

# Arsenite Inducible Regulatory Particle-Associate Protein

Subjects: **Biochemistry & Molecular Biology**

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Maintaining 26S proteasome activity under diverse physiological conditions is a fundamental requirement in order to maintain cellular proteostasis. Several quantitative and qualitative mechanisms have evolved to ensure that ubiquitin–proteasome system (UPS) substrates do not accumulate and lead to promiscuous protein–protein interactions that, in turn, lead to cellular malfunction.

AIRAP

PSMD14/RPN11

proteasome

protein misfolding

## 1. Introduction

Maintaining proteome functionality is a major challenge that the cell must resolve in order to survive the diverse conditions that it encounters. Changes in biophysical conditions and post-translational modifications all impact the correct folding of proteins, and cellular machineries have evolved to maintain proteome integrity. Promiscuous interactions of misfolded proteins are highly toxic to the cell and manifest in various pathologies [1]; therefore, restoring the proper configuration of misfolded proteins by chaperones, reducing enzymes and more constitutes a salvation pathway. In contrast, degradation pathways such as the ubiquitin–proteasome system (UPS) or the lysosomal pathway eliminate the toxicity of misfolded proteins by proteolytic degradation [2]. The 26S proteasome is the central module of the UPS to which all designated proteins, usually ubiquitinated with suitable configurations of ubiquitin, are delivered. The 26S proteasome is composed of a 20S catalytic particle and a 19S regulatory particle that enables recognition, deubiquitination and unfolding of a UPS substrate. In order to ensure the functionality of the basic 26S proteasome module, various modes of regulation exist in order to maintain functionality during acute and chronic conditions. We can artificially divide these modifications into several categories, including quantitative and qualitative changes in proteasome subunits (executed by transcriptional [3][4] [5] and translational [6] programs as well as post-translational modifications of proteasome subunits [7]) and changes in the composition of specific subunits, as exemplified in the case of the immunoproteasome (20Si) and the thymus proteasome (20St). In this mode of regulation, the exchange of catalytic subunits in response to acute signaling (IFN- $\gamma$ ) [8][9] or unique surroundings [10][11] enables modification of the proteasome to a more suitable peptide output. Another mode of regulation consists of alternating the entire cap composition. Alternative caps such as PA200 enable the engagement of the proteasome with specific substrates [12][13], while the PA28 (11S) enables proteasome modulation during specific conditions such as IFN- $\gamma$  [9][14][15]. The fourth mode of regulation that is diverse and continuously engaged with the proteasome involves shuttling factors. These accessory factors typically consist of a proteasome binding domain that usually consists of a ubiquitin-like (UBL) domain that binds one of the

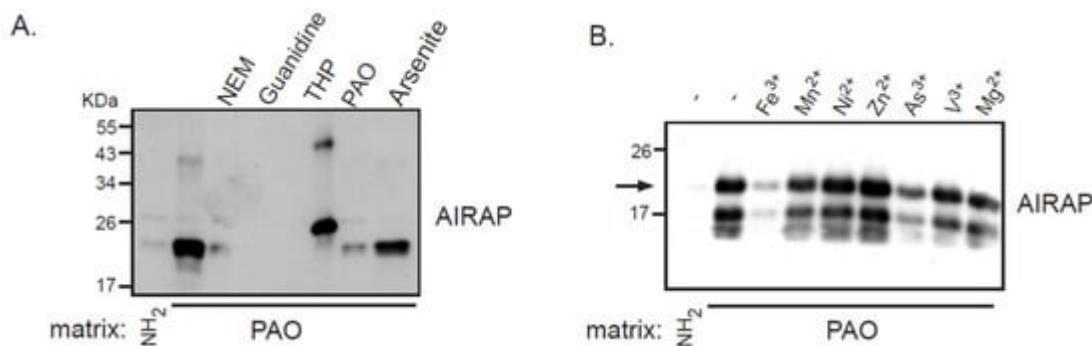
proteasome ubiquitin receptors and several ubiquitin binding domains (UBD) that account for ubiquitin chain affinity and specificity (for reviews, see [16][17][18]). The UBA–UBL mode of substrate delivery to the proteasome increases the diversity and flux of UPS substrates [19], induces proteasome activation [20] and supports the formation of liquid–liquid phase separation of proteasomes [21][22].

Originally identified as an arsenite inducible gene [23], AIRAP/Zfand2a was subsequently found to bind proteasomes with unique biochemical characteristics [24]. These include AIRAP–proteasome integrity in the absence of ATP and enhanced peptide hydrolysis. Unlike other proteasomal adaptors, AIRAP–proteasome binding is not obtained via UBA domains, nor does it bind ubiquitin. AIRAP knockout cells accumulate higher polyubiquitin levels in response to arsenite exposure (typical of UPS impairment) and proteasomes acquired from the knockout cells contain higher levels of polyubiquitinated substrates [24]. These findings rule out AIRAP's function as a shuttle factor but rather indicate it as a proteasome adaptor required for the efficient processing of UPS substrates by the proteasome upon arsenite-inflicted inhibition. In line with this notion, AIRAP expression was found to be induced upon proteasomal inhibition [25]; thus, AIRAP's role and gene expression programming point towards a mechanistic function in retaining proteasomal function upon inhibitory conditions, yet the mechanistic details of AIRAP function have not been revealed.

## 2. Recombinant AIRAP Binding to Arsenite

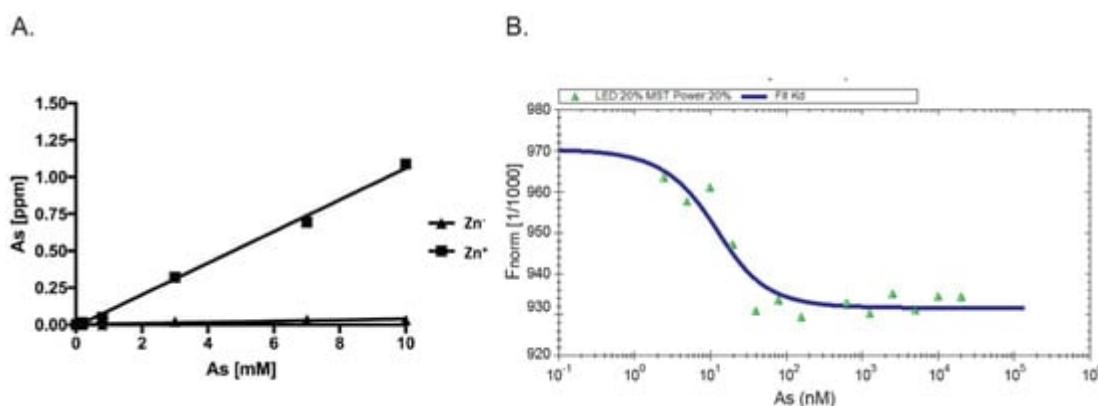
Arsenite-induced protein misfolding occurs through binding to vicinal cysteine residues [26]. AIRAP is a short polypeptide highly enriched with cysteine residues that coordinate through two AN1 type zinc fingers the binding of zinc residues. To evaluate the possibility of the direct binding of AIRAP to arsenite, we utilized an immobilized bivalent form of arsenite (phenylarsine oxide (PAO)) previously demonstrated to bind arsenite-binding proteins [27].

Recombinant AIRAP was incubated with PAO beads, unbound material was extensively washed, and AIRAP immunoblot was performed to evaluate the bound content (**Figure 1**). Several controls were performed to evaluate the nature of this binding, and the protein configuration was evaluated by preincubating the recombinant protein with guanidine. The modification of AIRAP cysteine content was evaluated by preincubation with *N*-ethylmaleimide (NEM) or the reducing agent THP. Finally, a trivalent arsenite ( $\text{As}^{3+}$ ) and a bivalent form of arsenite were used to evaluate their ability to outcompete the immobilized PAO binding to AIRAP. As evident from the results in **Figure 1A**, the binding of AIRAP was highly dependent on the protein's native structure (guanidine) and cysteine configuration (NEM and THP). Furthermore, AIRAP binding efficiently competed with a trivalent form of arsenite but not with a bivalent form (arsenite and PAO). To further investigate the specificity of AIRAP's ability to bind arsenite and the effect of different metals on AIRAP's ability to bind the PAO matrix, we incubated recombinant AIRAP with PAO matrix followed by incubation with various metals. As indicated in **Figure 1B**, metals such as  $\text{As}^{3+}$  and  $\text{Fe}^{3+}$  reduced AIRAP's binding to the PAO matrix (**Figure 1B**, lanes 3 and 7). These results imply a selective metal binding by AIRAP to the PAO matrix.



**Figure 1.** Recombinant AIRAP binding to a PAO matrix. **(A)** Recombinant mouse AIRAP was expressed as a C-terminus 6×His fusion protein and Ni-NTA affinity purified. Purified material was incubated with a control ( $\text{NH}_2$ ) or amino-phenylarsine (PAO) immobilized Sepharose 4B matrix in the presence of the cysteine alkylating *N*-ethylmaleamide (NEM), denaturing conditions (6M guanidine–HCl), reducing conditions (THP), sodium arsenite (As) or soluble amino-phenylarsine (PAO). Immobilized material was extensively washed, eluted by boiling with 2% SDS and subjected to SDS-PAGE and subsequent immunoblotting to evaluate AIRAP content. **(B)** The impact of metals on AIRAP binding to a PAO matrix was performed as in 1A by performing the binding in the presence of the indicated metals. The arrow indicates the full-length recombinant product of AIRAP.

As AIRAP is a zinc-binding protein, zinc may serve as a primary structural enhancer to enable the optimal positioning of cysteine residues to bind arsenite. This notion is further supported by our results with the PAO matrix (**Figure 1B**). To directly measure the protein's capacity to bind arsenite in a zinc-dependent manner, we measured arsenite binding to AIRAP by inductively coupled plasma optical emission spectroscopy (ICP-OES). Purified AIRAP was incubated with increasing concentrations of arsenite in the presence or absence of zinc, and arsenite content was evaluated. As seen in **Figure 2A**, AIRAP binding to arsenite was greatly enhanced when first incubated with zinc (marked  $\text{Zn}^+$ ). These results coincided with the indicated increase in AIRAP binding to the PAO matrix upon zinc incubation (**Figure 1B**). Analytical evaluation of this interaction was performed to determine the  $K_d$  value of AIRAP to arsenite using a micro-scale thermophoresis (MST) methodology [28]. A  $K_d$  of  $7 \text{ nM} \pm 0.87 \text{ nM}$  was determined (**Figure 2B**), demonstrating the high affinity of AIRAP toward arsenite.

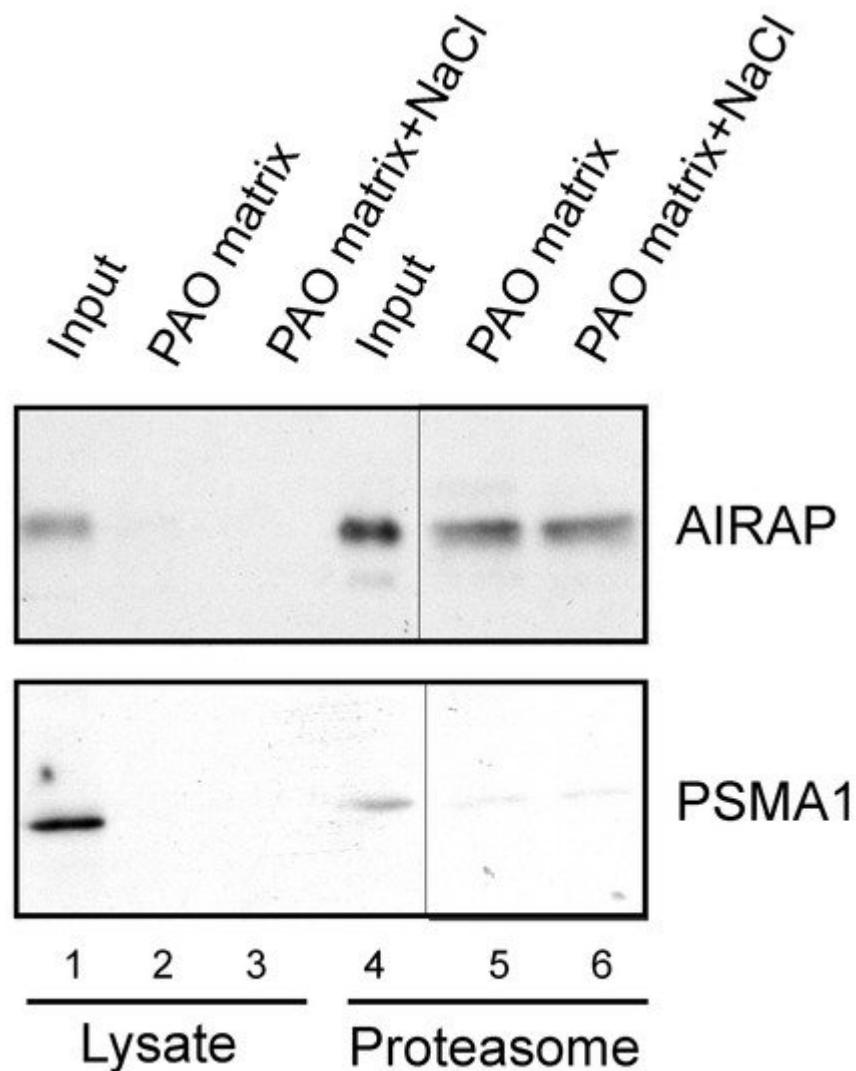


**Figure 2.** Dependence on zinc and MST measured affinity. **(A)** The presence of arsenite bound to AIRAP was determined by ICP-OES. Recombinant AIRAP was incubated in the presence ( $\text{Zn}^+$ ) or absence ( $\text{Zn}^-$ ) of zinc, prior

to arsenite exposure. Bound material was extensively washed and arsenite content determined. (B) Recombinant AIRAP (1  $\mu$ M) was incubated with a range of arsenite dilutions as indicated, and intrinsic fluorescence was measured by MST. Analysis revealed  $K_d$  values of  $7 \text{ nM} \pm 0.87 \text{ nM}$ .

### 3. AIRAP Binding to Arsenite In Vivo

After examining the *in vitro* binding of AIRAP to arsenite, we addressed the ability of endogenous AIRAP to bind arsenite. To this end, we used the cell lysate as a source for AIRAP. Upon endogenous expression of AIRAP (induced by arsenite treatment), AIRAP could not be detected on the PAO matrix from total lysate (**Figure 3**, lanes 2 and 3) but could be detected from purified proteasomes (**Figure 3**, lanes 5 and 6). It is important to note that the interaction was stable even after a high-salt wash, as would be expected from a metal-coordinated binding to the PAO matrix (lane 6). In addition, although catalytic particle subunit PSMA1 was present in the total lysate and the proteasome fractions (**Figure 3** lane 4), it was barely detectable in the PAO matrix-purified fractions; thus, PSMA1 served as a specificity control for the PAO matrix binding. We reasoned that the lack of AIRAP binding to the PAO matrix in the total lysate may have been due to the existence of other proteins that competed towards the PAO matrix binding, or it may have indicated a structural change that AIRAP may undergo upon proteasome binding. This transition may be intended towards optimal binding to arsenite only upon proteasome binding.

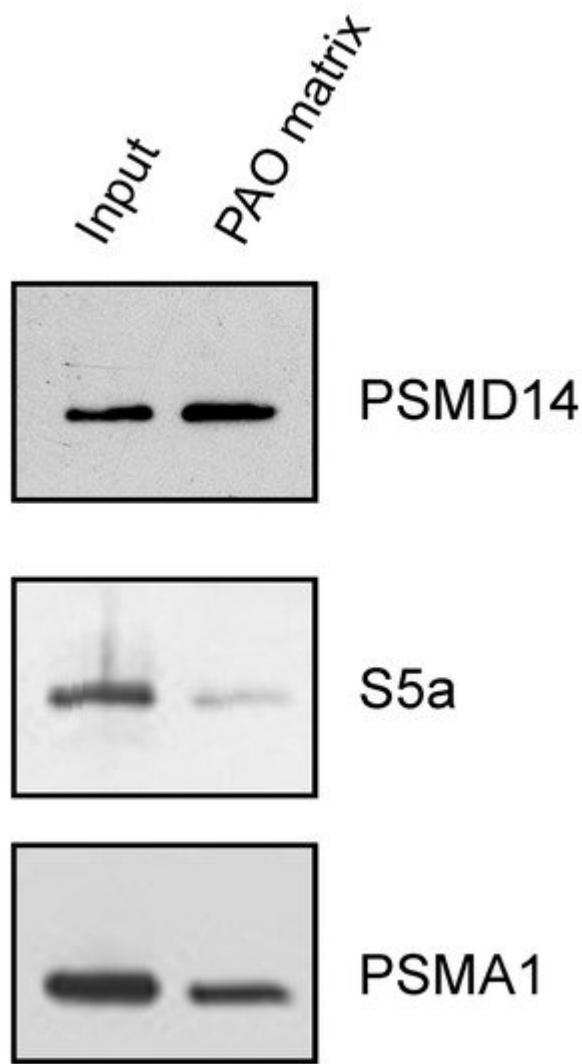


**Figure 3.** Endogenous proteasomal AIRAP binds arsenite. Cell lysates acquired from arsenite-treated cells were incubated with a PAO matrix. AIRAP and 20S proteasomal content (PSMA1) were evaluated by immunoblot (lanes 1–3). Alternatively, PAO matrix binding was performed using soluble proteasomes purified from arsenite-treated cells (lanes 4–6). Where indicated, PAO matrix was washed with high salt to evaluate impact on PAO binding (lanes 3 and 6).

#### 4. Endogenous Rpn11/PSMD14 Binds PAO Matrix More Strongly Than Other Proteasome Subunits

After establishing the ability of AIRAP to bind arsenite, we sought to understand the extent to which arsenite-binding proteins bind the proteasome and how this function may relieve UPS impairment upon arsenite exposure. We reasoned that the 26S proteasome has one metal-binding protein, Psmd14/Rpn11, that functions as a proteasomal deubiquitination enzyme [29][30]. Possible impairment of Psmd14 activity may occur if arsenite binding perturbs metalloprotease activity, as previously reported for Rpn11 metal selectivity [31]. This, in turn, will lead to polyubiquitin accumulation and impaired turnover of UPS substrates. We therefore sought to determine whether

Psmd14 can bind the PAO matrix. Affinity-purified soluble proteasomes were mixed with the PAO matrix and the Psmd14, S5a (an additional 19S subunit) and Psma1 (20S subunit) content was evaluated by immunoblot. As seen in **Figure 4**, Psmd14 was enriched on the PAO matrix. It is important to note that other proteasome subunits, while present, were not enriched as significantly as Psmd14 on the PAO matrix (compare PAO matrix lane and input lane for all evaluated proteasome subunits). These results are in line with Psmd14 being the proteasomal target for AIRAP binding.



**Figure 4.** PSMD14 binding to PAO. Soluble proteasomes were incubated with PAO matrix, extensively washed and PSMD14, 19S (S5a), and 20S (PSMA1) content revealed by immunoblot.

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