

A Transcription Factor IIA-Binding Site Differentially Regulates RNA Polymerase II-Mediated Transcription in a Promoter Context-dependent Manner

Juan Wang^{§,*}, Shasha Zhao^{§,*}, Wei He[§], Yun Wei[§], Yang Zhang[§], Henry Pegg[†], Paul Shore[†], Stefan G.E. Roberts^{#,1}, Wensheng Deng^{§,2}

From the [§]Institute of Biology and Medicine, Wuhan University of Science and Technology, Wuhan City, Hubei Province, 430065, China; the[†]The School of Biological Sciences, University of Manchester, Manchester, M13 9PL, UK and the [#]School of Cellular and Molecular Medicine, University of Bristol, Bristol, BS8 1TD, UK.

Running title: Role of TFIIA recognition element in transcription

*: J.W. and S.Z. contributed equally to this work.

1: Address correspondence to Stefan GE Roberts, stefan.roberts@bristol.ac.uk

2: Address correspondence to Wensheng Deng, dengwensheng@wust.edu.cn, Tel: 18571542248

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Abstract

RNA polymerase II polymerase II (pol II) is required for the transcription of all protein-coding genes and as such represents a major enzyme whose activity is tightly regulated. Transcriptional initiation therefore requires numerous general transcriptional factors and cofactors that associate with pol II at core promoter to form a pre-initiation complex. Transcription factor IIA (TFIIA) is a general cofactor that binds TFIID and stabilizes the TFIID-DNA complex during transcription initiation. Previous studies showed that TFIIA can make contact with the DNA sequence upstream or downstream of the TATA-box, and that the region bound by TFIIA could overlap with the

elements recognized by another factor, TFIIB, at adenovirus major late core promoter. Whether core promoters contain a DNA motif recognized by TFIIA remains unknown. Here we have identified a core promoter element upstream of the TATA box that is recognized by TFIIA. A search of the human promoter database revealed that many natural promoters contain a TFIIA recognition element (IIARE). We show that the IIARE enhances TFIIA-promoter binding and enhances the activity of TATA-containing promoters, but represses or activates promoters that lack a TATA box. Chromatin immunoprecipitation assays revealed that the IIARE activates transcription by increasing the recruitment of pol II, TFIIA, TAF4 and p300 at TATA-dependent promoters. These findings extend

our understanding of the role of TFIIA in transcription, and provide new insights into the regulatory mechanism of core promoter elements in gene transcription by pol II.

Eukaryotic transcription initiation by RNA polymerase II requires numerous transcription factors and cofactors to nucleate at core promoter to form a pre-initiation complex (1, 2). The core promoter plays a critical role during transcriptional initiation and contains a number of DNA sequence elements such as the TATA box, the initiator, the downstream promoter element, the TFIIB recognition elements (BREs) and others (1-4). These elements are recognized by general transcription factors and cofactors (5-9), and assist to direct and orientate pre-initiation complex formation at the promoter. Core promoter elements not only regulate the activity of transcription but also determine transcription start site selection (4, 10 and 11). Genome-wide studies have revealed that many genes lack so-called ‘canonical’ core promoter elements, suggesting that other core promoter elements remain to be discovered. Indeed, a recent study showed that core promoter element, DTIE, directs transcription start site selection of genes with TATA-less promoters (12). Nevertheless, it has been proposed that core promoter elements may not be essential to transcription of some genes *in vivo* (13), suggesting that canonical core promoter elements fine-tune physiological responses for specific genes (4). It has been shown that many ‘noncanonical’ promoters instead contain epigenetic marks including histone modifications (H3K4me3 and H3K27me3) and DNA marks such as enhancers, CpG islands and ATG deserts (14-16). The mechanisms by which canonical or noncanonical core promoters regulate transcriptional initiation are not fully understood.

Transcription factor TFIIA comprises of three subunits, α , β and γ ; TFIIA α/β and TFIIA γ are encoded by different genes (17-19). The precursor of TFIIA α/β can be digested by *taspase1*, but uncleaved TFIIA α/β remains active in transcriptional regulation (20). Recent studies showed that the cleavage of TFIIA by *taspase 1* is involved in a number of molecular and biological processes (21-24). Although TFIIA was originally characterized as a general transcription factor, TFIIA is dispensable in transcription *in vitro* (25-26); perhaps, TFIIA is better to be described as a general cofactor because it acts as an anti-repressor or co-activator in transcriptional regulation (27-31). TFIIA can counteract the inhibitory roles of TAF1 and BTAF1 during TBP binding to the TATA box as well as the repressive effects of NC2 and HMGB1 on transcription (27, 28). TFIIA has also been shown to stabilize TFIID binding to DNA by interacting with transcriptional activators, TBP-associated factors (29, 30, 32-34) and TBP-related factors (35-37). It has been proposed that TFIIA induces the disassociation of TBP dimers and promotes the association between TBP and the TATA-box promoter (38). TFIIA stabilizes the TBP-TATA box complex through direct contact on the face of TBP opposite to the TFIIB-binding side (39, 40). Studies using cryo-electron microscopy revealed that TFIIA and the transcription activator Rap1 cooperatively commit TFIID in transcription initiation (41).

Previous studies showed that human TFIIA can make specific contacts with the DNA immediately upstream of the TATA box (4, 39 and 42-44). In our previous work, mutations of BRE consensus bases within the AdML promoter inhibit the formation of TFIIA-TBP-DNA complexes, suggesting that a sequence-specific TFIIA DNA-binding region might overlap with the sequence of the BRE (45). In this study, we confirm that TFIIA makes direct contact with the

sequence immediately upstream of the TATA box at the adenovirus major late (AdML) promoter using combined molecular approaches. Using this information, we identified a core promoter element upstream of the TATA box that is recognized by TFIIA. We show that the TFIIA recognition element regulates transcription activity in a promoter context-dependent manner and determine the mechanism by which the TFIIA recognition element regulates transcription at the AdML promoter.

Results

TFIIA makes direct contact with the DNA sequence upstream of the TATA box at the AdML promoter — Our previous work showed that mutations of BRE^u and BRE^d consensus bases within the AdML promoter (AdML-mBRE^{ud}, Fig. 1A) inhibit the formation of a TFIIA-TBP-DNA complex when compared to wild type AdML promoter, suggesting that a TFIIA DNA-binding region could overlap with the BRE^u or BRE^d of the AdML promoter (45). To exclude the possibility that mutations of BRE^{ud} consensus bases might affect TBP binding to the promoter DNA, bandshift assays were performed using TBP and radio-labelled promoter DNA fragments. As shown in Fig. 1B, the formation of TBP-DNA complexes showed minimal difference between the wild type AdML and AdML derivatives, indicating that mutations of the BRE^u or BRE^d consensus bases within the AdML promoter do not affect the formation of a TBP-DNA complex. To confirm this observation, protein-DNA binding assays were performed using HEK293T nuclear extract and promoter DNA immobilized on magnetic beads. TFIIA showed decreased binding to the AdML-mBRE^{ud} promoter when compared with the wild type AdML promoter. However, TBP binding to DNA showed little difference between these two promoters (Fig. 1C), confirming that mutations of BRE^{ud}

consensus bases do not affect TBP binding to DNA, but significantly reduce TFIIA recruitment. To determine the specific contribution of the BRE^u and BRE^d elements to TFIIA recruitment, bandshift assays were performed using native hTFIIA, recombinant hTBP and radio-labelled DNA fragments containing either a mutated BRE^u or a mutated BRE^d (Fig.1A). The result shows that mutations in the BRE^u but not the BRE^d reduced the formation of a TFIIA-TBP-DNA complex compared with the wild type AdML promoter (Fig. 1D). This suggests that TFIIA DNA-binding region overlaps with the BRE^u at the AdML promoter. To determine if TFIIA interacts with specific bases within the BRE^u, DNA methylation interference assays were performed using native hTFIIA, recombinant hTBP and radio-labelled AdML promoter DNA fragments. The data show that two G bases immediately upstream of the TATA box displayed weak signals at the positive strand of AdML (the black solid dots in lane IIA, Fig. 1E). This observation was verified by density quantification for the G tracks (-33 and -34 in G tracks, Fig. 1F); indicating that TFIIA can interact with at least two bases upstream of the TATA box. Taken together, these data confirm that TFIIA makes direct contact with the sequence immediately upstream of the TATA box at the AdML promoter.

Derivation of TFIIA recognition element upstream of the TATA box and its prevalence in natural promoters — Previous studies using photocrosslinking showed that TFIIA subunits, α and β , interact with a number of bases immediately upstream of the TATA box (on both positive and negative strands) at the AdML promoter (42). Our methylation interference analysis revealed the contacts between TFIIA and the two G bases of the positive strand upstream of the AdML TATA box (the black solid dots in Fig. 2A). Using this

information, we generated a core promoter DNA library that contains randomized bases between position -41 and -32 within AdML promoter derivatives (Fig. 2A). The randomized DNA was labelled with [α - 32 P]- ATP and subjected to protein-DNA binding selection through bandshift assays (SELEX). After seven rounds of SELEX, the efficiency of selection was examined by bandshift assay. Fig. 2B shows that formation of a TFIIA-TBP-DNA complex was significantly enhanced, indicating that the affinity of TFIIA to DNA was increased by SELEX. The DNA fragments were then cloned into pGEM3 and 50 clones sequenced across the promoter region. A TFIIA-binding consensus sequence was determined by the frequency of occurrence for each base. As shown in Fig. 2C, the consensus sequence exhibited strong preference for G or T, with exclusion of A and C at the region between -41 and -39 and the region between -35 and -32. Based on this information, we derived a consensus sequence:

5'-G-T/G-G-G/C/A-G/C-A/G-T-G/T-G/T-T-3'.

Here we refer to the consensus sequence upstream of the TATA box as TFIIA recognition element (this element is abbreviated as IIARE, intentionally distinguished from the ARE (Androgen Response Element)). The IIARE displayed four-base identity to the BRE^u (-38 to -32), and six-base identity to wild type AdML (-41 and -32). To gain insight into the prevalence of the IIARE in natural promoters, we searched the promoters from human promoter database (<http://epd.vital-it.ch/>).1957 TATA-containing promoters and 2058 TATA-less promoters were used to analyse the presence of IIARE consensus bases in the region between -41 and -32 at the natural promoters. The result shows that 50.3 % of the TATA-containing promoters contain over 5 consensus bases of the IIARE, 26.5% of promoters over 6 consensus bases and 9.96% of promoters

over 7 consensus bases; However, the IIARE showed a higher prevalence at the TATA-less promoters, particularly where the number of IIARE consensus bases is ≥ 7 or 8 (Table 1). The prevalence of the IIARE in TATA-less natural promoters is similar to the 5-8 % of natural promoters that contain a TATA box (1). These results suggest that the IIARE is potentially present in a large proportion of natural promoters. Whether the IIARE in these promoters is functional requires further study.

The IIARE mutation in TATA-dependent promoters inhibits TFIIA recruitment and promoter activity—

To determine if the IIARE consensus that we derived affects the affinity of TFIIA to DNA, we generated AdML promoter derivatives that contain an optimal IIARE(AdML-IIARE) or a defective IIARE (AdML-mIIARE) (Fig.3A). Promoter DNA was immobilized with streptavidin magnetic beads and binding assays were performed using reconstituted hTFIIA and recombinant hTBP. Stably bound proteins were detected by Western blot. As shown in Fig. 3B, TFIIA exhibited decreased binding to the promoter containing a defective IIARE (AdML-mIIARE) when compared with the promoter containing an optimal IIARE (AdML-IIARE). Thus, the IIARE within AdML promoter derivatives enhances TFIIA binding to DNA, which is consistent with the observation from the random selection (Fig. 2B). To determine the effect of the IIARE on the activities of these promoters, *in vitro* transcription assays were performed using HEK293T nuclear extract and reporter vectors driven by either the AdML-IIARE or AdML-mIIARE promoter. Since the vector does not contain any eukaryotic promoter except the tested promoter, transcripts can be detected using RT-qPCR as described previously (46). The AdML-mIIARE promoter showed significant reduction in transcriptional

activity in either the absence or presence of Gal4-VP16 when compared with the AdML-IIARE promoter or the wild type AdML promoter (Fig. 3C; left and right panel). To verify these observations, transient transfection of the promoter-driven reporters was performed with HEK293T cells in the absence or presence of Gal4-VP16 and promoter activity determined by luciferase assay. The IIARE defective promoter, AdML-mIIARE, showed reduced activity regardless of the co-expression of Gal4-VP16 when compared with the AdML-IIARE promoter or the AdML-WT promoter (Fig. 3D). Taken together, these data indicate that the IIARE within the AdML promoter is required for efficient TFIIA recruitment and positively regulates promoter activity.

We next determined if our observations can be extended to natural promoters that contain a IIARE consensus sequence. A search of the human promoter database revealed that the cytochrome P450 1A2 (hCYP1A2) promoter contains a typical IIARE and a canonical TATA box (Fig. 4A). DNA fragments containing either the wild type hCYP1A2 core promoter or a mutant IIARE derivative were synthesized and cloned into the reporter vector pGL3-basic. Using these DNA fragments and vectors-based sequences, protein-DNA binding assays, *in vitro* transcription assays and luciferase assays were performed as described in Fig.3. The results from these experiments showed that an intact IIARE within the hCYP1A2 promoter is required for TFIIA recruitment (Fig. 4B) and efficient promoter activity (Fig. 4C-E), confirming a positive role for the IIARE in TATA-containing promoters.

The promoters of AdML-WT and hCYP1A2-WT also contain a BRE^u consensus sequence, raising the possibility that the IIARE-defective promoters also affect the BRE^u consensus bases. To exclude the possibility of

interference by mutation of BRE^u, the natural promoter that directs expression of farnesylidiphosphate synthase (FDPS1) was tested. The FDPS1 promoter contains a TATA box and five IIARE consensus bases, but only one BRE^u consensus base (-38, Fig. 4A). An FDPS1 promoter mutant derivative was generated through altering four of the IIARE consensus bases (Fig 4A). The wild type FDPS1 promoter and IIARE-defective derivative were then cloned into pGL3-basic, followed by reporter assay. The results show that mutations of the IIARE consensus sequence within the FDPS1 promoter significantly reduced expression of the reporter gene (Fig. 4F). However, we note that inhibition of promoter activity in the mIIARE derivative can be partially alleviated by co-expression of Gal4-VP16 (Fig 4F). This effect is likely due to TFIIA interaction with activator (29) providing an alternative mechanism for TFIIA recruitment. Taken together, these data suggest that the IIARE plays a positive regulatory role in both basal and activated transcription at promoters containing a TATA box.

The IIARE is a negative element in AdML-IIARE promoter derivatives that lack a TATA box — Several previous studies have showed that core promoter elements can work alone or in combination with other elements in transcription regulation (4), and that the function of core promoter elements depends on the context of the promoter (9). Indeed, TFIIA can stabilize TBP binding to the TATA box (39, 40). We therefore asked whether the role of the IIARE in gene transcription is dependent on the TATA box. To answer the question, AdML derivatives that lack a TATA-box, but contain an optimal IIARE (AdML-IIARE-mTATA) or a defective IIARE (AdML-mIIARE-mTATA), were synthesized (Fig 5A) and cloned into pGL3-basic. *In vitro* transcription assays were performed with these

vectors. Intriguingly, the promoter of AdML-mIIARE-mTATA showed a significant increase in transcription activity in the absence or presence of Gal4-VP16 when compared with the AdML-IIARE-mTATA promoter (Fig. 5B). This suggests that the IIARE plays an inhibitory role in the absence of a TATA box. To confirm this observation, transient transfection assays were performed with HEK293T cells with the promoter derivatives. The data show that the AdML-mIIARE-mTATA promoter was significantly more active than the AdML-IIARE-mTATA promoter in both the absence and presence of Gal4-VP16 (Fig. 5C, D). To exclude the possibility that mutation of the TATA-box might solely account for this observation, luciferase assays were performed using the reporter vector driven by either the AdML-IIARE-TATA or AdML-IIARE-mTATA promoters. The data confirm that mutation of the TATA-box significantly reduced the activity of the reporter gene (Fig. 5E, F), suggesting that the TATA-box mutation does not contribute to the increased activity caused by mutation of IIARE.

We next determined whether the IIARE functions as a negative element in a natural promoter that lacks a TATA box. To this end, two natural TATA-less promoters, from the PCDHB17 and C1QTNF7 genes, were identified by searching the human promoter database. Core promoter DNA fragments of the PCDHB17 and C1QTNF7 genes and their derivatives containing a mutant IIARE were synthesized (Fig. 5A). They were cloned into pGL3-basic vector and then transfected into 293T cells. Unexpectedly, mutation of the IIARE significantly reduced expression of the reporter genes driven by the PCDHB17 or C1QTNF7 promoters (Fig. 5G, H), suggesting that the IIARE positively regulates transcription of the PCDHB17 and C1QTNF7 promoters. Unlike at the AdML derivatives that lack a functional TATA element,

these results are consistent with those from the promoters containing the TATA box. To understand how a defective TATA-box affects TFIIA binding to the promoter, protein-DNA assays were performed using promoter DNA fragments and recombinant proteins TFIIA and TBP. The data show that the promoters lacking the TATA box were still bound by TFIIA regardless of the presence of TBP (Fig. 5I, J). However, mutation of the IIARE did not affect TFIIA recruitment to the AdML derivative promoters that lack a functional TATA box (Fig. 5I). In contrast, mutation of the IIARE in the PCDHB17 promoter (which is naturally TATA-less) reduced TFIIA recruitment, (Fig. 5J). Taken together, these data imply that the IIARE regulates transcription activity in a promoter context-dependent manner.

The IIARE positively regulates transcription by increasing recruitment of pol II, TFIIA, TAF4 and p300 at the AdML-IIARE promoter—We have determined that the IIARE can play a positive and negative role in transcription at the AdML-IIARE and AdML-IIARE-mTATA promoters respectively (Fig. 3C, D, and Fig. 5B and C). Whether this observation can be reproduced *in vivo* remains unclear. We therefore generated stable 293 cell line derivatives using Flp-In system as shown in Fig. 6A. The Flp-In 293 stable cell lines were used for luciferase assays in both the absence and presence of Gal4-VP16. The results show that mutation of the IIARE significantly reduced the activity of the reporter gene driven by the AdML-mIIARE promoter compared to the AdML-IIARE promoter (Fig. 6B, C). In contrast, compared to the AdML-IIARE-mTATA derivative, we observed an increase in activity of the reporter gene driven by the AdML-mIIARE-mTATA promoter (Fig. 6D, E). These results are consistent with those from the assays *in vitro* (Fig. 3, Fig. 5B and C).

To understand how the IIARE regulates transcription of the AdML-IIARE or

AdML-mIIARE promoters, ChIP assays were performed by using the Flp-In 293 stable cell lines and the antibodies indicated in Fig. 6F. As shown in Fig. 6F and G, the occupancy of TFIIA significantly decreased at the promoter of AdML-mIIARE compared with that at the promoter of AdML-IIARE. This is consistent with the results from protein-DNA binding assays *in vitro* (Fig. 3B). Strikingly, the occupancies for pol II, TAF4 and p300 at the promoter of AdML-mIIARE decreased 2-3 fold respectively compared with recruitment at the AdML-IIARE promoter, suggesting that the IIARE enhances recruitment of these factors to the promoter of AdML-IIARE. However, the occupancies for TBP, TFIIB, TAF1 and BTAF showed little difference between the AdML-IIARE and AdML-mIIARE promoters.

To understand how the IIARE affects transcription of the ADML promoter derivative that lack a TATA box, ChIP assays were performed with the stable cell lines integrated with these promoters. Fig. 6H shows that the occupancy of TFIIA showed little change between the promoters of AdML-IIARE-mTATA and AdML-mIIARE-mTATA, which is consistent with the observation for protein-DNA binding assay (Fig. 5I). However, the occupancies of pol II or TAF1 at the AdML-IIARE-mTATA promoter were significantly lower than that at the AdML-mIIARE-mTATA promoter (Fig. 6H, I), suggesting that the IIARE inhibits the transcription activity by reducing the recruitment of pol II and TAF1 at the AdML-IIARE-mTATA promoter. The occupancies of TFIIB, TBP, TAF4, P300 did not show a significant difference between the promoters except BTAF1 recruitment was slightly reduced at the mutant IIARE derivative. The data show that TBP was recruited at both of the promoters *in vivo* even though they contained a mutant TATA box. TBP can bind other general

factors and cofactors such as TFIIB and TAFs, which likely provides an alternative means of recruitment independent on the TATA element (1). Taken together, the data in Fig. 6 suggest that the IIARE enhances transcription of the AdML promoter by increasing the recruitment of pol II, TFIIA, TAF4 and p300 at the promoter, but inhibits transcription of the AdML promoter that lacks a TATA element by reducing the recruitment of pol II and TAF1 at the promoter.

Discussion

Previous studies revealed that TFIIA makes direct contacts with up to 9 base pairs upstream of the TATA box (42, 47) and 2 base pairs downstream of the TATA box (42). Crystal structure analyses of TFIIA-TBP-DNA using yeast TBP and TFIIA core domains showed that the β -barrel of TOA 1 interacts with 3-5 bases upstream of the TATA box, but not with the bases downstream of the TATA box (39, 44). In this study, we have confirmed that TFIIA makes direct contacts with the DNA sequence upstream of the TATA box (Fig. 1D, E), in particular with the two G bases immediately upstream of the TATA box on the positive strand of the AdML promoter (Fig. 1E, F). This result is in agreement with the previous findings (43) and suggests the presence of TFIIA recognition region upstream of the TATA box at the AdML promoter. Using the information, we then defined a TFIIA recognition element (IIARE) upstream of the AdML TATA box through SELEX and DNA sequencing. This element displays strong preference for G and T, and against A and C at its two ends (Fig. 2C). In this regard the IIARE consensus sequence is similar the BRE^d where TFIIB also favours G or T, although the specific sequence of consensus bases is distinct (9). The IIARE is distinct from the BRE^u (8), in which TFIIB prefers the bases G and C. Another feature of the IIARE is that the central region (-38 to -36)

displays less preference to G or T than the flanking bases (-41 to -39 and -35 to -32) (Fig. 2C). Previous photocrosslinking studies showed that TFIIA β mainly cross-linked to the sequence immediately upstream of the TATA box, whereas TFIIA α was mostly cross-linked with the sequence further upstream of TFIIA β -cross-linked region. This spacing between the TFIIA α and TFIIA β contact regions (42) could potentially contribute to the difference in base preference within the IIARE (Fig. 2C).

We found mutation of the IIARE severely affected TFIIA binding to promoter DNA and reduced promoter activity regardless in both the absence and presence of the transcriptional activator Gal4-VP16 (Fig. 3, Fig. 4 and Fig. 5G, J). This coincided with reduced binding of TFIIA to the promoter DNA. However, when artificial AdML promoter derivatives that lack a TATA box were tested for the same experiments, we observed a negative function for the IIARE (Fig. 5B-D). Thus, our results suggest that the IIARE modulates transcription in promoter context-dependent manner. Previous studies showed that both BRE^u and BRE^d can also positively or negatively regulate the activity of core promoter (9, 48). The BRE^d regulates transcription activity in a promoter context-dependent manner (9); whereas the BRE^u plays different roles in basal and activator-dependent transcription (48). These observations suggest that the role of core promoter elements in transcription relies on specific conditions, either in the context of the core promoter or the interplay of gene-specific transcriptional regulators. Indeed, previous *in vivo* studies suggest that the function of a core promoter element depends on the specific physiological environment (13).

ChIP assays showed that mutation of the IIARE significantly reduced the recruitment of pol II, TFIIA, TAF4 and p300 at the AdML promoter

(Fig. 6F, G). Others have reported that the transcriptional coactivator p300 promotes the formation of a TFIIA-TBP-DNA complex through acetylation of TFIIA (49). In addition, TFIIA binds to TAF4 to assist PIC assembly at core promoter (33). Thus, IIARE-dependent increase in the occupancy of TFIIA could subsequently enhance the recruitment of TAF4 and p300 at the promoter with an optimal IIARE. TAF1 and BTAF1 have been shown to compete with TFIIA to bind the TBP-DNA complex and inhibit transcription (50, 51). However, the recruitment of TAF1 and BTAF1 was not affected by mutation of the IIARE within the AdML promoter that contains a functional TATA box (Fig. 6F, G). Surprisingly, mutation of the IIARE within the ADML promoter containing a defective TATA element increased the recruitment of both pol II and TAF1 (Fig. 6H, I). It is possible that mutation of the IIARE combined with defective TBP-TATA interactions increases the potential of TAF1 to bind the Initiator element. In addition, TAF1 can play dual roles in transcriptional regulation through phosphorylation of histones to activate transcription (1, 52). It is therefore possible that TFIIA-IIARE contacts modulate the recruitment of TAF1 in the absence of TBP-TATA interactions.

Our ChIP assays revealed that the subunits of TFIID differentially associated with either the ARE-containing promoter or ARE-defective promoter. Previous studies suggest that TAFs can form multiple complexes, some of which do not contain TBP (1). Moreover, protein-coding genes show distinct requirements for individual TAFs (53), and TAFs are differentially recruited at the promoters of many active genes as stem cell differentiation progresses (54). Therefore, it is plausible that TFIIA-IIARE interactions can influence the TAF-dependency of the core promoter and thus their requirement at individual genes.

In this study we identified a core promoter element upstream of the TATA box that is recognized by TFIIA. The IIARE modulates transcription in a promoter context-dependent manner. The IIARE enhances transcription by increasing the recruitment of pol II, TFIIA, TAF4 and p300 at the AdML promoter. The IIARE can acts as a negative element at the AdML promoter when the TATA element is rendered inactive. In this context the IIARE inhibits the association of TAF1 and pol II with the AdML-IIARE. These findings not only extend our understanding of the role of TFIIA in gene transcription, but also provide new insights into the regulatory mechanism of core promoter elements in transcription by pol II.

Experimental procedures

Plasmids, proteins and site-directed mutagenesis—Human cDNAs for the TFIIA subunits (α/β and γ) and TBP were cloned into the plasmid pET30a. The cDNA for HA-tagged TFIIA α/β was cloned into pcDNA 3.1(+). The promoter DNA fragments for AdML, hCYP1A2, FDPS1 PCDHB17, C1QTNF7 and their derivatives were initially cloned into the pGEM3 plasmid. A DNA fragment comprised of nine Gal4 DNA recognition sites was inserted immediately upstream of the promoter derivatives; the nine Gal4 DNA-binding sites along with promoter DNA fragment was then transferred to the reporter vector pGL3-basic. The cDNA encoding Gal4-VP16 was cloned into pET30a and pcDNA 3.1 (+). Recombinant proteins for TFIIA α/β , TFIIA γ , TBP and Gal4-VP16 were expressed with *E. coli* BL21 (DE3; Agilent) and purified with Nickel agarose as described in the manufacturer's manual (Qiagen). Reconstituted TFIIA was obtained through denaturing the recombinant proteins TFIIA α/β and TFIIA γ with protein denature buffer (50 mM Tris-Cl (pH7.4), 1 mM EDTA, 100 mM NaCl, 0.1 mM PMSF and 7

M Urea) for 4 hours, and then renaturing the proteins by dialysis (20 mM HEPES (pH7.6), 1 mM EDTA, 100 mM NaCl, 0.1 mM PMSF, 10% Glycerol and 0-7 M Urea); the concentration of urea in dialyzing buffer was decreased at 1M/5 hours from 7M to 0M during the dialysis. Native TFIIA was purified from HEK293T nuclear extract as described previously (55). All chemicals were purchased from Sigma-Aldrich.

Bandshift assays and methylation interference—

One microgram of core promoter DNA for wild type AdML-BRE promoter and its derivatives (AdML-mBRE^{ud},

AdML-mBRE^d and AdML-mBRE^u) was labeled with 2.5 μ Ci of [γ -³²P]-ATP (GE Healthcare) and 2.5 U of T4 polynucleotide kinase (Promega). Bandshift assays were performed with 0.1 pmol of radio-labelled AdML core promoter DNA, 5 or 10 μ g of native human TFIIA (hTFIIA) protein and 10 μ g of recombinant human TBP (hTBP) protein as described previously (45). Methylation interference was performed using 1 pmol of radio-labelled AdML core promoter DNA, 50 μ g of native human TFIIA protein and 50 μ g of recombinant hTBP protein as described previously (9).

Immobilized protein-DNA binding assays and Western blot

—Overlapping oligonucleotides for wild type AdML core promoter (AdML-WT) and its derivatives or wild type hCYP1A2 p, PCDHB17 and their derivatives were synthesized and modified with biotin at the 5' end of the positive strands (TianyiHuiyuan Co.). One microliter of 25M forward and reverse oligonucleotides were used for overlapping PCR in 50 μ L reaction mixture. The PCR product was purified by the PCR clean kit (Axygen) and immobilized with Streptavidin magnetic beads (Thermo) in 100 μ L 1*PBS buffer at 4 °C overnight. The beads were then washed with 1*PBS buffer 3 times and collected with a magnetic stand

(Promega). The DNA-immobilized beads were incubated with HEK293T nuclear extract or reconstituted hTFIIA and recombinant hTBP at 4°C for 4 hours. The beads were washed with buffer D (20 mM Hepes, 100 mM KCl, 0.2 mM EDTA, 1mM DTT, 0.2 mM PMSF, 20% Glycerol) 3 times, DNA-bound proteins were eluted with 1*SDS loading buffer and the samples analyzed by Western blot. The antibodies for Western blot were purchased from Abcam (TFIIA: ab50821, TBP: ab51841 and β -Tubulin: ab6046) and Santa Cruz Biotech (Gal4: SC-577).

Random selection—The oligonucleotides for the AdML derivative used in random selection was synthesized as follows:

5'-CGTGACCGGGTGTTCCTNNNNNNNNNT
ATAAAAGGGGGTGGGGGCGCGTTCGTCCTC
ACTCTCTCCGCATCGCTGT-3'. This DNA
fragment contains ten consecutive randomized
nucleotides immediately upstream of the TATA
box, the randomized region was synthesized with
equimolar of nucleotides (Taiyihuiyuan Co.). The
primers used for PCR and cloning were as follows:
RSF:

5'-CGCGGATCCCGTGACCGGGTGTTCCT-3';

RSR:

CGCGAATTCACAGCGCATGCGAATTCAT
G-3'. PCR primers, RSF and RSR, contain
restriction sites recognized by BamH I and EcoR I
respectively to assist DNA fragment cloning.
Random Selection (SELEX) was performed as
described previously (9). Briefly, in the first cycle,
0.1 μ L of 25 μ M AdML template and 1 μ L of 25
 μ M primers (RSF and RSR respectively) were
used for PCR in 50 μ L reaction mixture containing
200 μ M of dNTP, 1 μ L of [α -³²P]- dATP (2.5 μ Ci)
and 5 U of Taq DNA polymerase (Roche). The
radioactive PCR product was purified on a 5%
native polyacrylamide gel. Bandshift assays were
performed using 0.2 pmol of purified DNA
fragment, 10 μ g of native hTFIIA and 10 μ g of

recombinant hTBP as described previously (45).
The DNA was recovered from the shifted band and
amplified by PCR within 50 μ L reaction mixture as
described above. The amplified radioactive DNA
was used for the second round of selection. After
seven rounds of selection in total, the recovered
DNA fragments were amplified by PCR without
the addition of radioactive isotope, followed by
digestion with BamH I and EcoR I and cloned into
pGEM3. Fifty clones in total were screened and
used for sequencing of the promoter region. The
frequency of occurrence for each base over the
randomized region (from -41 to -32) was counted
and the consensus bases were determined by the
frequency of occurrence.

In vitro transcription assays—*In vitro* transcription
assays were performed using the non-radioactive
method as described in our recent work (46). In
this method, pGL3-basic reporter gene vector
driven by promoter or promoter derivative was
used for DNA template. One hundred nanograms
of DNA template were used for RNA synthesis in
the reaction mixture (25 μ L Hela nuclear extract, 3
 μ L 100 mM MgCl₂, 3 μ L 10 mM NTP) in the
absence or presence of 1 μ g of Gal4-VP16. When
the reaction finished, the samples were subject to
DNA template depletion, hybridization and primer
extension, transcript was detected by using
RT-qPCR and our uniquely designed primers (46).
Relative transcriptional activity was analyzed by
Bio-Rad CFX Manager 3.0 software.

Cell culture, transfection and luciferase assay—
HEK293T cells were cultured in 12-well plates
with high-glucose DMEM complete medium.
Transient transfection for HEK293T cells or Flp-In
293 stable cell lines was performed with Turbofect
reagent (Thermo). Luciferase assays were
performed with the dual-light detection system
(Thermo). Briefly, HEK293T cells were
co-transfected with 1.5 μ g of reporter vector driven
by wild type promoter or its IIARE derivatives, 1

µg of vector expressing β-galactosidase and 0.5 µg of vector expressing Gal4-VP16 or empty vector pcDNA3.1 (+). 36 hours post-transfection, the cells were harvested and disrupted with the lysis buffer provided by the manufacturer (Thermo). After centrifugation at 1200 rpm for 5 minutes, 3 µL of the cell lysate were used to detect luciferase activity with the dual-light detection system. Luciferase activity was normalized by the activity of β-galactosidase within the same sample. Relative luciferase activity was obtained by comparing luciferase activity between wild type promoter and its IIARE-defect promoter, in which the basal activity of wild type promoter was arbitrarily set as 1.

Generation of Flp-In 293 stable cell lines and ChIP assay—Flp-In 293 stable cell lines were generated with the Flp-In system (Invitrogen). A Flp-In 293 host cell line that has been integrated with the fragment of FRT-LacZ-Zeocin was provided by Taylor lab (The University of Manchester). The DNA fragments for AdML-IIARE-Luc, AdML-mIIARE-Luc, AdML-IIARE-mTATA-Luc and AdML-mIIARE-mTATA-Luc were cloned into the vector pcDNA5/FRT respectively. The vector pcDNA5/FRT bearing a promoter-driven reporter gene (pcDNA5/FRT-pro-luc) and the vector expressing a recombinase (pOG44) were co-transfected into 293 host cell line with

lipofectamine 2000 (Thermo). 48 hours post-transfection, selection was initiated by adding hygromycin to the cells at the final concentration of 200 µg/mL. The cell line from a single colony was obtained by diluting the cells into 96-well plates. The positive cell lines were determined by detecting the expression of luciferase gene.

Chromatin immunoprecipitation (ChIP) assays for pol II and transcription factors or cofactors were performed using Flp-In 293 stable cell line that was integrated with promoter-driven luciferase gene as described previously (45). The ChIP samples were detected by qPCR using Bio-Rad Real Time Detection System and analyzed by Bio-Rad CFX manager 3.0. The relative occupancy was obtained by comparing the enrichment of promoter DNA from the ChIP sample of each factor with that from Input sample, 1 ng of genomic DNA was used to perform qPCR for input, which is equivalent to 0.01% of the sample used for the ChIP assay of individual factor. Fold change was obtained by comparing the relative occupancy of a factor between the promoters with or without ARE. The antibodies for ChIP assays were purchased from Santa Cruz Biotech(pol II: SC-21751, TBP: SC-204, TFIIB: SC-225, TAF1: SC-735, TAF4: SC-136093, BTAf1: SC-8139 and p300: SC-584) and Abcam (TFIIA: Ab50821).

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Conflict of interest

The authors declare that they have no conflict of interest with the contents of the article.

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Author contribution

J.W. and S.Z. performed *in vitro* transcription assays and reporter assays, western blot, ChIP assays and data analysis. W.H. cloned natural promoters and derivatives. Y.W. expressed and purified TFIIA and derivatives. H.P. assisted protein-DNA binding assay. P.S. provided facilities and consumables for performing some of experiments such as bandshift assays. Y.Z. cloned the AdML-IIARE promoter and its derivatives. S.G.E.R. designed and supported a part of experiments and revised the manuscript. W.D. designed most of experiments, performed bandshift assays, random selection, protein-DNA binding assays and data analysis, gained the fund and wrote the manuscript.

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Footnotes

TFIIA: transcription factor IIA; NC2: negative cofactor 2; TFIID: transcription factor IID; Gal4-VP16: an recombinant activator consist of Gal4 DNA-binding domain and herpes virus protein 16 activation domain; TBP: TATA-binding protein; TAF1: TBP-associated factor 1; BTAF1: B-TFIID TBP –associated protein 1; TAF4: TBP-associated factor 4, HMGB1: high mobility group box 1 protein; IIARE: TFIIA recognition element; BRE^u: TFIIB recognition element upstream of the TATA box, BRE^d: TFIIB recognition element downstream of the TATA box, AdML: adenovirus major late promoter; AdML-BRE^{ud}: adenovirus major late promoter containing TFIIB recognition elements at upstream and downstream of the TATA box; AdML-BRE^u: Adenovirus major late promoter containing a TFIIB recognition elements at upstream of the TATA box; AdML-IIARE: adenovirus major late promoter containing a TFIIA recognition element, ADML-mIIARE: adenovirus major late promoter containing a mutated TFIIA recognition element; AdML-IIARE-mTATA: adenovirus major late promoter containing a TFIIA recognition element

and a mutated TATA box; AdML-mIIARE-mTATA: adenovirus major late promoter containing a mutated TFIIA recognition element and a mutated TATA box; hCYP1A2: human cytochrome p450 1A2; FDPS1: farnesyl diphosphate synthase 1; ChIP: chromatin immunoprecipitation. PCDHB17: Protocadherin beta 17; C1QTNF7: C1q and tumour necrosis factor –related protein 7.

Tables

Table 1. The prevalence of IIARE in human natural promoters

Number of IIARE consensus bases	≥ 5	≥ 6	≥ 7	≥ 8
Percentage \pm SD in TATA-containing natural promoters	50.3 ± 0.095	26.5 ± 0.062	9.96 ± 0.023	2.4 ± 0.014
Percentage \pm SD in TATA-less natural promoters	58.4 ± 0.134	28.7 ± 0.095	15.5 ± 0.041	7.9 ± 0.021

Fig1

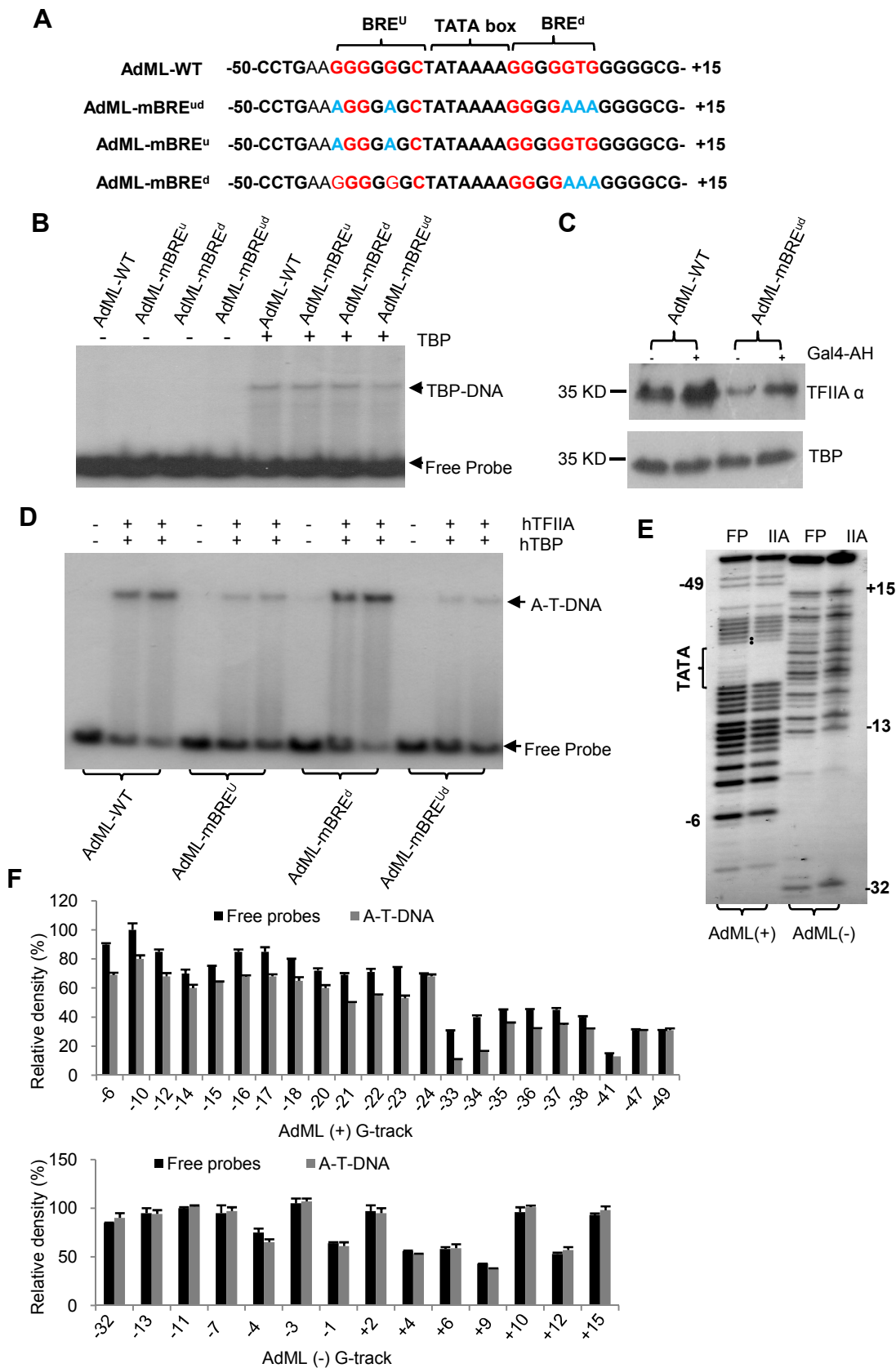


Fig. 1 TFIIA makes direct contacts with the sequence upstream of the TATA box at the AdML promoter. A, The DNA sequences for wild type AdML and AdML derivatives showing the TATA box (bold), BRE consensus bases (red) and BRE mutated bases (light blue). B, Mutations of the BRE^u (or BRE^d) consensus bases in AdML promoter did not affect the formation of TBP-DNA complex. Banshift assays were performed with 200ng TBP and 1μCi radio-labeled wild type AdML or AdML derivatives and detected by autoradiography. C, Mutations of the BRE^{ud} consensus bases in AdML promoter reduced TFIIA recruitment to promoter DNA but did not reduce TBP binding. Protein-DNA binding assays were performed using DNA-immobilized streptavidin magnetic beads and HEK293T nuclear extract, followed by incubation, washing and elution. The eluted samples were detected by Western blotting with the antibodies against TFIIA and TBP respectively. Equivalent volume of elution (10μL) was loaded in each lane for Western blotting . GAL4-AH: an activator comprised of a GAL4 DNA-binding domain and an acidic peptide (alpha helix). D, Mutations of the BRE^u consensus bases in AdML promoter decreased the formation of TFIIA-TBP-DNA complex. EMSA was performed using recombinant hTBP, native hTFIIA and radioactive wild type AdML or AdML derivatives and detected by autoradiography. A-T-DNA: TFIIA-TBP-DNA complex; the sequence for AdML wild type and its derivatives are indicated as in A. E, Methylation interference assays showing TFIIA contacts with the bases immediately upstream of the TATA box. The TATA box (left bracket) and TFIIA contacting sites (solid black dots) are indicated in AdML positive strand. FP: free probe, IIA: TFIIA. F, Relative density of G tracks for both AdML promoter positive and negative strands in the methylation interference assays performed in E. The relative density for the G-tracks of both free probe and the DNA from TFIIA-TBP-DNA complex was analyzed by Image J.

Fig 2

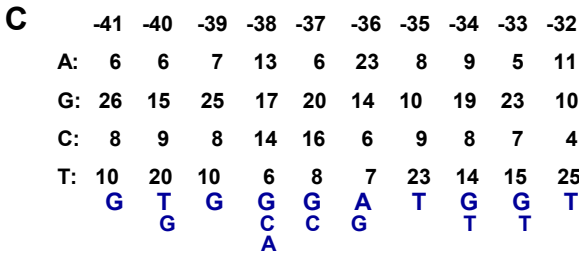
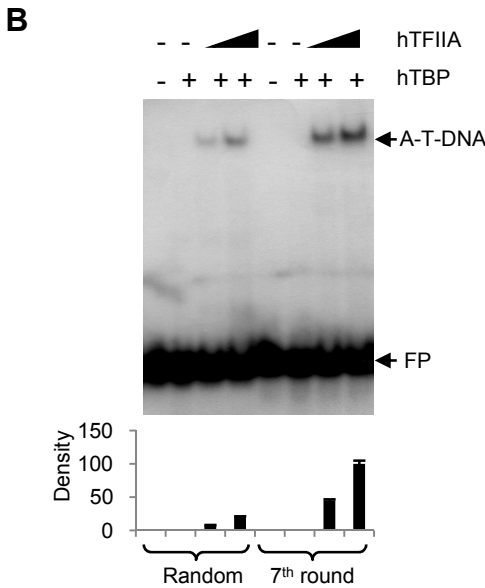
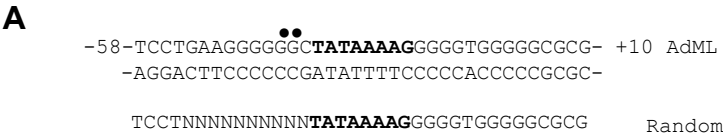


Fig. 2 A TFIIA recognition element upstream of the TATA box derived from random selection and DNA sequencing. A, The sequences for wild type AdML and randomized AdML positive strand. TFIIA contacting bases in the AdML promoter are marked with solid black dots and the TATA box is shown in bold character on the positive strand. The randomized bases (10 consecutive Ns) are located upstream of the TATA box (bold). B, Bandshift assay showing increased enrichment of the TFIIA-TBP-DNA complex after seven rounds of selection compared with the original randomized DNA pool (Random). EMSA was performed with 100 ng recombinant hTBP and native hTFIIA (50 or 100 ng) and detected by autoradiography. Relative density for EMSA is presented (bottom panel). FP: Free probe, A-T-DNA: TFIIA-TBP-DNA complex. C, The frequency of base occurrence at each position over randomized region (-41 nt to -32 nt) and the consensus bases of TFIIA recognition region (below). 50 clones in total were screened and sequenced after seven rounds of selection. The consensus bases are determined by the frequency of occurrence (>25%) at each position. The size of letter reflects the frequency of occurrence for each base.

A

AdML-WT -58-CCGGGTGTTCTCTGAAGGGGGGCT**TATAAAA**GGGGGTGGGGGCGCGTTC-+10

AdML-IIARE -58-CCGGGTGTTCTCT**GTGGGATGGT****TATAAAA**GGGGGTGGGGGCGCGTTC-+10

AdML-mIIARE -58-CCGGGTGTTCTCT**CCCTTCCCCCT****TATAAAA**GGGGGTGGGGGCGCGTTC-+10

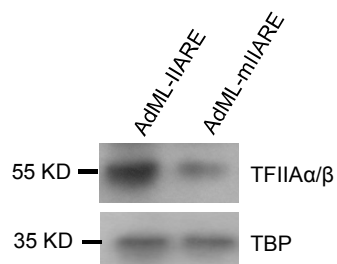
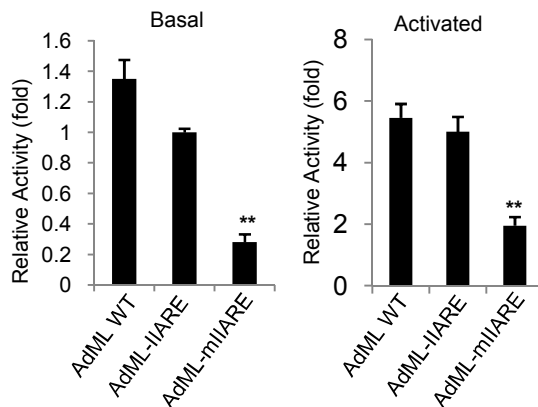
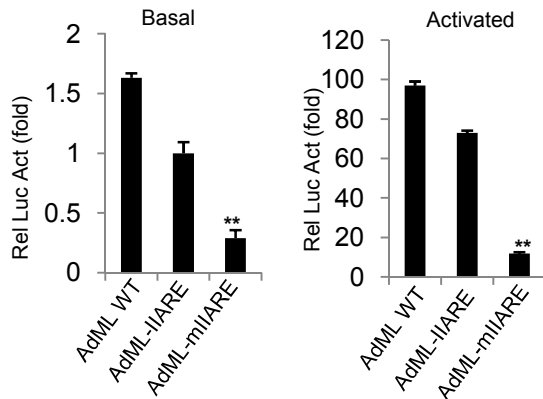
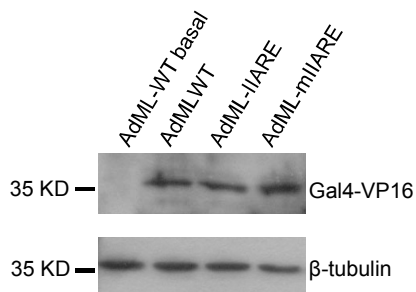
B**C****D****E**

Fig. 3 Mutation of the IIARE in the ADML promoter reduces TFIIA binding to DNA and promoter activity. A, The DNA sequences for the wild type AdML core promoter and its derivatives showing the TATA box (bold), IIARE consensus bases (blue) and mutated IIARE consensus bases (red). B, Mutation of the IIARE upstream of the TATA box reduces TFIIA binding to the promoter DNA. Protein-DNA binding assays were performed with recombinant hTFIIA, hTBP and promoter DNA-immobilized streptavidin magnetic beads. The eluted samples were analyzed by Western blotting with the indicated antibodies. Equivalent volume of elution (10 μ L) were loaded in each lane for Western blotting. C, *In vitro* transcription assays showing the relative activity for the promoters of AdML-WT, AdML-IIARE and AdML-mIIARE in the absence (left) or presence (right) of GAL4-VP16. Relative activity was obtained by comparing the basal activity of the AdML-IIARE promoter with that of the AdML-WT promoter or that of AdML derivative promoters in the absence or presence of Gal4-VP16, where the basal activity for the AdML-IIARE promoter was arbitrarily set as 1. D, Luciferase assays showing the effect of IIARE mutation on reporter activity. Relative luciferase activity was obtained by comparing the basal activity of the AdML-IIARE promoter with that of the AdML-WT promoter or AdML derivative promoters with or without the co-expression of Gal4-VP16. The basal activity for the AdML-IIARE promoter was arbitrarily set as 1. E, Immunoblotting of the samples used in D. Western blot was performed using antibodies against Gal4 or β -tubulin. Equivalent amounts of protein (10 μ g) were loaded in each lane for Western blotting. Each *column* in C and D represents the mean \pm SE of three independent experiments. Significant difference was analyzed by comparing the activity of the IIARE-containing promoter with that of the IIARE-defect promoter. * represents $P < 0.05$, ** represents $P < 0.01$, P value was obtained by student's *t* test.

Fig 4

A

hCYP1A2-WT -58-AATCTGATAGG**GGGCGGTGTT**TATAAAAAGGCCACTCACCTA- +10
hCYP1A2-miARE -58-AATCTGATAGG**CCCTTC**CCCTATAAAAAGGCCACTCACCTA- +10
FDPS1-WT -58-CATTGGGAGTT**GTAGTCTAAT**TATATATTTCCGCCCTTAGTGT- +10
FDPS1-miARE -58-CATTGGGAGTT**AAAGTCAAA**TATATATTTCCGCCCTTAGTGT- +10

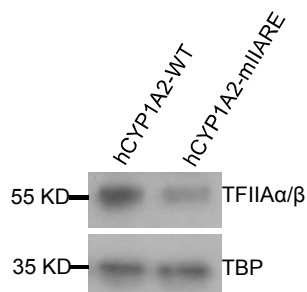
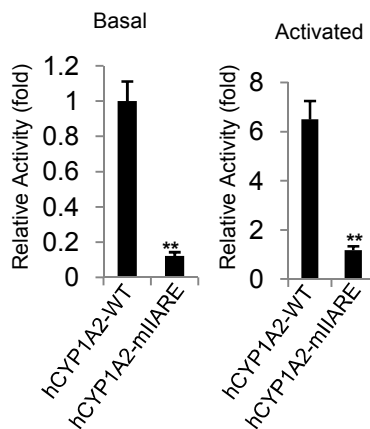
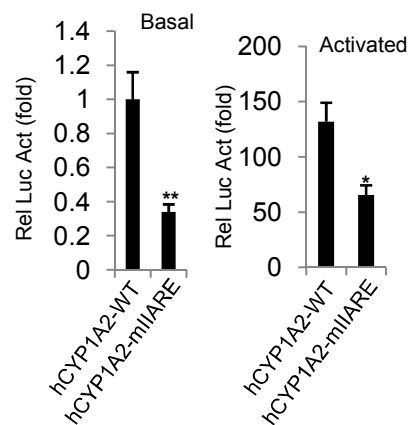
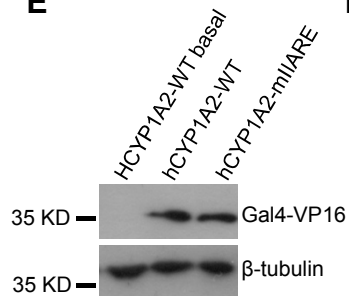
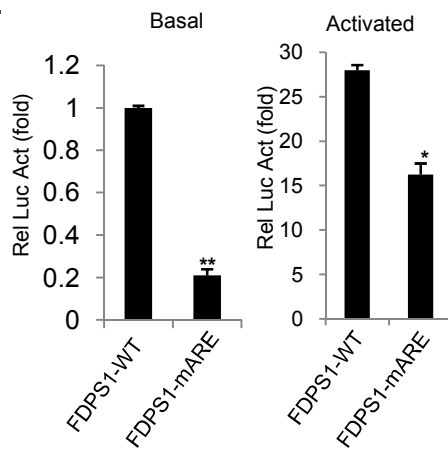
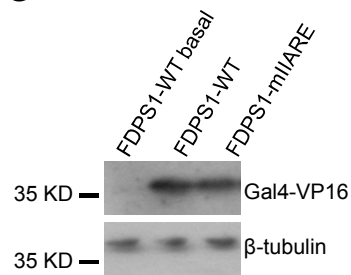
B**C****D****E****F****G**

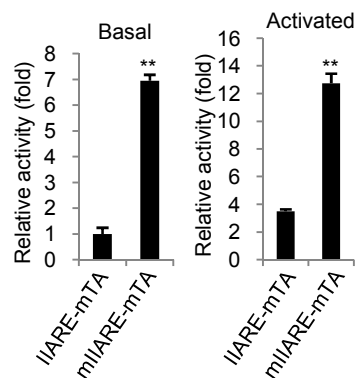
Fig. 4 Mutation of the IIARE in natural promoters reduces TFIIA binding to DNA and promoter activity. A, The DNA sequences for natural core promoters and their derivatives showing the TATA box (bold), the IIARE consensus bases (blue) and the mutated IIARE consensus bases (red). B, Mutation of the IIARE reduces TFIIA binding to the hCYP1A2 promoter. C, *In vitro* transcription assays showing the relative activity for the promoters of hCYP1A2-WT and hCYPEA2-mIIARE in the presence or absence of GAL4-VP16.. D, Luciferase assays showing the effect of mutation of the IIARE on the expression of a reporter gene driven by the human CYP1A2 gene promoter or its derivatives. E, Immunoblotting of the samples to confirm the co-expression of Gal4-VP16 used in C. F, Luciferase assays showing the effect of IIARE mutation on the expression of a reporter gene driven by the human FDPS1 gene promoter. G, Immunoblotting of the samples with the co-expression of Gal4-VP16 used in E. Protein-DNA binding assays, *in vitro* transcription assays, luciferase assays and Western blotting were performed as in Fig 3. Each *column* in C, D and F represents the mean \pm SE of three independent experiments. Significant difference was analyzed by comparing the activity of the IIARE-containing promoter with that of the IIARE-defect promoter. * represents $P < 0.05$, ** represents $P < 0.01$, P value was obtained by student's *t* test.

Fig 5

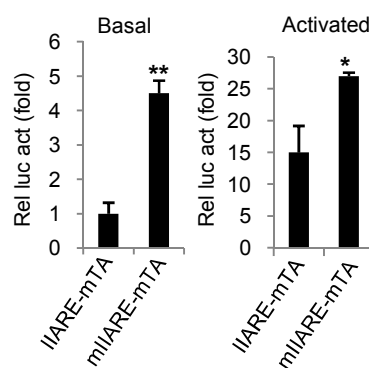
A

IIARE-mTA -58-CCGGGTGTTCTCTG**TGGGATGGT****CGATCCG**GGGGGTGGGGGCGCGTTC-+10
mIIARE-mTA -58-CCGGGTGTTCTCTC**CCTTCCCCC****GATCCG**GGGGGTGGGGGCGCGTTC-+10
PCDHB17-IIARE -58-TCTCCTTAAAAGCT**GTGCGG****TTTTCTGGC**AGCTCCAAAAGGAAACACT-+ 10
PCDHB17-mIIARE -58-TCTCCTTAAAAGC**AAAGCGGG****AAATCTGGC**AGCTCCAAAAGGAAACACT-+ 10
C1QTNF7-IIARE -58-TGGGAGAAGGAGT**GGGAGTTGT****TTAAACT**GAGAAAGACAGTTGCAAA -+10
C1QTNF7-mIIARE -58-TGGGAGAAGGAGT**AGAGG****AAATTA**AACTGAGAAAGACAGTTGCAAA--+10

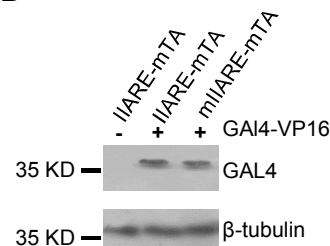
B



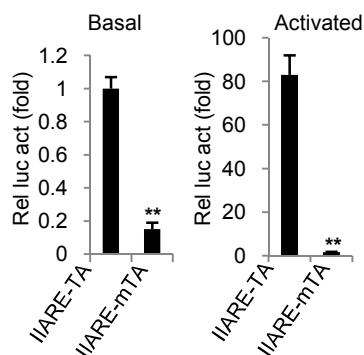
C



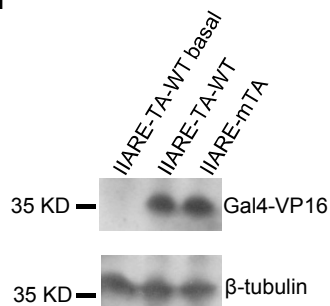
D



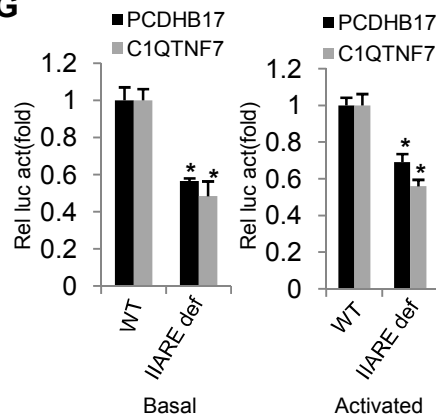
E



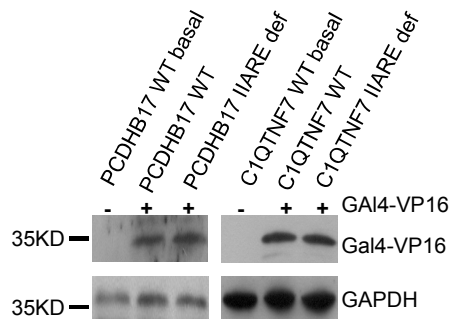
F



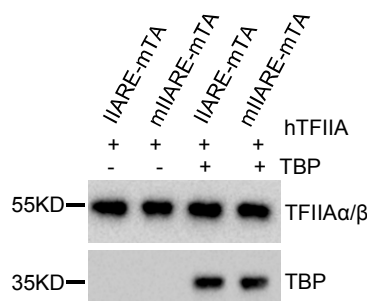
G



H



I



J

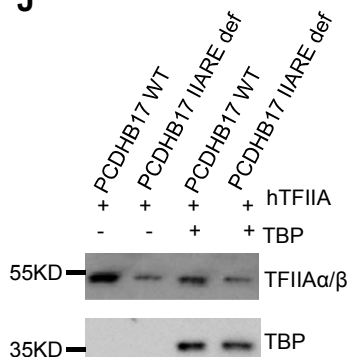


Fig. 5 The effect of IIARE mutation on the transcriptional activity of the AdML promoters that lack a TATA box. A, The DNA sequences for core promoters lacking a TATA box and their derivatives showing the IIARE consensus bases (blue), the mutated IIARE consensus bases (red) and the sequences around the TATA region (-31- -25; bold). B, Mutation of the IIARE significantly increased transcription activity *in vitro* of the AdML promoter derivatives lacking a TATA box. C, Mutation of the IIARE within the AdML-IIARE -mTATA promoter significantly increased the expression of the reporter gene. D, Immunoblotting analyses to conform the co-expression of Gal4-VP16 in the samples used in C. E, Luciferase assays showing the effect of mutations in the TATA-box on the expression of a reporter gene driven by the IIARE-containing AdML promoter. F, Western blotting of the samples to confirm co-expression of Gal4-VP16 in the samples used in E. G, Luciferase assays showing the effect of mutation of the IIARE on the activity of natural promoters that lack a the TATA box. H, Western blotting to confirm the co-expression of Gal4-VP16 in the samples used in G. I and J, Western blotting showing the effect of IIARE mutation on TFIIA binding to the promoters that lack of the TATA box. *In vitro* transcription assays, luciferase assays, protein-DNA binding assays and Western blotting were performed as in Fig 3. Abbreviation of the tested promoters is as follows: IIARE-mTA: AdML-IIARE-mTATA; mIIARE-mTA: AdML-mIIARE-mTATA; IIARE-TA: AdML-IIARE-mTATA. Each *column* in B, C, F and G represents the mean \pm SE of three independent experiments. Significant difference was analyzed by comparing the activity of the IIARE-containing promoter with that of the IIARE-defect promoter . * represents $P < 0.05$, ** represents $P < 0.01$, P value was obtained by student's *t* test.

Fig 6

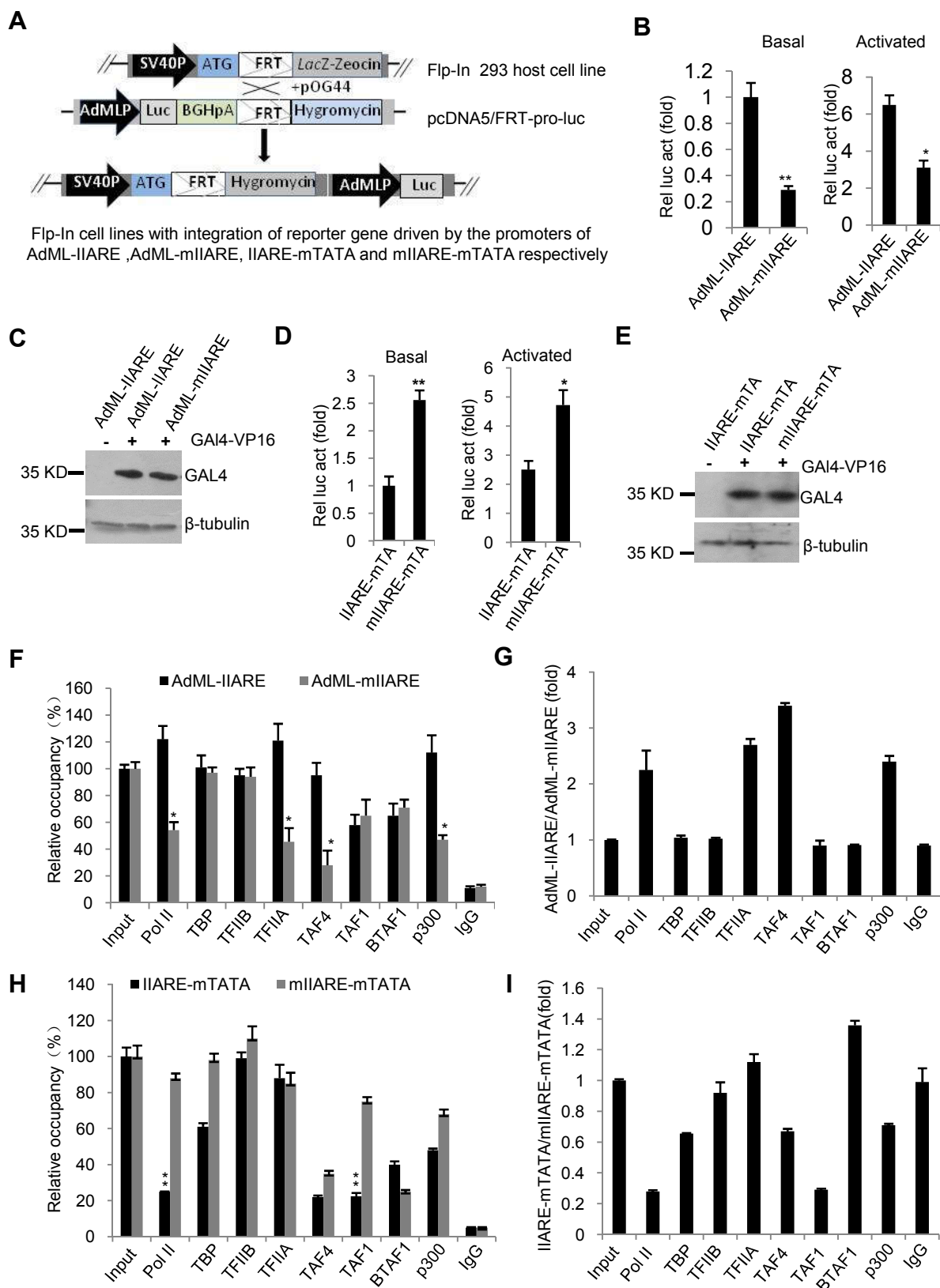


Fig. 6 The IIARE positively modulates transcription by increasing the recruitment of pol II, TFIIA, TAF4 and p300 at the promoter of AdML-IIARE. A, The scheme for the generation of Flp-In 293 stable cell lines that integrate promoter-directed luciferase gene derivatives. B, Mutation of the IIARE within the promoter of AdML-IIARE significantly reduced reporter gene expression *in vivo*. Flp-In 293 stable cell lines were co-transfected with pcDNA- β -Gal and pcDNA-Gal4-VP16 or control pcDNA3.1, the cell lysate was used to detect luciferase activity. Relative luciferase activity was obtained as in Fig. 3D. C, Immunoblotting analyses of the samples to confirm expression of Gal4-VP16 in samples used in part B. Immunoblotting was performed as in Fig 3E.. D, Luciferase assays showing the effect of IIARE mutation on reporter gene expression for the AdML derivatives that lack a the TATA box. Luciferase assays were performed as in B. E, Immunoblotting analyses for the samples to confirm expression of Gal4-VP16 used in D. Immunoblotting was performed as in Fig 3E. F, ChIP-qPCR assays showing the relative occupancy for the indicated factors at the promoters of AdML-IIARE or AdML-mIIARE. G, Comparison of the relative occupancy for indicated factors between the promoters of AdML-IIARE and AdML-mIIARE. H, ChIP-qPCR assays showing the relative occupancy for the indicated factors at the promoters of AdML-IIARE-mTATA or AdML-mIIARE-mTATA. I, Comparison of the relative occupancy for indicated factors at the promoters of AdML-IIARE-mTATA and AdML-mIIAR-mTATA. The relative occupancy was obtained by comparing the enrichment of promoter DNA from the ChIP sample for individual factor with that from input. 1 ng of genomic DNA was used to perform qPCR for input, which is equivalent to 0.01% of sample used for ChIP assay of individual factor. Each *column* in B, D, F and H represents the mean \pm SE of three independent experiments.* represents $P<0.05$, ** represents $P<0.01$, P value was obtained by student's *t* test.

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