

Chromatin affinity-precipitation using a small metabolic molecule: its application to analysis of *O*-acetyl-ADP-ribose

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Abstract In the cell, many small endogenous metabolic molecules are involved in distinct cellular functions such as modulation of chromatin structure and regulation of gene expression. *O*-acetyl-ADP-ribose (AAR) is a small metabolic molecule that is generated during NAD-dependent deacetylation by Sir2. Sir2 regulates gene expression, DNA repair, and genome stability. Here, we developed a novel chromatin affinity-precipitation (ChAP) method to detect the chromatin fragments at which small molecules interact with binding partners. We used this method to demonstrate that AAR associated with heterochromatin. Moreover, we applied the ChAP method to whole genome tiling array chips to compare the association of AAR and Sir2. We found that AAR and Sir2 displayed similar genomic binding patterns. Furthermore, we identified 312 potential association cluster regions of AAR. The ChAP assay may

therefore be a generally useful strategy to study the small molecule association with chromosomal regions. Our results further suggest that the small metabolic molecule AAR associates with silent chromatin regions in a Sir2-dependent manner and provide additional support for the role of AAR in assembly of silent chromatin.

Keywords Chromatin affinity-precipitation (ChAP) · *O*-acetyl-ADP-ribose (AAR, OAADPR) · Sir2 · Chromatin immunoprecipitation (ChIP) · Silent chromatin

Introduction

In addition to proteins, many noncoding RNA molecules and small metabolites, such as NAD, GTP, ADPR, NAM, and cAMP, may play important regulatory roles in the cell. Similar to the widely studied small noncoding RNA molecules, which are involved in the regulation of gene expression, small cellular metabolites and biosynthetic molecules, such as cAMP, tRNA^{Tyr}, FMN, and NAD, play important roles in modulation of gene expression in the cell [16, 25, 53]. These small regulatory molecules are involved in either changing the activity of transcription factors/cofactors [16, 55], the feedback loop of transcriptional RNA level [12, 32], or changes in chromatin structure [25, 30, 39]. Therefore, small metabolic molecules might be also involved in epigenetic processes. However, although some metabolites have been well studied and have well-known biological functions and mechanisms, many remain poorly characterized and the details of their molecular mechanisms, even their fundamental biological functions, still remain to be determined.

O-acetyl-ADP-ribose (OAADPR or AAR for simplicity) is a small metabolic molecule. The cellular AAR

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concentration has been reported to be around 560 nM [27]. AAR is generated during NAD-dependent deacetylation by silent information regulator 2 (Sir2) family proteins [4, 5, 30, 38, 44, 48]. Sir2 is involved in many distinct cellular functions, including the regulation of gene expression, genome stability, metabolism, and aging [4, 8, 13, 21, 26, 33, 37, 42, 44, 46, 49, 52, 53]. There are several other AAR-producing enzymes, such as Hst1, in the cell. Although AAR is produced by Sir2-like proteins from bacteria, yeast, and humans and is shown to block/delay oocyte maturation [5], bind to chromatin-related proteins [24], activate ion channels [15], and affect cellular reduction-oxidation status [51], its precise biological functions and molecular mechanisms are not fully known.

In eukaryotic cells, nuclear DNA is packaged into chromatin. Post-translational modifications of histones and association with histone-binding proteins organize the genome into distinct states that are referred to as heterochromatin and euchromatin [22, 33, 40]. The known heterochromatic DNA domains of *Saccharomyces cerevisiae* are the silent mating type loci (*HML* and *HMR*) and the telomeric DNA regions [33, 43]. In addition, silencing factors are involved in stabilization of the ribosomal DNA (rDNA) tandem repeats [6, 45]. The Sir proteins, Sir2, Sir3, and Sir4, are required for establishing and maintaining silent chromatin domains at telomeres and the mating type loci [1, 14, 23, 41]. These Sir proteins form a nucleosome binding complex called the SIR complex [34–36, 47]. AAR has been shown to promote a change in stoichiometry and structure of the SIR complex in vitro [30]. In addition, NAD-dependent deacetylation of nucleosome arrays purified from yeast, coupled to AAR synthesis, results in the formation of SIR-nucleosome filaments that require all three Sir proteins [39]. Because binding of the SIR complex to nucleosome arrays in these experiments is not sufficient for filament formation, we proposed that AAR induces a structural change that mediates filament formation. These results suggest that AAR binds to at least one of the Sir proteins and indicate that both deacetylation and AAR synthesis contribute to SIR complex assembly in vitro and that both events may be required for the formation of silent chromatin in vivo. In addition, AAR increases the affinity of the Sir3 protein for nucleosomes in in vitro binding assays [31]. However, it has been reported that the requirement for Sir2 and AAR can be partially bypassed in some transcriptional silencing assays [9, 54].

In this study, we developed a novel chromatin affinity-precipitation (ChAP) method, which allowed us to study the ability of immobilized small molecules to precipitate specific chromatin fragments. Using AAR as an example, the results demonstrated that AAR associated with silent heterochromatic regions in a Sir2-dependent manner.

Moreover, we used the ChAP method to compare the association of AAR and Sir2 via high resolution whole genome tiling array chips. We found that AAR displayed a similar genomic binding pattern to that of Sir2. We also identified 312 additional association cluster regions for AAR. These results indicate that AAR can associate with specific target(s) in silent chromatin and establish ChAP as a general method for probing the chromosomal targets of metabolites.

Materials and methods

Yeast strains

The yeast strains used for this study were SF10 (BJ459, *MATa ura3-52 trp1 lys2-801 leu2Δ1 pep4Δ::HIS3 prb1Δ1.6R can1*) and W303-1a (*MATa ade2-1 can1-100 his3-11,15 leu2-3,112 trp1-1 ura3-1*). Strain *sir2Δ::HIS3* (JRY3433) was previously described [34, 42].

Purification of proteins and ^{32}P -O-acetyl-ADP-ribose (AAR), gel electrophoresis, Western blotting, quantification, and HDAC fluorescent activity assay

Proteins were purified as previously described [30, 35]. ^{32}P -AAR was prepared by deacetylation of an acetylated histone H4 peptide using Hst2 and ^{32}P -NAD. Protein samples were separated on SDS-PAGE and stained by Coomassie brilliant blue R520 or silver nitrate to visualize protein bands. Western blotting was performed using either the ECL Western blot detection system (Amersham Biosciences) or the Western Lightning Chemiluminescence Reagent (PerkinElmer Life Sciences). AlphaEaseFc (Alpha Innotech) was used to quantify the density of protein or DNA bands on the gel and of small molecule spots on the TLC plate. HDAC fluorescent activity assay was performed following the manufacturer's instructions (Merck/Biovision kit) except a reaction buffer containing 1 mM NAD was used. Briefly, 85 μl of enzyme was mixed with 10 μl of HDAC assay reaction buffer and added to 5 μl of fluorescent substrate. After 30 min incubation at 37°C, the reaction was stopped by adding 10 μl of lysine developer and incubation at 37°C for 30 min. The signals were recorded by reading samples in a fluorescence plate reader (Ex/Em: 350–380/440–460 nm).

Deacetylation reactions, TLC, HPLC, and MALDI-TOF mass spectrometry

Approximately 2.5 mg GST-Sir2 was immobilized on glutathione-Sepharose 4B (Amersham Biosciences). The immobilized GST-Sir2 was incubated with 1 mM β -NAD $^{+}$

and 1.5 mM synthetic penta-acetylated histone H4 peptide (SGRGK^{Ac}GGK^{Ac}GLGK^{Ac}GGAK^{Ac}RHRK^{Ac}, Yao-Hong Biotechnology) in 1.75 ml of 50 mM HEPES-KOH (pH 7.0), 300 mM KCl, 1 mM Mg(OAc)₂ for 2 h at 30°C and then overnight at 4°C. Protein bound to the solid beads was packaged into a column, and the enzymatic reaction products were eluted and either directly tested by TLC or further analyzed by HPLC and MALDI-TOF mass spectrometry using standard procedures as previously described [30].

Dot blotting

Serial dilutions of protein samples were spotted on a PVDF membrane (50–200 µmol) and were incubated with ³²P-AAR at 4°C overnight. The membranes were then washed twice with 20 mM Hepes-KOH pH 8.0, 300 mM KCl, 1 mM MgCl₂, and 0.05% NP40, followed by two washes with 20 mM Hepes-KOH pH 8.0, 100 mM KCl, 1 mM MgCl₂, and 0.025% NP40 at room temperature. The membranes were then subjected to autoradiography and quantification by phosphorimaging using QuantityOne software (BioRad).

BIAcore surface plasmon resonance analysis

Real time protein-small molecule interactions were examined using a BIAcore 3000 or BIAcore 2000 instrument (BIAcore). Sir2 or BSA was individually immobilized on different flow cell of a CM5 sensor chip using an amine-coupling kit (BIAcore). The experimental procedures for interaction assays and data analysis were previously described [29, 30].

Affinity pull-down and chromatin affinity-precipitation (ChAP)

Activated Affigel 10 resin (BioRad) was washed with ice cold H₂O three times and equilibrated in cold buffer A (50 mM Hepes-KOH pH 8.0, 300 mM KCl, 1 mM MgCl₂, 0.05% NP40, 0.5% BSA, 0.5% casamino acids) three times. To immobilize a small molecule on the beads, 10 mM individual NAD, ADPR, NAM, AAR, ATP, ADP, or AMP was coupled to the beads at 4°C for 4–6 h, followed by washes with cold buffer B (50 mM Tris-HCl pH 7.0, 150 mM KCl, 1 mM MgCl₂, 0.05% NP40, 0.1% BSA, 0.5% casamino acid) three times. To block the remaining coupling sites, cold 1 M ethanol-amine-HCl pH 8.5 was incubated with beads for 1–2 h. The beads were then washed with cold buffer B three times.

For affinity pull-down assays, purified protein or whole cell lysate extract, containing 0.1% BSA and 0.1%

casamino acid, was added to each immobilized resin and incubated at 4°C overnight. Before elution with buffer C (4× SDS-PAGE Laemmli loading dye without Bromophenol blue), beads were washed five times with cold buffer B.

For ChAP, the treatment of cell lysate extract was as previously described for ChIP assays [39]. Each immobilized resin was incubated with cell lysate at 4°C overnight. Beads were then washed once with cell lysis buffer (50 mM Hepes-KOH pH 7.5, 300 mM NaCl, 1 mM EDTA, 0.5% Triton X-100, 0.1% sodium deoxycholate, 0.1% SDS, and protease inhibitors); twice with 50 mM Hepes-KOH pH 7.5, 250 mM NaCl, 1 mM EDTA, 0.25% Triton X-100, 0.1% sodium deoxycholate, 0.05% SDS, and protease inhibitors; twice with 10 mM Tris-HCl pH 8.0, 0.25 M LiCl, 0.5% NP40, 0.5% sodium deoxycholate, and 1 mM EDTA; twice with 10 mM Tris-HCl pH 8.0, 0.25 M LiCl, and 1 mM EDTA; and once with TE (10 mM Tris-HCl pH 8.0 and 1 mM EDTA pH 8.0) at room temperature. Beads were eluted by incubating with 100 µl of 50 mM Tris-HCl pH 8.0, 10 mM EDTA, and 1% SDS at 65°C for 15 min. Eluates were transferred to a fresh tube and pooled with a final bead wash of 150 µl TE with 0.67% SDS. For input DNA, 200 µl TE with 1% SDS was added to 50 µl cell lysate extract. To reverse the cross-links, all samples were incubated at 65°C overnight, followed by the addition of 250 µl TE, 5 µg glycogen, and 100 mg proteinase K, and incubation at 37°C for 2 h. After adding 55 µl of 4 M LiCl, the samples were extracted once with phenol/chloroform and once with chloroform. DNA was then precipitated by two volumes of ethanol, washed with 75% ethanol, air dried, and resuspended in 50 µl TE with RNase A (20 µg/ml for input DNA and 2 µg/ml for affinity-precipitated DNA) and incubated at 37°C for 1 h. Gel conditions, quantification of PCR amplified specific bands, and primer pairs—except for PCR1 of YBR079C: 5'TGAGCACGCTTCTGTCTTTC3', 5'GAAGGCTGAATTGG AAGCTG3' and PCR2 of YBR079C: 5'ACAGCGATTTTTCGATGTCC3', 5'GGTTTTTGTGCTGCCTCTC3'—were as previously described [20, 42].

Chromatin affinity-precipitation on chip (ChAP on chip)

The precipitated chromatin DNA fragments were obtained as described above. Then, following the NimbleGen standard protocol instructions, samples were processed from the steps of post IP, ligation-mediated PCR (LM-PCR), labelling with either cy5 or cy3, hybridization with *S. cerevisiae* whole genome tiling array chip (NimbleGen), to the step of scanning chip and analysis of data using the NimbleScan program and SignalMap software.

Results

Association of AAR with Sir2

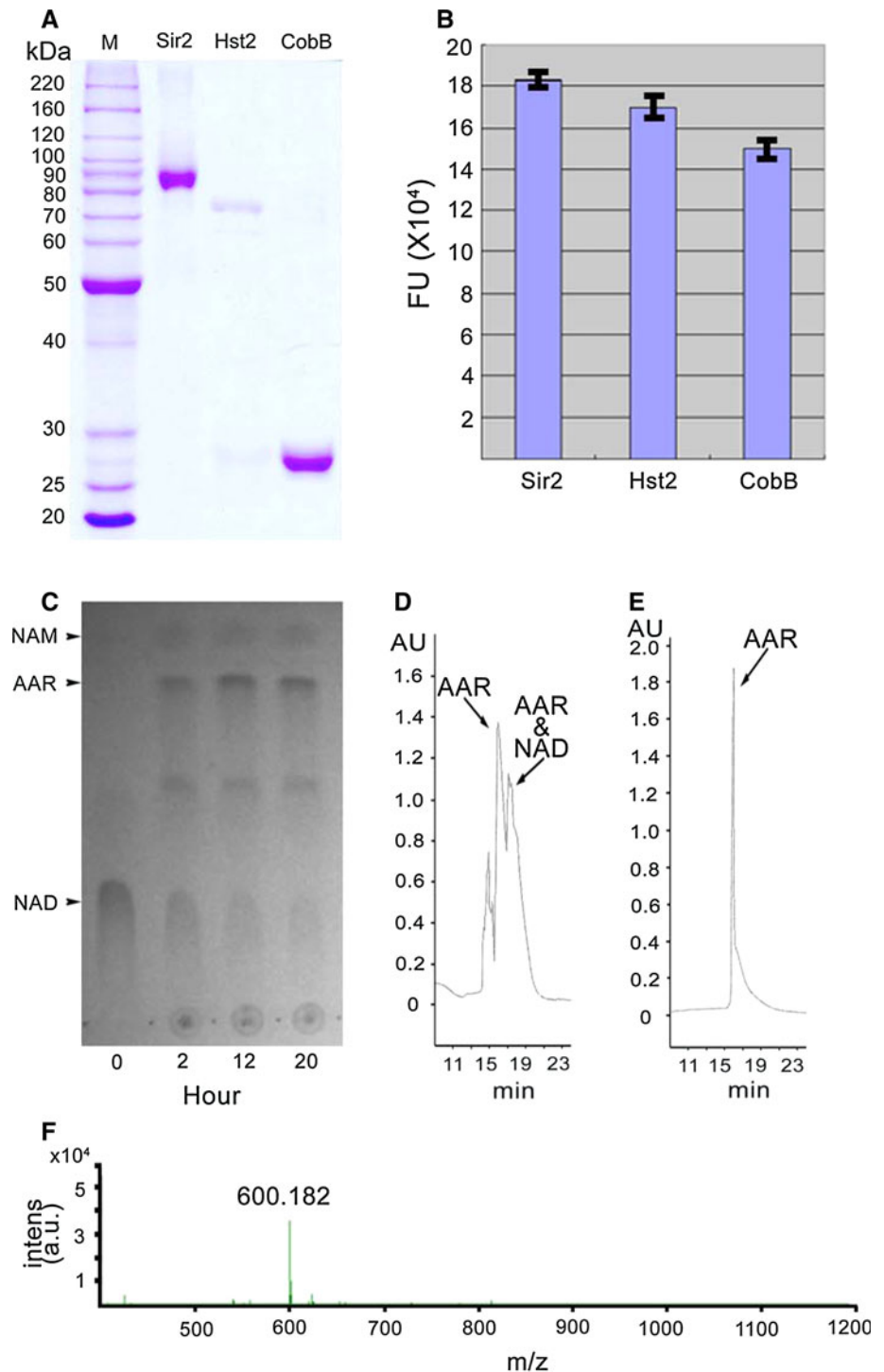
Based on the ability of Sir2 to generate AAR, a Sir2-AAR interaction is postulated. However, intriguing as this proposed interaction is, a definitive proof is lacking. Therefore,

we were interested to investigate the physical interaction between AAR and Sir2.

In order to obtain sufficient quantities of AAR, we optimized the condition of purification of AAR. We were able to produce AAR by enzymatic synthesis using either Sir2 or its homolog proteins, Hst2 and CobB (Fig. 1a). As shown in Fig. 1b, Sir2 was slightly more efficient in

Fig. 1 Purification of AAR.

a Coomassie-stained SDS polyacrylamide gel showing purified Sir2, Hst2, and CobB. **b** HDAC fluorescent activity assays showing the deacetylation activities of Sir2, Hst2, and CobB, respectively. **c** TLC separation of NAD-dependent deacetylation reaction products. The reaction times, reactant (NAD), and products (AAR and NAM) are indicated on the *bottom* and *left*, respectively. **d** Purification of NAD-dependent deacetylation reaction products on a C18 HPLC column. The peaks were collected and the positions of AAR and NAD as determined by MALDI-TOF mass spectrometry are indicated. **e** Several collected AAR peaks shown in **d** were combined to be re-purified by HPLC. **f** Mass spectrum showing the molecular mass of AAR from the peak shown in **e**



deacetylation of the substrate than Hst2 and CobB. Therefore, we chose Sir2 for the further large scale reactions. We also used thin layer chromatography (TLC) to monitor the results of the enzyme reaction over time. Under our conditions, after 2 and 12 h, around 54 and 85% of NAD was metabolized to AAR and NAM by Sir2, respectively. After 20 h reaction, there was still about 5% unconsumed NAD (Fig. 1c). As previously described [30], we purified the reaction products on a C18 HPLC column. Two peaks of interest, one containing AAR and the other containing AAR and NAD, were individually collected (Fig. 1d). The pure AAR peak collections were then combined, checked, and re-purified by HPLC again (Fig. 1e) and also confirmed by matrix assisted laser desorption/ionization-time of flight (MALDI-TOF) mass spectrometry (Fig. 1f).

We used a dot blotting assay to rapidly determine the physical binding of AAR to Sir2 that was immobilized on a PVDF membrane. We blotted increasing amounts of each protein (50–200 μ mol) onto membranes, which were then incubated with 32 P-labeled AAR. As shown in Fig. 2a, consistent with the Hst2-AAR crystal structure [57], we detected binding of AAR to Hst2, even for the lowest Hst2 concentration (50 μ mol) on the membrane. In the control, we observed no binding to BSA and C-terminal fragment of Sir3 at the same concentration range (data not shown). AAR also associated with Sir2 with an apparently higher efficiency than with Hst2 (Fig. 2a).

For studying the specificity of small molecule interaction with protein, we used an affinity pull-down approach. Several small molecules such as NAD, ADP-ribose (ADPR), nicotinamide (NAM), ATP, ADP, AMP, and AAR were immobilized on solid resins, which were then used to examine the specificity of affinity pull-down. Sir2 and Hst2 associated with AAR, as well as NAD, but bound little or no ATP, ADP, or AMP (Fig. 2b, c and data not shown). Furthermore, not only yeast Sir2 and Hst2 but human SirT1 were also able to undergo affinity pull-down by AAR but not by AMP from whole cell lysate extract (Fig. S1). Nevertheless, Sir2 associated with ADPR with a similar efficiency to that observed for AAR. Sir2 associated with NAM with lower efficiency. Both association of Hst2 with ADPR and with NAM showed a lower efficiency than Hst2-AAR association.

To measure the real time binding of small molecule interaction with Sir2, we performed BIAcore surface plasmon resonance (SPR) assays. Sir2 and control BSA were individually immobilized to a CM5 sensor chip, and small molecules were then used as the analyzers. Binding was observed between Sir2 and AAR but not between Sir2 and ADP or Sir2 and AMP (Fig. 2d and data not shown). The on and off rates for the association of AAR, ADPR, NAD, ADP, and AMP with Sir2 are summarized in

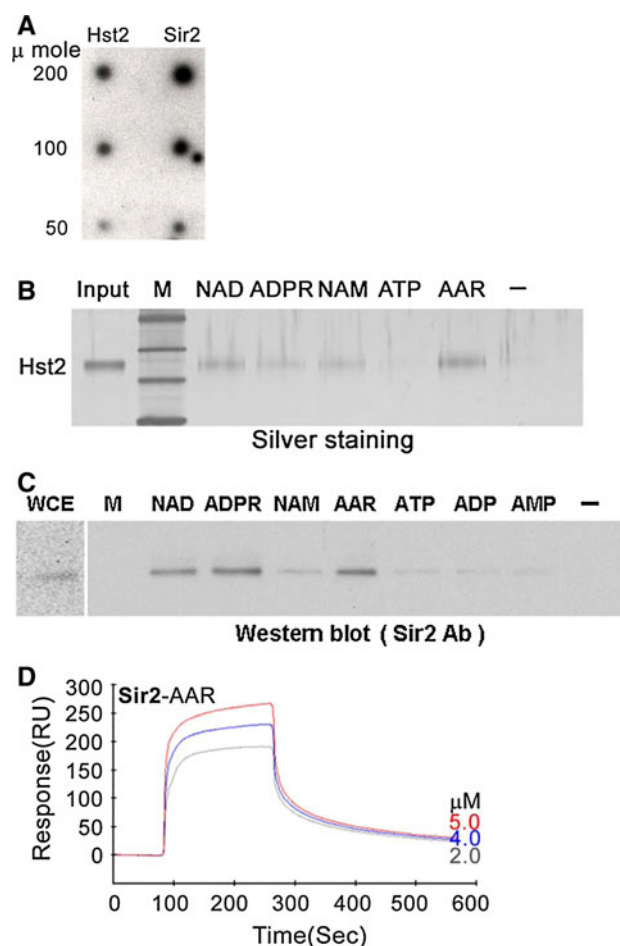


Fig. 2 Association of small molecules with Hst2 and Sir2. **a** Dot blotting assays showing the binding of 32 P-AAR to the indicated amounts of Hst2 and Sir2 immobilized on a PVDF membrane. **b** and **c** Affinity pull-down assays showing the binding of Hst2 and Sir2 to the indicated small molecules, which were immobilized on beads. **d** BIAcore surface plasmon resonance (SPR) experiments showing the association of AAR with immobilized Sir2. The concentrations of AAR are indicated on the sensorgram (right). *M* Protein standard marker, *WCE* whole cell extract, – bead only control

Table 1. Affinity calculation yielded K_D values of ~ 300 nM for the Sir2-AAR interactions. The K_D value for the association of NAD with Sir2 was ~ 580 nM. However, ADPR bound to Sir2 with lower affinity (K_D of ~ 3 μ M) (Table 1).

Affinity precipitation of heterochromatin fragments by AAR

Chromatin immunoprecipitation (ChIP) [19] has allowed significant progress in the study of in vivo protein-DNA interactions. We modified the typical ChIP protocol to assess the ability of small molecules such as AAR, NAD, and ATP to associate with sheared chromatin fragments. Briefly, this method, referred to as chromatin affinity-

Table 1 Association of AAR, ADPR, NAD, ADP, and AMP with immobilized Sir2

Interaction	ka (1/Ms)	kd (1/s)	KD (M)
Sir2-AAR	$(1.23 \pm 0.02) \times 10^3$	$(3.74 \pm 0.35) \times 10^{-4}$	$(3.04 \pm 0.26) \times 10^{-7}$
Sir2-ADPR	$(8.17 \pm 0.06) \times 10^2$	$(2.41 \pm 0.12) \times 10^{-3}$	$(2.95 \pm 0.16) \times 10^{-6}$
Sir2-NAD	$(1.70 \pm 0.45) \times 10^4$	$(9.18 \pm 0.91) \times 10^{-3}$	$(5.75 \pm 1.33) \times 10^{-7}$
Sir2-ADP	NM	NM	NM
Sir2-AMP	NM	NM	NM

NM Not measurable

precipitation (ChAP) involves the immobilization of small molecules to an Affigel resin via primary amine cross-linking. This resin is then used for affinity-precipitation and analysis of chromatin fragments as is the case with the standard ChIP assay.

Under our experimental conditions, as shown in Fig. 3a and b, using actin as a control, we observed enrichment signals of silent chromatin fragments, HMR-E, HML-E, HMR-a, HML- α , NTS1/5S, and NTS2/18S, using the AAR resins compared to resins that contained either ATP or other small molecules (about 1.5- to 2- vs. 1-fold enrichment over input). Interestingly, for NAD, the enrichment signals were detected on HMR-E, NTS1/5S, and NTS2/18S but not on HML-E, HMR-a, and HML- α . In contrast, for ADPR, the enrichment signals were detected on HMR-E, HML-E, HML- α , NTS1/5S, and NTS2/18S but not on HMR-a. AAR-dependent DNA precipitation required chromatin, as no enrichment was observed using naked DNA fragments (Fig. 3c). These results indicated that we were able to specifically precipitate silent heterochromatin fragments using the distinct immobilized small molecules such as AAR.

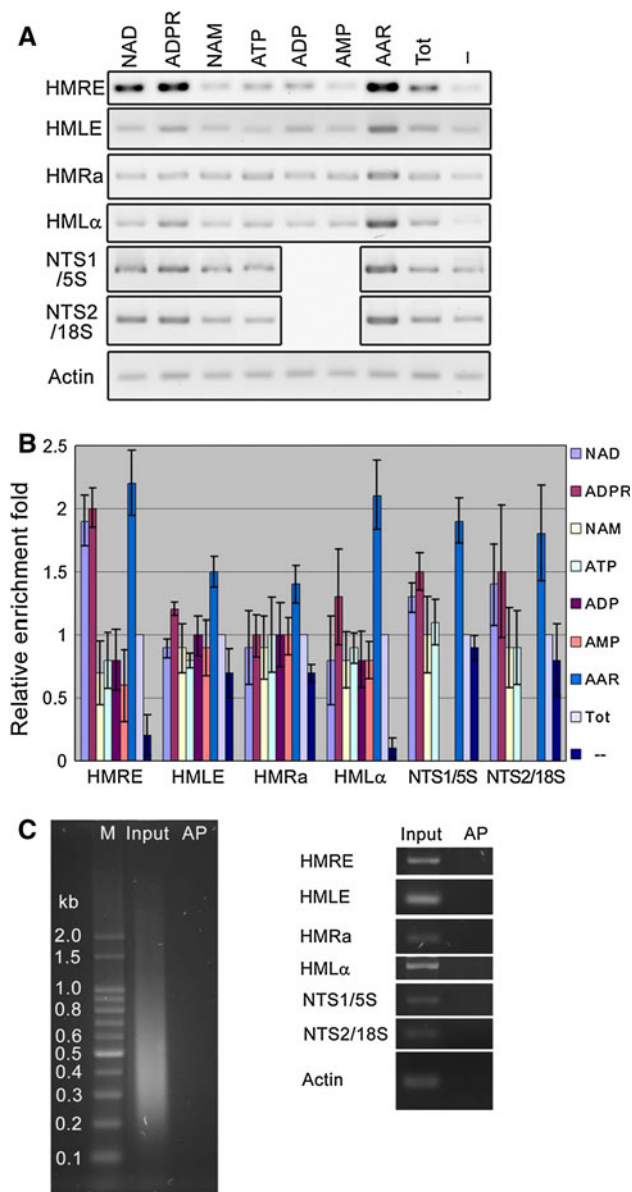
Previous studies have shown that deletion of any of the *sir* genes results in a disruption of heterochromatin [1, 18, 41]. To determine whether the association of AAR with chromatin required heterochromatin formation, we performed ChAP assays in wild type cells and cells that carried a deletion of the *sir2* gene. As our results suggest that Sir2 is an AAR binding partner, AAR should not be able to affinity-precipitate silent chromatin fragments in *sir2* deletion cells. As shown in Fig. 4, we observed an enrichment of four different silent chromatin fragments, the HMR-E and HML-E silencers, the HMR-a, and HML- α regions in ChAP from two different *sir2*⁺ strains, but these enrichments were not detected in *sir2* deletion cells. No significant difference in the signal enrichment of the above fragments was observed between *sir2*⁺ and *sir2* deletion cells in ChAP assays with AMP beads (Fig. 4). Although the enrichment of silent chromatin fragments in AAR pull-downs appeared to be weak (about 1.5-fold enrichment over input), the dependence on Sir2 suggested specificity in this assay. The association of AAR with silent chromatin fragments was consistent with our binding results presented in Fig. 2.

Genome-wide localization of AAR and Sir2

We next used the ChAP and ChIP assays in combination with whole genome tiling array chips to determine the genome-wide localization of AAR and Sir2. Genome-wide localization of Sir2 has been previously reported [28], but not using high resolution tiling arrays (“microarray of ~13 k unique spots at a resolution of 2 kb” vs. “tiling array of 385 kb probes with 50-mer probes at 32 bp median spacing”).

In general, AAR co-localized with Sir2 not only at all telomeric regions but also at almost all other chromosomal regions (Figs. 5 and S2). Interestingly, at several chromosome ends, including 1L, 1R, 2R, 3L, 3R, 4L, 6R, 9R, 10R, 11L, 11R, 13R, 14R, and 15L, both Sir2 and AAR display a narrower localization pattern (around only 1 kb from the terminus) compared to their distributions on other chromosomal ends (Fig. 5b). However, the localizations of Sir2 and AAR on each chromosome end ranged from ~1 to ~14 kb and, moreover, the broader distribution regions displayed a closely connected bi- or tri-peak distribution pattern (Figs. 5a, b and S2).

We performed further data analysis by calculating and filtering out peaks with false discovery rate (FDR) by using NimbleScan software. Other than the known silent chromatin regions, this analysis identified 312 potential AAR association clusters that spanned 481 genes, about 6.7% of yeast genes, and 40 intergenic regions (Table S1). We also used the PCR method to confirm some of these newly identified AAR association gene regions (Fig. S1), and the results were consistent with our ChAP on chip results. However, these newly identified cluster regions were mostly co-associated with Sir2 (Figs. 5 and S2). Interestingly, at some regions, we observed AAR association in the apparent absence of Sir2, particularly near the end parts of cluster distribution (e.g., Chromosome 2, ~397 k, Fig. 5c), suggesting that the association may be mediated by other AAR receptor proteins in these regions. This feature was consistent with the PCR results shown in Fig. 5d. We also confirmed this co-association feature by using *sir2* deletion cells and almost all enrichment signals of AAR and Sir2 disappeared, except a few weak enrichment signals on some regions, such as YIR019C (Figs. S2 and S3). However, compared to the telomeric DNA regions, in which the highest enrichment was over 30-fold



in Sir2 immunoprecipitations, these newly identified regions were usually enriched less than 8-fold (the cut-off of enrichment fold was artificially set at $2^{1.5}$ for the

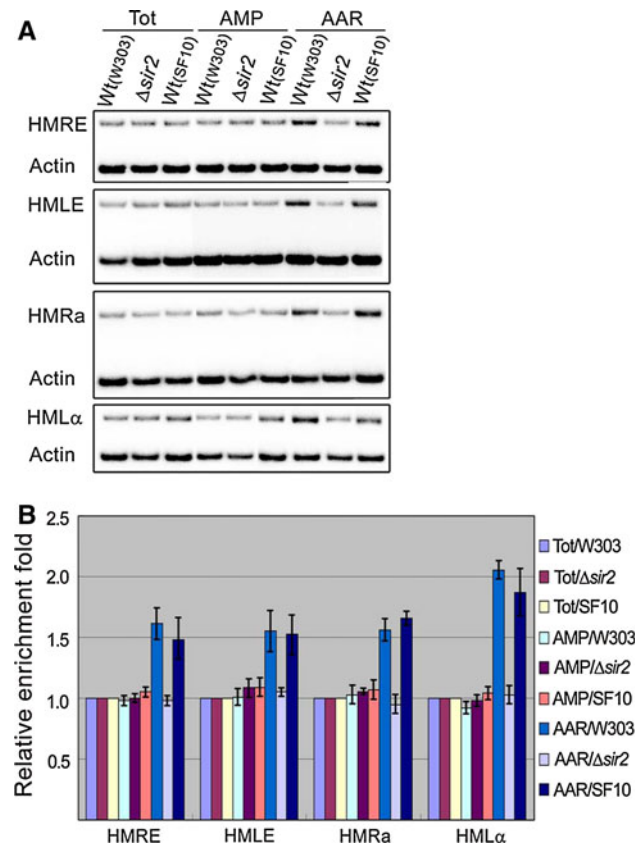
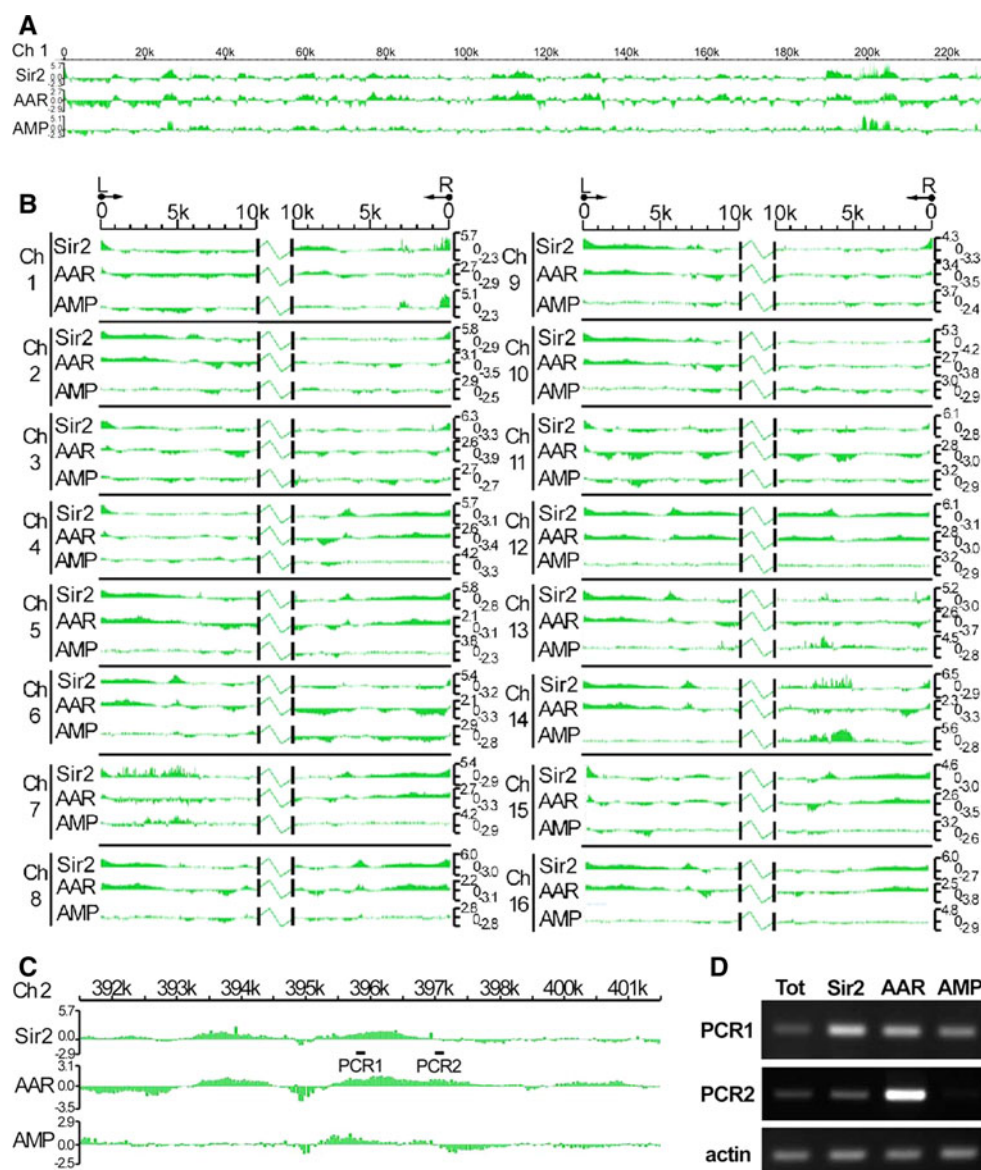


Fig. 4 Sir2-dependent AAR association with silent chromatin. **a** ChAP assays showing the association of AAR with silent chromatin fragments in *sir2* deleted cells as compared with *sir2*⁺ wild type cells. AMP was used as small molecule control, and actin primer was used as internal control and for normalization of signal. **b** Quantification of ChAP experiments showing the relative fold enrichments of AAR at silent chromatin fragments in *sir2*⁺ and *sir2* deleted cells. Averages and standard errors are shown for the results of three to four experiments. *Tot*, *Wt(W303)*, *Wt(SF10)*, and Δ *sir2* represent total cell lysate, two *sir2*⁺ (wild type), and *sir2* deleted strains, respectively

threshold value of AAR affinity-precipitation) (Figs. 5 and S2). As shown in supplemental Table S2, these AAR-associated regions were diverse and contained genes involved in many biological functions such as transferase activity, transcription regulation, protein binding, DNA binding, and signal transduction. At the same time, other ontological analyses of, for example, biological processes and cellular components also showed a similar diversity. However, it remains to be determined whether the Sir2 protein and AAR play any role in the regulation of these loci.

Our results showed that all chromosome ends were associated with Sir2 (Figs. 5a and S2), whereas previously little or no Sir2 enrichment was observed for telomeric regions of chromosomes 3R, 4L, 9R, and 16R [28]. However, most of the previously identified nonheterochromatic targets of Sir2 [28] were also enriched in our tiling array experiments (Fig. 5, Table S2).

Fig. 5 Genome-wide distribution of Sir2 and AAR. **a** Chromosomal display of association patterns of Sir2 and AAR on the whole chromosome 1. **b** Chromosomal display of association patterns on the telomere regions. For each chromosome, Sir2, AAR, and AMP data, respectively, are presented for the 10 kb region from the left (L) and right (R) chromosome end. **c** The different distributions of enrichment signals of Sir2 and AAR around position 397 k of chromosome 2. Chromosome number and the relative ratio of enrichment signal (\log_2) are indicated. **d** The different enrichment signals of Sir2 and AAR on the same region shown in **c**. The detected positions of PCR1 and PCR2 are also indicated in **c**. Actin primer was used as internal control and for normalization of signal. *Tot* represents total cell lysate



Discussion

Our results show that the ChAP method provided a general useful assay for studying the association of small molecules with chromosomal DNA fragments. Using ChAP assay, we have provided direct physical evidence for the interaction of a small metabolic molecule, AAR, with the Sir2.

Both AAR-Sir2 and AAR-Hst2 interactions are consistent with the known ability of these enzymes to use NAD as a cofactor during deacetylation and with the observation that NAM is an inhibitor of Sir2 [3]. Presumably, the binding of AAR to Sir2 and Hst2 reflects its affinity for the known NAD binding pocket in the active site of these enzymes [2, 56, 57]. However, our results do not rule out the possibility that AAR association with chromatin

fragments involves a Sir2-mediated heterochromatin formation.

Although AAR is produced by Sir2-like proteins from bacteria, yeast, and humans, its biological target(s) are not fully known. One of the nuclear sirtuins, Hst2, a Sir2 homolog in yeast, has been shown to co-crystallize with both AAR and a histone peptide [57]. A histone variant macroH2A1.1, mH2A1, has been identified as a target of ADP-ribose and/or AAR binding [24].

The phylogenetic conservation of Sir2-like enzymes and their reaction mechanism suggests that AAR may play similar regulatory roles in other systems. Sir2-like proteins have been implicated in several different regulatory pathways, and in mammalian systems, the SirT1 protein deacetylates p53, tubulin, and PCAF [4, 7, 10, 11]. AAR had been reported to directly bind to the cytoplasmic

domain of TRPM2 and to activate the TRPM2 channel [15]. Furthermore, a recent report showed that AAR binds to three glycolytic enzymes, GAPDH, PGK, and ADH, suggesting that AAR may be involved in the cytosolic regulation of glycolysis [51]. AAR, whose production is coupled to deacetylation of these and as yet to be identified substrates, may act as a signaling or second messenger molecule to regulate downstream events. In this regard, AAR may be similar to cyclic ADP-ribose (cADPR), which is a second messenger that regulates calcium signaling in mammalian cells [17].

On the whole genome tiling array chip, at several chromosome ends, such as 4L, 9R, and 11R (Fig. 5b), an absence of Sir2 and AAR broader spreading along the chromosomal end was shown. This might imply that Sir2 was just involved in a short distance (~ 1 kb) of spreading but not further longer spreading at these telomeres. Under our experimental conditions, although Sir2 and AAR displayed similar genomic distribution patterns, some regions only associated with either Sir2 or AAR (Figs. 5c and S1), suggesting that AAR might not associate with all Sir2 interaction regions. Perhaps, other chromosome-bound Sir2-like proteins, such as Hst1, mediate the association of AAR with these chromosome regions.

It has been reported that the requirement for Sir2 and AAR in silencing can be bypassed when Sir3 is fused with Hos3 as a chimera protein [9] or when Sir3 is overexpressed in cells containing histone H4 lysine to arginine mutations that mimic deacetylated H4 [54]. Our studies provide a starting point for further dissection of the AAR binding site, which is required in order to assess the importance of AAR-Sir binding in silent chromatin assembly in vivo. Nevertheless, in addition to AAR interaction with Sir2, we have also found that AAR binds to Sir3 (unpublished data), suggesting that multiple interaction sites of the SIR complex may contribute to AAR binding.

The genome-wide ChAP approach described here is useful not only for identification of chromosome-associated AAR targets but also for the identification of non-chromosomal targets of AAR. In the nucleus, the results of genome-wide screening show that AAR primarily co-precipitates with genomic fragments that are associated with Sir2 (Figs. 5 and S2). However, some AAR-associated regions did not contain Sir2 (Fig. 5c) and may be bound by other AAR target protein(s), such as other Sir2-like proteins or Sir3. To fully investigate the possible biological function(s) of AAR, the identification of its other targets, including both nuclear and cytoplasmic targets, is required. Affinity precipitation using a small molecule coupled to a solid resin was used successfully to identify the first histone deacetylase [50]. AAR coupled to a solid resin, combined with mass spectrometry identification of the

bound proteins, should be useful in identification of additional AAR targets.

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