Pol III-transcribed RNAs, a pulse-chase experiment was performed in wild-type and iwr1-2 strains using <sup>3</sup>H-uridine. Samples were taken at various times after the chase began, and total RNA was separated on a denaturing polyacrylamide gel. <sup>3</sup>H-labeled tRNA and 5S rRNA were quantified after being normalized to the 5.8S rRNA. The iwr1-2 mutant showed reduced levels of tRNA and 5S rRNA at time point 0; however, there was no further reduction during the 4 hr chase, indicating no decreased stability of tRNA and 5S rRNA (Fig. 3A). To investigate whether the iwr1-2 mutation influences the rate of transcription by Pol III, wild-type and iwr1-2 strains were cultured with <sup>3</sup>H-uridine and samples removed at various time points after <sup>3</sup>H-uridine addition and analyzed for tRNA, 5S rRNA, and 5.8S rRNA accumulation. In the iwr1-2 mutant the incorporation of <sup>3</sup>H-uridine into tRNA and 5S rRNA was reduced to 43% and 64% (k-value) (Fig. 3B) compared to wild-type, whereas the rate of 5.8S rRNA synthesis was virtually unaffected (94%). We conclude that Iwr1 is required for the efficient production of tRNA and 5S rRNA.

The **IWR1 gene is genetically linked to the Pol III machinery**

From the screen for mutants requiring the RIT1 gene, we also obtained strains with mutations in the RPC160 or the RPB5 gene. The RPC160 gene encodes the large subunit of RNA polymerase III, and the RPB5 gene encodes one of five subunits that are shared among all three RNA polymerases [27]. In addition to the RIT1 dependence, the rpc160-101 and rpb5-101 mutants showed a reduced growth rate and a 35% and 31% reduction in total RNA levels, respectively (data not shown). To further investigate the role of Iwr1 in transcription by Pol III, we combined the iwr1-2 allele with an rpc160-101 or an rpb5-101 mutation. The combinations were lethal, consistent with a role of Iwr1 in transcription by Pol III (Fig. 4A). Increased gene dosage of RPC160, but not RPB5, in

**Figure 2. The iwr1-2 mutant is defective for Pol III transcription and translational initiation.** (A) Northern blots of RNA isolated from wild-type (UMY2220) and iwr1-2 (UMY2808) strains that were probed for tRNA<sub>Met</sub><sup>wt</sup>, tRNA<sub>Met</sub><sup>m</sup>, and 5.8S rRNA. (B) Ethidium bromide staining of total RNA from wild-type (UMY2220) carrying pRS316 and iwr1-2 (UMY2808) strains carrying pRS316 or pRS316-IWR1. Band intensities were normalized to the 5.8S rRNA signals, and the amount of each RNA species was expressed relative to the corresponding value in the wild-type strain, which was set to 1. (C) Polysome profiles of total extracts isolated from wild-type (UMY2220) carrying pRS316, and iwr1-2 (UMY2808), carrying pRS316 or pRS316-IWR1 were analyzed by sedimentation in sucrose gradients. The polysome to monosome ratio (P/M) was calculated.

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the iwr1-2 mutant suppressed the RIT1 dependence and the low-level of total tRNA and 5S rRNA, and it partially suppressed the growth defect at 30° and 37° (data not shown, Fig. 4B).

Maf1 is the main regulator for signaling pathways to mediate repression of transcription by Pol III (For review see [13]). A maf1 mutant is unable to repress transcription by Pol III in response to various stress signals. To investigate whether the reduced rate of Pol III transcription observed in the iwr1-2 mutant is mediated through Maf1, we introduced a maf1 null allele into the iwr1-2 mutant. The double mutant showed similar growth phenotype (Fig. 4C) and steady-state levels of tRNA and 5S rRNA as the single iwr1-2 mutant (Fig. 4D). Together these genetic data support a role for Iwr1 in transcription by Pol III and show that the iwr1 mutant affects transcription by Pol III independently of the key negative regulator, Maf1.

Iwr1 is a nuclear protein that may interact weakly with Pol III

To investigate the intracellular localization of Iwr1, we epitope-tagged the genomic copy of IWR1 with the hemagglutinin epitope (HA). By indirect immunofluorescence, Iwr1-HA localized predominantly to the nucleus (Fig. 5A). Immunoprecipitation of HA-tagged Rpe160 co-precipitated Iwr1, suggesting a physical interaction between Iwr1 and Pol III (Fig. 5B). Gavin et al. (2002) have previously shown that Iwr1 co-purified with the Pol II specific subunit, Rpb3. However, no Rpb3 was detected in our Rpe160 immunoprecipitate, excluding the possibility that contaminating Pol II subunits were responsible for the presence of Iwr1 (Fig. 5B). Immunoprecipitation of a 13-myc-tagged Iwr1 co-precipitated Rpe160, but with very low efficiency (data not shown). By an Rpe160 immunodepletion experiment we showed that only 5–10% of total Iwr1 was associated with Rpe160 (Fig. 5C). To confirm that the Pol III complex remained intact throughout the experiment, we used the Pol III specific subunit Rpe82 as a control (Fig. 5C). Even though Iwr1 and Pol III do co-precipitate, the low level of association suggests an indirect or weak interaction.

Reduced Pol III occupancy on tRNA genes in an iwr1 mutant

In addition to the polymerase, the Pol III transcription machinery consists of the multi-subunit transcription factors TFIIIB and TFIIIC [1,4,12]. TFIIIC consists of six subunits, which recognize the A- and B-boxes and promote recruitment of the initiation factor, TFIIIB, to the region upstream of the transcriptional start site. The recruited TFIIIB cooperates with TFIIIC to recruit the RNA polymerase to the transcriptional start site [28]. To investigate whether lack of Iwr1 function affects the Pol III transcription factor occupancy profile at Pol III-transcribed
loci, we performed chromatin immunoprecipitation (ChIP) on wild-type and iwr1D strains using antibodies against the TFIIIB components TBP and Bdp1, the TFIIIC component Tfc4, and the Pol III-specific polymerase subunit Rpc34. We assayed the occupancy of each factor at a variety of Pol III loci: SCR1H, SNR6, tC(GCA)B, EMT5, IMT2, and IMT4. In the iwr1D strain, Bdp1 occupancy is essentially unchanged, TBP occupancy is reduced by 30 to 50%, Rpc34 occupancy is slightly decreased, and Tfc4 occupancy is normal to slightly increased (Fig. 6). This suggests that Iwr1 facilitates the recruitment of TBP and the RNA polymerase to Pol III-transcribed loci. We note that Bdp1 and TBP, which are both subunits of TFIIIB, show different occupancy profiles in the iwr1D strain; it is possible that deletion of Iwr1 affects the stability of the TBP-DNA association, while Bdp1 occupancy is maintained through its contacts with other components of the Pol III apparatus.

Iwr1 is important for recruitment of TBP to Pol I, Pol II, and Pol III promoters

Because TBP participates in transcription by all three RNA polymerases and shows diminished occupancy at Pol III loci in the iwr1D strain, we also tested TBP occupancy by ChIP at Pol I- and Pol II-transcribed loci in wild-type and iwr1A strains. We found that TBP occupancy at Pol II loci is approximately two-fold lower in an iwr1A mutant strain, indicating that the iwr1A defect is not specific to Pol III transcription (Fig. 7A). Furthermore, occupancy by TBP at the Pol I-transcribed rDNA locus (RDN37) is also somewhat lower in the mutant strain (Fig. 7A). Iwr1 is therefore important for recruitment of TBP to loci transcribed by all three nuclear RNA polymerases in yeast.

Pol II occupancy is reduced in the iwr1A mutant

In light of the known interaction between Iwr1 and RNA Pol II [20,21] and the effect of the iwr1A mutation on TBP occupancy at Pol I-transcribed loci, we tested occupancy by the Pol II subunit Rpb1 in wild-type and iwr1A strains. We performed ChIP analysis on ten different Pol II-transcribed rDNA loci (RD37) in strains deleted for the IWR1 gene. We observed a similar occupancy decrease on the Pol II loci as observed on Pol III-transcribed loci, i.e., recruitment of the polymerase was reduced in the iwr1 null strain (Fig. 7B). Thus, two independent observations, namely reduced association of the polymerase and of TBP at Pol II-transcribed loci, strongly suggest that Iwr1 functions in transcriptional initiation by Pol II. After completion of our

Figure 4. The IWR1 gene is genetically linked to Pol III transcription. (A) Wild-type (UMY2220), iwr1-2 (UMY2808), rpc160-101 (UMY2802), iwr1-2 rpc160-101 (UMY2984), rpc5-101 (UMY2804), and iwr1-2 rpc5-101 (UMY2986) strains all carrying pRS316-IWR1 were grown in synthetic complete (SC) medium, serially diluted, spotted onto SC and SC+5-FOA plates. (B) Total RNA isolated from wild-type (UMY2220) and iwr1-2 (UMY2808) strains carrying either pRS316 or pRS316-RPC160 was analyzed as in Figure 2B. (C) Growth of wild-type (UMY2220), maf1Δ (UMY2975), iwr1-2 (UMY2808), and maf1Δ iwr1-2 (UMY3059) strains on YEPD plates. (D) Total RNA isolated from wild-type (UMY2220), maf1Δ (UMY2975), iwr1-2 (UMY2808), and maf1Δ iwr1-2 (UMY3059) strains was analyzed as in Figure 2B. doi:10.1371/journal.pone.0020829.g004
work, it was reported that Iwr1 is directly involved in the import of Pol II into the nucleus [23].

The Pol III transcription defect is not caused by a Pol II transcriptional defect

To determine whether the polymerase occupancy decrease at Pol II-transcribed loci resulted in lower transcript levels of Pol II-transcribed genes, we analyzed the levels of eleven mRNAs by rtPCR (Fig. 8A). We included genes encoding components of the Pol III transcription machinery to determine whether the decreased abundance of these Pol II-transcribed factors might be the indirect cause of the observed defect in Pol III transcription. Strikingly, we did not observe significant differences in the steady-state level of RNA synthesized by Pol II in iwr1 and wild-type cells (Fig. 8A). The TBP and polymerase recruitment defect at Pol II-transcribed loci in the iwr1 Δ strain does not alter the steady-state level of the mRNAs tested, suggesting that a post-transcriptional mechanism is able to compensate for any resulting decrease in transcription. Furthermore, the normal levels of the mRNAs encoding the Pol III transcription factors Brf1, Tfc6, Rpc160, and Rpc34 make it clear that the observed decrease in transcription by Pol III in the iwr1 Δ strain is not an indirect effect of diminished transcription of the Pol III machinery itself. In addition, western blot analysis of HA-tagged Rpc160 revealed no difference in the levels of Rpc160 between wild-type and iwr1-2 strains (Fig. 8B). Thus, the defect in Pol III transcription in the iwr1 mutant strain is not merely a trivial consequence of diminished Pol III transcription factor availability.

Discussion

We isolated the IWR1 gene in a genetic screen for mutants requiring the RIT1 gene for growth. The RIT1 gene encodes an initiator tRNA specific 2'-O-riboosyl phosphate (Ar(p)) modification enzyme [29], and the presence of the Ar(p) modification prevents tRNAMet from being utilized in translation elongation [16–18]. Under conditions in which the tRNAMet steady-state level is reduced, the Ar(p) modification is required to maintain a sufficient level of eIF2:GTP:tRNAMet ternary complex to promote efficient translational initiation [19]. In the iwr1-2 mutant the tRNAMet steady state level of tRNAMet is reduced ten-fold, thereby dramatically affecting initiation of translation (Fig. 2A and C). In an iwr1 mutant the RIT1 gene is essential, as use of tRNAMet lacking Ar(p) for translation elongation would reduce the already limited pool of tRNAMet available for initiation to below the minimum level required for cell survival.

We found an iwr1 mutant strain to have a significant reduction in steady state levels of tRNA and 5S rRNA. Interestingly, the large decrease in Pol III transcript levels in the iwr1 Δ strain (Fig. 2) occurs even though there is only a moderate reduction in the association of transcription factors with Pol III-transcribed loci (Fig. 6). It therefore appears that, while Iwr1 has a modest effect on transcription factor association, it exerts a more significant influence on the process of transcription itself. Accordingly, we observed a reduced rate of Pol III-transcribed RNA synthesis in the iwr1 Δ mutant strain (Fig. 3). Interestingly, the occupancy profile of the Pol III general transcription factors in the iwr1 Δ strain (namely decreased polymerase occupancy accompanied by a slight increase in TFIIIC occupancy) is reminiscent of that observed in yeast strains subjected to conditions repressive for Pol III transcription [30].
The reduction in both TBP and polymerase association with Pol II-transcribed loci in the iwr1 mutant is indicative of a defect in preinitiation complex formation and hence likely in the rate of transcriptional initiation. Indeed, after the work in this paper was completed, it was demonstrated that Iwr1 is directly involved in the nuclear import of Pol II [23]. However, we observe no significant consequence for steady-state mRNA levels. This suggests that, in contrast with the situation at Pol III-transcribed loci, there is a compensatory mechanism, presumably at the level of transcript stability either during or after completion of the RNA, to regulate mRNA levels. A similar discordance between Pol II occupancy and RNA levels has also been observed in yeast cells lacking either the Swi/Snf nucleosome remodeling complex [31] or the Asf1 histone chaperone [32], suggestive of a general compensation mechanism that is not specific to Iwr1. Similarly, although TBP occupancy is lower at the Pol I-transcribed rDNA locus in the iwr1 strain, we saw no decrease in Pol I-transcribed ribosomal RNA levels, including levels of precursor RNA (Fig. 8). Taken together, these results indicate that, although Iwr1 influences transcription factor occupancy at promoters controlled by all three RNA polymerases, iwr1 mutant strains are specifically defective in accumulating Pol III-transcribed RNAs, thereby explaining the Pol III-specific phenotypes.

Figure 6. Occupancy profile of Pol III factors in the iwr1 null strain. Chromatin immunoprecipitation was performed on chromatin from wild-type and iwr1Δ strains using antibodies against TBP, Bdp1, Tfc4, and Rpc34. Immunoprecipitation efficiency was determined for each Pol III locus by dividing the yield of PCR product in the immunoprecipitated sample by the amount of product obtained from the input control. Relative occupancy values were determined by dividing the immunoprecipitation efficiency at each locus by the immunoprecipitation efficiency at a negative control locus.

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In this regard, in the plant Arabidopsis thaliana, the Iwr1 homolog affects transcription by Pol IV and Pol V, which are involved in RNA-directed DNA methylation [33]. It is unlikely that the reduction in Pol III- and Pol I-containing preinitiation complexes is due to a defect in transcription by Pol II, because mRNA levels of all genes tested (including many encoding components of Pol III factors) are unaffected in an iwr1 mutant strain.

Our efforts to perform ChIP on the Iwr1 protein itself with either N- or C-terminal epitope tags were not successful, suggesting that Iwr1 may not function directly at promoters or that its interaction is transient. The likelihood of an indirect function or transient interaction is supported by the low co-precipitation efficiency between Iwr1 and Pol III, and by the displacement of Iwr1 from Pol II by transcription initiation factors and DNA [23]. Interestingly, Iwr1 has been found in large-scale experiments to have genetic interactions with a significant number of chromatin modifying and remodeling proteins, including several components of the Swi/Snf complex, four members of the Swr1 complex, which exchanges histone variant H2AZ for H2A, and H2AZ itself [34]. A connection of Iwr1 to chromatin structure is further suggested by the compensatory mechanism for mRNA levels that is also observed in strains lacking Swi/Snf or Asf1 [31,32]. These observations lead to the speculation that the effect of Iwr1 on TBP recruitment might be attributable to a possible influence on promoter accessibility mediated through these interactions with chromatin-associated proteins. Alternatively, Iwr1 might affect the nuclear import of TBP, the TBP-containing complex TFIIIB that is required for transcription by Pol III, or other factors affecting transcription and/or chromatin structure.
Figure 7. Occupancy profile of TBP and RNA polymerase in the wild-type and *iwr1* null strain. (A) TBP occupancy at the Pol I-transcribed RDN37 locus and at various Pol II-transcribed loci in wild-type and *iwr1* strains. (B) Occupancy by the large subunit of RNA Polymerase II at various loci in wild-type and *iwr1* strains. The occupancy values are expressed as folds over the values at an ORF-free control region. *a-* corresponds to promoter region.
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Figure 8. Relative expression of RNAs in *iwr1* and wild-type strains. (A) Relative expression of various regions of the long Pol I-transcribed region (left) and of a variety of Pol II-transcribed RNAs (right). The regions on the left were chosen to represent both precursor RNA (regions removed during RNA maturation) and mature RNA. (B) Western blot analysis of the RPC160-3HA (UMY3031) and the *iwr1-2* RPC160-3HA (UMY3241).
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Supporting Information

Methods S1 Additional methodological details beyond those in the Materials and Methods section.

(PDF)

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References


Author Contributions

Conceived and designed the experiments: AE ZM KS AB. Performed the experiments: AE ZM XF JL. Analyzed the data: AE ZM KS AB. Contributed reagents/materials/analysis tools: AE ZM XF JL. Wrote the paper: AE ZM KS AB.