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Permalink
https://escholarship.org/uc/item/22z1b4k5

Journal
Nucleic acids research, 40(20)

ISSN
0305-1048

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Publication Date
2012-11-01

DOI
10.1093/nar/gks826

Peer reviewed
X-ray structure of the fourth type of archaeal tRNA splicing endonuclease: insights into the evolution of a novel three-unit composition and a unique loop involved in broad substrate specificity

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Received July 23, 2012; Revised and Accepted August 8, 2012

ABSTRACT

Cleavage of introns from precursor transfer RNAs (tRNAs) by tRNA splicing endonuclease (EndA) is essential for tRNA maturation in Archaea and Eukarya. In the past, archaeal EndAs were classified into three types (α₀², α₄ and α₂β₂) according to subunit composition. Recently, we have identified a fourth type of archaeal EndA from an uncultivated archaeon Candidatus Micrarchaeum acidiphilum, referred to as ARMAN-2, which is deeply branched within Euryarchaea. The ARMAN-2 EndA forms an ε₂ homodimer and has broad substrate specificity like the α₂β₂ type EndAs found in Crenarchaea and Nanoarchaea. However, the precise architecture of ARMAN-2 EndA was unknown. Here, we report the crystal structure of the ε₂ homodimer of ARMAN-2 EndA. The structure reveals that the ε protomer is separated into three novel units (εᴺ, ε and εᶜ) fused by two distinct linkers, although the overall structure of ARMAN-2 EndA is similar to those of the other three types of archaeal EndAs. Structural comparison and mutational analyses reveal that an ARMAN-2 type-specific loop (ASL) is involved in the broad substrate specificity and that K161 in the ASL functions as the RNA recognition site. These findings suggest that the broad substrate specificities of ε₂ and α₂β₂ EndAs were separately acquired through different evolutionary processes.

INTRODUCTION

Transfer RNA (tRNA) is an adapter molecule that acts as a translator of genetic information from nucleotide sequence of messenger RNA to amino acid sequence of protein. Because tRNA needs to go through a maturation process in order to synthesize proteins correctly and smoothly, tRNA maturation is essential for life. tRNA splicing, which removes introns and joins exons in precursor (pre)-tRNA, is an important process in tRNA maturation.

Many interruptions of pre-tRNA with introns have been found in all three domains of life. In Eukarya, most introns are predominantly located in the canonical position between nucleotide positions 37 and 38 in the anticodon loop of tRNA, while archaeal introns are located not only in the canonical position but also in various non-canonical positions including the D-and T-loops, the variable region and the aminoacyl stem (1). In some cases, single archaeal pre-tRNAs include two or three introns, called multiple-introns, in non-canonical positions (2,3). The introns in eukaryotic cytoplasmic and archaeal pre-tRNA are removed by a tRNA splicing endonuclease (EndA) (4–6). The eukaryotic EndA consists of four subunits (SEN2, SEN15, SEN34 and SEN54) (7,8). In contrast, archaeal EndAs are classified into three types by subunit composition, namely, homotetramer (ε₄), homodimer (ε₂) and heterotetramer (ε₂β₂) (9) and Figure 1]. Figure 1 shows the structures and characteristics of the three types of archaeal EndAs. The ε subunit in the archaeal EndAs is a catalytic subunit and
Figure 1. Structures and characteristics of three types of archaeal EndAs. The subunit interactions are represented by cartoon models on the left side. The β–β interaction responsible for inter/intraunit formation, the L10 loop and pocket responsible for dimer/tetramer formation are highlighted. The catalytic triads are marked by green circles. The right panels show the ribbon models of EndAs. The sources of EndAs are shown in parentheses and the substrate specificities of EndAs are shown next to the parentheses: (A) α4 type MJ-EndA; (B) α2 type AFU-EndA; (C) αβ2 type APE-EndA. Full names of the archaea species are as follows: MJ, Methanocaldococcus jannaschii; AFU, Archaeoglobus fulgidus; APE, Aeropyrum pernix. The secondary structure diagrams of substrate RNA motifs. The splicing sites are indicated using arrows. B and H represent bulge and helix, respectively. The h and h’ indicate the helices close to the 3-nt bulge on the exonic side and intronic side, respectively. (D) Left, strict BHB motif; Right, relaxed BHB motifs (hBH and HBh’).
shares homology with SEN2 and SEN34 subunits of
eukaryotic EndA, implying a common evolutionary
origin between the eukaryotic and archaeal EndAs
(7,10). Both the eukaryotic and archaeal EndAs are
proposed to use a similar cleavage chemistry to that of
ribonuclease A (6,11). The archaeal \( \alpha \) subunit has a cat-
alytic triad, L-10 loop and pocket (Figure 1). In the case of
\( \alpha_2 \) EndAs, the \( \alpha \) subunit contains two \( \alpha \) units joined by a
polypeptide linker (Figure 1B). In Figure 1, the locations
of the catalytic triad in archaeal EndAs are marked by
green circles. The negatively-charged L-10 loop and posi-
tively-charged pocket contribute to subunit interaction
and are conserved in the three types of archaeal EndAs.
However, the substrate recognition mechanisms of EndAs
are different to some extent. The eukaryotic EndA
requires the mature domain of pre-tRNA for the recogni-
tion of cleavage sites in the canonical position (12),
although the three types of archaeal EndAs can remove
introns with a bulge–helix–bulge (BHB) motif irrespective
of the existence of the pre-tRNA mature domain.
Furthermore, the \( \alpha_2 \beta_2 \) type of archaeal EndA possesses
a broad substrate specificity that recognizes relaxed BHB
motifs of various lengths and disruption of either the 5'-
or 3'-bulge in the BHB (so-called BBh' and hBH) as well as
the strict BHB motif (Figure 1C and D). In contrast, the
\( \alpha' \) and \( \alpha_2 \) type EndA recognize only the strict BHB motif
(13–19). The BBh' and hBH motifs are often found at
non-canonical positions in introns of pre-tRNAs from
Crenarchaeae and Nanoarchaeae, consistent with the posses-
sion of the \( \alpha_2 \beta_2 \) type EndA (3). Some of these pre-
tRNAs are spliced into two or three gene fragments at
different loci and are called split or tri-split tRNAs,
respectively (20,21). Furthermore, permuted tRNA, in
which the 5’ and 3’ halves of the coding sequences
separated by intervening elements have their positions
switched, has been discovered in some genera of
Crenarchaeae (22). Only the \( \alpha_2 \beta_2 \) type of archaeal EndA
with broad substrate specificity has the ability to excise
non-canonical introns, suggesting the coevolution of dis-
rupted tRNA gene diversity and EndA architecture
(16,21). In Crenarchaeal EndAs, the Crenarchaeae-
specific loop (CSL) is conserved in the catalytic \( \alpha \)
subunit (19). In our previous study, it was revealed that
the CSL is responsible for the broad substrate specificity
and that a conserved Lys residue in the CSL functions as
the substrate recognition site (23).

Recently, we found a fourth type of archaeal
EndA from an uncultivated archaeon Candidatus
Micarchaeum acidiphilum, referred to as ARMAN-
(Archaeal Richmond Mine Acidophilic Nanoorganism)-2
(24), which was discovered in an acid mine drainage site
at Iron Mountain in Northern California (25). Our bio-
chemical and bioinformatic analyses have led us to
propose that the ARMAN-2 EndA has a novel three-unit
architecture? The precise architecture of three-unit inter-
actions will provide new insights into the molecular evo-
lution of archaeal EndA. Furthermore, remarkably, the
ARMAN-2 EndA possesses a broad substrate specificity
that cleaves introns with both strict and relaxed BHB
motifs despite lacking the CSL region. What structural
properties of ARMAN-2 EndA confer the broad substrate
specificity? Structural determination of ARMAN-2
EndA is necessary to address these issues. We present
herein an X-ray crystal structure of ARMAN-2 EndA,
demonstrating a novel three-unit arrangement of the \( \epsilon_2 \)
homodimeric complex. Our structural comparison of
ARMAN-2 (\( \epsilon_2 \)) and the other three types (\( \alpha' \), \( \alpha_2 \)
and \( \alpha_2 \beta_2 \)) of archaeal EndAs shows that the ARMAN-2
EndA possesses an ARMAN-2 type-specific loop (ASL).
Our structure-guided mutagenesis study identified the
catalytic residues and revealed that the ASL is responsible
for the broad substrate specificity. Furthermore, our study
suggests that the Lys residue in the ASL plays the same
role as the Lys residue in the CSL for the broad substrate
recognition and that the ASL has been acquired by a dis-
tinctly independent evolutionary pathway to the CSL.

MATERIALS AND METHODS

Protein expression and purification

A pET-23b vector (Novagen) harboring an ARMAN-2
EndA gene attached to a 6× His tag at its C-terminus has
been previously constructed (24). The plasmid
was used for overexpressing the recombinant ARMAN-2
EndA in Escherichia coli Rosetta 2(DE3) strain
(Novagen). Escherichia coli cells harboring the plasmid
were grown in LB media supplemented with 100 µg/ml
of ampicillin at 37°C, and then isopropylthio–β-
galactoside (IPTG) was added to a final concentration of
0.5 mM when the cells density reached OD600 = ~0.8.
After cultivation at 20°C for 24h, the cells were harvested
by centrifugation (6000 rpm at 4°C for 20 min).
The cells were suspended in 15 ml buffer A [20 mM Tris–HCl (pH 7.6),
200 mM KCl, 20 mM imidazole 10 mM 2-mercaptoethanol and 5% glycerol]
supplemented with protease inhibitor cocktail (Roche) and then disrupted
with an ultrasonic disruptor (model VCX-500, Sonics &
Materials., Inc., USA). A fraction of E. coli proteins was
denatured by heat treatment at 50°C for 20 min and
removed by centrifugation (18 000 rpm at 4°C for
20 min). The supernatant was loaded onto a Ni–NTA
Superflow column (Qiagen) equilibrated with buffer
A and then the enzyme was eluted by buffer A containing
500 mM imidazole. The eluted fractions were collected
and then loaded onto a HiTrap Heparin–Sepharose
column (GE Healthcare) equilibrated with buffer B
[20 mM Tris–HCl (pH 7.6), 50 mM KCl, 10 mM
2-mercaptoethanol and 5% glycerol]. The bound protein was eluted by a linear gradient of buffer B from 50 mM to 1 M KCl. The eluted fractions were collected and then concentrated to ~3 ml volume using Amicon Ultra-15 centrifugal filter units. Finally, the concentrated protein was applied to a HiLoad 16/60 Superdex 75 pg column (GE Healthcare) equilibrated with buffer C [20 mM Tris–HCl (pH 7.6), 700 mM NaCl, 10 mM 2-mercaptoethanol and 5% glycerol]. The single peak fractions were collected. Mutant genes were generated using the QuickChange site-directed mutagenesis kit (Stratagene), and the mutations were verified by DNA sequencing. Mutant proteins were expressed and purified in the same manner as the wild-type protein. The recombinant Archaeoglobus fulgidus (AFU)-EndA and its chimera mutants were prepared as reported previously (23). The protein purities were confirmed by SDS–PAGE (Supplementary Figure S1).

Crystallization

The single-peak fractions from the Superdex-75 gel-filtration column were pooled and then concentrated to ~10 mg/ml using Vivaspin 15 SR centrifugal filter units (Sartorius stedim biotech). Initial trials for crystallization of the ARMAN-2 EndA were performed by the hanging-drop vapor diffusion method using a Crystal Screening Kit (Hampton Research). The drop solution was equilibrated against 200 μl of reservoir solution at 22°C. A few crystals were obtained under some of the tested conditions which contained PEG 3350 as the precipitant. Based on the initial crystallization conditions, we then searched for optimum conditions. When the ARMAN-2 EndA protein solution was mixed with an equal volume of a crystallization solution that contained 18% PEG3350 and 0.2 M tri-ammonium citrate (pH 7.0), crystals grew within 5 days at 22°C producing full-sized rectangular-shaped (200 × 100 × 100 μm) crystals. For the experimental phase determination by single-wavelength anomalous dispersion (SAD) method, the crystal was soaked in mother liquor supplemented with 0.4 mM KPtCl4 at 22°C for 16 h. Cryo-protection of the native and Pt-induced crystals was achieved by stepwise transfer to the respective artificial mother liquor containing 25% glycerol. The crystals were then flash-frozen in liquid nitrogen.

Data collection and structure determination

X-ray diffraction data sets from native crystals (λ = 1.0000) and SAD data sets from Pt-induced crystals (λ = 1.0717) were collected at 100 K on the BL38B1 beamline at SPring-8 (Hyogo, Japan). All data sets were processed, merged and scaled using the HKL2000 program (26). Using the deduced Pt-SAD data set, all 19 Pt positions were identified and refined in the orthorhombic space group $P 2_12_12_1$, and the initial phase was calculated by using AutoSol in PHENIX (27), followed by automated model building using RESOLVE (28). The resulting map and partial model were used for manually building the model using COOT (29). The model was further refined by using PHENIX (27). Using the native data set and the refined model as a search coordinate, the structure of the ARMAN-2 EndA was determined by molecular replacement with the Phaser program (30). The model was further manually built with PHENIX (29) and refined with PHENIX (27). The structure of ARMAN-2 EndA was refined to $R_{work}/R_{free}$ of 21.8%/25.7% at 2.25 Å resolution (Table 1). The space group of the crystal belonged to $P 3_2_1$, where two ARMAN-2 EndA molecules are present in an asymmetric unit. The final model contained residues 2-387 (chain A and B) and 152 water molecules. The final model of the ARMAN-2 EndA structure was further checked using PROCHECK (31), showing the quality of the refined model. Ramachandran plots (%) of the ARMAN-2 EndA structure are tabulated in Table 1. The structure factor and coordinates have been deposited in the Protein Data Bank (PDB code 4FZ2). All structural figures were generated by PyMOL (DeLano Scientific, Palo Alto, CA).

### Intron-cleavage assay by the splicing endonuclease

The transcripts of ARMAN-2 pre-tRNA[^Cys] (GCA) were prepared using T7 RNA polymerase as described in our previous report (24). Splicing reactions were performed as follows. 1.0 μg EndA was mixed with 0.2 nmol transcripts in 50 μl buffer D [50 mM

### Table 1. Data collection and refinement statistics

|                          | ARMAN-2 EndA | ARMAN-2 EndA Pt-derivative |
|--------------------------|--------------|-----------------------------|
| **Data collection**       |              |                             |
| Space group              | $P 3_2$      | $P 2_12_12_1$               |
| Cell dimensions          |              |                             |
| $a$, $b$, $c$ (Å)        | 112.02, 112.02, 81.08 | 75.62, 85.17, 140.19      |
| $\alpha$, $\beta$, $\gamma$ (°) | 90, 90, 90 | 90, 90, 90          |
| Resolution (Å)           | 50 to 2.25   | 50 to 2.05                 |
| Resolution (Å)           | (2.33–2.25)  | (2.07–2.00)                |
| $R_{merge}$ a            | 6.5 (49.2)   | 5.6 (35.1)                 |
| $I/\sigma(I)$            | 39.2 (4.7)   | 12.8 (10.6)                |
| Completeness (%)         | 99.4 (95.4)  | 99.7 (98.6)                |
| Redundancy               | 5.9 (5.5)    | 12.9 (11.7)                |
| **Refinement**           |              |                             |
| Resolution (Å)           | 37.4–2.25   |                             |
| No. reflections          | 49748        |                             |
| $R_{work}/R_{free}$ b    | 21.8/25.7    |                             |
| No. atoms                | 6302         |                             |
| Protein                  | 6454         |                             |
| Water                    | 152          |                             |
| Avg. B-factors (Å²)      | 51.5         |                             |
| R.m.s.d.                 |              |                             |
| Bond lengths (Å)         | 0.006        |                             |
| Bond angles (°)          | 1.0          |                             |
| Ramachandran plot (%)    | 90.2         |                             |
| Most favored             | 9.1          |                             |
| Additional allowed       | 0.7          |                             |
| Generously allowed       | 0.0          |                             |
| Disallowed               | 0.0          |                             |

The value in the parentheses is for the highest resolution shell.  

\[ R_{merge} = \Sigma \Sigma_j <\hat{r}(h)> - <\hat{r}(h)>/\Sigma_j <\hat{r}(h)> \]  

where $<\hat{r}(h)>$ is the mean intensity of symmetry-equivalent reflections.  

\[ R_{work} = \Sigma (I_F(\text{obs}) - I_F(\text{calc})/\Sigma I_F(\text{obs}) \]

\[ R_{free} = R\text{-factor for a selected subset (10%) of reflections that was not included in earlier refinement calculations}\]
Tris–HCl (pH 7.6), 5 mM MgCl₂, 6 mM 2-mercaptoethanol and 50 mM KCl] and incubated at 50°C. Aliquots (10 µl) were removed at 0, 10, 30 and 60 min and were analyzed by 15% PAGE/7 M urea. The gel was stained with 0.05% toluidine blue.

RESULTS AND DISCUSSION

Overall structure

We initially crystallized an ARMAN-2 EndA to obtain structural information. Two different space groups were found in different ARMAN-2 EndA crystals under the same crystallization conditions. One crystal belonged to the orthorhombic space group P2₁2₁2₁, whereas the other belonged to the trigonal space group P3₂₁. In this study, we determined the structure of ARMAN-2 EndA from the latter crystal at 2.25 Å resolution (Figure 2A). Although the structure from the former crystal could be solved at 2.00 Å resolution, it exhibited many disordered regions due to the effects of crystal packing (data not shown). The final model of ARMAN-2 EndA contains two molecules per asymmetric unit. The two molecules are structurally almost identical (R-factor = 21.8 and Rₚᵣₑเอก = 25.7 in Table 1).

The ARMAN-2 EndA is composed of two ε protomers producing a homodimeric subunit structure, ε₂ (Figure 2A). The overall shape of the ε₂ homodimer structure is like a rectangular parallelepiped. The ε protomer consists of 10 α helices and 23 β strands (Figure 2B and C). Furthermore, the structure can be separated into three units, the αN unit (2–97 residues; orange), the α unit (126–288 residues; pink) and the βC unit (301–387 residues; cyan). The three units are connected by two linkers, linker 1 (98–125 residues; black) and linker 2 (289–300 residues; black): linker 1 connects the αN and α units, whereas linker 2 connects the α and βC units. The αN unit (orange) is composed of a mixed anti-parallel and parallel β sheet (β₁–β₅), four α helices (α₁–α₄) and one β strand (β₆). The β strand of the αN unit, the β strand of linker 1 and five β strands (β₁₈, β₁₉, β₂₁, β₂₂ and β₂₃) of the βC unit participate in forming one mixed anti-parallel and parallel β sheet. This β₇–β₂₃ interaction probably prevents structural fluctuation of linker 1. The βC unit (cyan) consists of one β sheet (β₁₈, β₁₉, β₂₁, β₂₂ and β₂₃), one α helix (α₁₀) and one β strand (β₂₀). The β sheet is structurally sandwiched by two α helices (α₄ and α₁₀), thereby stabilizing the unit interaction between the αN and βC units. Furthermore, the β₂₀ and β₂₃ strands interact with the β₁₆ and β₁₇ strands in the α unit, respectively. These two anti-parallel β sheets connect the βC and α units. The α unit (pink) can be separated into two subdomains, the N-terminus and C-terminus. The N-terminal subdomain is composed of a mixed anti-parallel and parallel β sheet (β₈–β₁₂ and three α helices (α₅–α₇), and the C-terminal subdomain is composed of a mixed anti-parallel and parallel β sheet (β₁₃–β₁₅ and β₁₇), two α helices (α₈–α₉) and one β strand (β₁₆). The five α helices (α₅–α₉) are placed around the two β sheets. Thus, these intra-unit interactions probably contribute to maintenance of the structural integrity of ARMAN-2 EndA. The configuration of secondary structures in the αN and βC unit overlaps with that of the α unit (Figure 2D). Thus, this configuration is commonly observed in the α and β subunits of the ε₂β₂ EndAs (19,23,32). Furthermore, our structure-based sequence alignment analysis has shown that the overlap region of the αN, α and βC units is found in the N-terminal subdomain of the α subunit in the ε₂ type EndAs, the entire domain of the α subunit in the ε₂β₂ type EndAs and the C-terminal subdomain of the β subunit in the ε₂β₂ type EndAs (Supplementary Figure S2), although the βC unit includes an amino acid sequence (373–387 in ARMAN-2 EndA), which is found in the α subunit instead of the β subunit in the case of Crenarchaeal ε₂β₂ type EndAs (Supplementary Figure S2B). Based on these structural observations, we have redefined the fourth type of archaeal ε₂ EndA, where ε is three-units (αN–α–βC).

Structural comparison with three types of archaeal EndAs

Our current structural study clarified the ε₂ subunit structure of ARMAN-2 EndA (Figure 3). The architecture of the three units and subunit interactions were far beyond our previous expectations because two long linkers connect the three units in ARMAN-2 EndA. This architecture is not observed in the other three types of EndAs (Figure 1). Nevertheless, the overall shape and size of the ε₂ structure of ARMAN-2 EndA is very similar to those of the other three types of archaeal EndAs (Figure 1 and Supplementary Figure S3). In addition to the three types of archaeal EndAs, a structural homology search by the Dali sever (33) confirms that the structure of βC and α units in ARMAN-2 EndA is homologous to that of a subunit (SEN15) of human EndA and that of prokaryotic DNA restriction enzymes.

As shown in Figure 1 and 3, two β–β strand interactions at the domain interface are conserved in all four types of archaeal EndAs (24). The interactions are shown to be responsible for intra/subunit interactions such as the α–α subunit assembly in the α₄ type EndA, the α–α domain assembly in the ε₂ type EndA and the β–β subunit assembly in the ε₂β₂ type EndA (11,19,32). However, in the case of ε₂ ARMAN-2 EndA, the β–β strand (β₂₂–β₂₃) interaction does not directly contribute to the interaction between the α and βC units since the β₂₂ and β₂₃ strands are parts of the C-terminal βC unit (Figures 2C and 3A). Instead, two anti-parallel β strand interactions (β₂₀–β₁₆ and β₂₃–β₁₇) connect the α units since the β₂₂ and β₂₃ strands are parts of the C-terminal βC unit (Figures 2C and 3A). Furthermore, the β₆–β₁₈ strand interaction appears to contribute to the assembly of αN and βC units together with formation of a sandwich by the β sheet of the βC unit and two α helices (α₄ and α₁₀). Because the three-unit architecture is connected by two linkers, the linkers enable it to easily form a complete ε protomer as compared to the α–α subunit assembly in ε₂ type EndA and the α–β subunit assembly in ε₂β₂ type EndAs. Therefore, the linkers play an important role in the three-unit architecture. In contrast, as previously expected from our bioinformatics study (24), the three-unit molecule assembles with another through the interaction of a negatively-charged L10 loop with a
Figure 2. Crystal structure of ARMAN-2 EndA. (A) Ribbon stereo diagram of the overall structure of the functional ε2 homodimeric complex, where ε stands for the union of three units (αε–βε–βε'). The αε unit, α unit, βε unit and two linker regions are colored orange, pink, cyan and black, respectively. The N- and C-terminal ends are labeled as N and C, respectively. (B) Ribbon diagram of the ε protomer. The secondary structures of the α helix and β strand are labeled (in order) as the α and β, respectively. The αε unit, α unit, βε unit and two linker regions are colored as described above, respectively. (C) A secondary structure topology diagram of the ε protomer. The α helices and β strands are represented by circles and triangles, respectively. The αε, α and βε units are colored as in Figure 2A and B. (D) Superimposition diagram of Cα atoms of the αε (orange) and βε (cyan) units onto that of the α unit (pink) of ARMAN-2 EndA.
positively-charged pocket of the \( \alpha \) unit in the opposing three-unit molecule (Figure 3B). These electrostatic interactions are observed in other EndA structures (Figure 1), suggesting the structural and/or functional importance of these interactions. Figure 3B shows the molecular interaction of the L10 loop and positively-charged pocket in the ARMAN-2 EndA. Two salt-bridge interactions (D357–K228 and E359–R234) are observed. The positively and negatively-charged amino acid residues are conserved in almost all EndAs as reported previously (24). In fact, our previous mutagenesis study has shown that the D357A mutant of ARMAN-2 EndA barely cleaves the introns from pre-tRNA\(^{\text{Ile}}\) and pre-tRNAC\(^{\text{Cys}}\) (24), suggesting that the salt-bridge interaction (D357–K228) is required for the formation of functional \( \varepsilon_2 \) homodimer of ARMAN-2 EndA. Although we could not observe the dissociation of \( \varepsilon_2 \) homodimer into \( \varepsilon \) protomer in the D357A mutant under our cross-linking analysis (24), the breakage of the salt-bridge interaction (D357–K228) probably induce the conformational change of the side chain of K228. In the ARMAN-2 EndA structure, the side chain of K228 is located close to that of R275, which is the putative RNA recognition residue as described in the next section, at the 4.2 \( \AA \) distance between the K228 C\( ^\gamma \) and R275 N\( _2 \). Thus, the conformation of the side chain of R275 can also be changed by the electrostatic repulsion between the K228 and R275 in the D357A mutant, resulting in the loss of intron-cleavage activity.

The active site

It has been reported that three catalytic residues (tyrosine, histidine and lysine) as well as two substrate recognition residues (two arginines) are conserved in the \( \alpha \) subunit of the EndA from Euryarchaeal (6,11). Our structural study and amino acid sequence alignment strongly suggest that the Y236, H251 and K282 residues are the catalytic residues and that the R275 and W384 are the possible substrate recognition residues in the case of the ARMAN-2 EndA (Supplementary Figure S2B). These five residues are located on the enzyme surface around the expected catalytic pocket (Figure 4A) and can be arranged at similar locations to that of their counterpart residues in the three types of archaeal EndAs (Supplementary Figure S3). To clarify whether the catalytic triad of ARMAN-2 EndA is indeed formed...
by the three predicted residues (Y236, H251 and K282), we constructed three alanine mutants (Y236A, H251A and K282A) and then performed an intron-cleavage assay with the mutants. Prior to the mutant study, we optimized the assay conditions by using the wild-type ARMAN-2 EndA and two pre-tRNA transcripts (pre-tRNA\textsuperscript{Ile} and pre-tRNA\textsuperscript{Cys}) previously used as substrates (24). The pre-tRNA\textsuperscript{Ile} and pre-tRNA\textsuperscript{Cys} contain a strict BHB motif at the canonical position and relaxed BHB motif at a non-canonical position, respectively (Figure 4B and C). The ARMAN-2 EndA completely removed the introns from both the pre-tRNA\textsuperscript{Ile} and pre-tRNA\textsuperscript{Cys} within 60 min. Next, we assayed for the removal of intron by the three mutants, Y236A, H251A and K282A. All three mutants failed to remove the introns from both the pre-tRNA\textsuperscript{Ile} and pre-tRNA\textsuperscript{Cys} (Figure 4D and E), suggesting that the Y236, H251 and K282 residues play an important role as the catalytic triad of ARMAN-2 EndA. Of the RNA recognition residues (R275 and W384) of ARMAN-2 EndA, the typtophan is only conserved in the\(a\textsubscript{2}\beta\textsubscript{2}\) type EndAs from Crenarchaea and Nanoarchaea (Supplementary Figure S2B). Instead of the tryptophan residue, an arginine residue is conserved in the\(a\textsubscript{0}\beta\textsubscript{0}\) and\(a\textsubscript{4}\beta\textsubscript{4}\) EndAs from Euryarchaea. Two arginine residues in the\(a\textsubscript{0}\beta\textsubscript{0}\) EndA capture the adenine base in the first bulge of the BHB motif by cation–π interactions (6,11).

Figure 4. The active site of ARMAN-2 EndA. (A) Close-up view of the active site. The catalytic triad comprised of three catalytic residues (Y236, H251 and K282) and two putative RNA recognition residues (R275 and W384) are shown by stick model (green). (B) Time-dependent cleavage of ARMAN-2 pre-tRNA\textsuperscript{Ile}(UGU) by the wild-type ARMAN-2 EndA. Predicted secondary structure of ARMAN-2 pre-tRNA\textsuperscript{Ile}(UGU) labeled with two arrows indicating the splicing sites is shown at the left side of the gel. (C) Time-dependent cleavage of ARMAN-2 pre-tRNA\textsuperscript{Cys}(GCA). Predicted secondary structure of ARMAN-2 pre-tRNA\textsuperscript{Cys}(GCA) indicating the splicing sites is shown at the left side of the gel. (D) Cleavage activities of the wild-type and three mutants (Y236A, H251A and K282A) using ARMAN-2 pre-tRNA\textsuperscript{Ile}(UGU). (E) Cleavage activities of wild-type and three mutants (Y236A, H251A and K282A) using ARMAN-2 pre-tRNA\textsuperscript{Cys}(GCA). Reaction mixtures were separated on 15% polyacrylamide/7 M urea gels. The cleaved products are shown using arrows at the right side of the gel.
A tryptophan residue can act as an alternative for the arginine because its indole ring can form a hydrophobic interaction with the nucleotide instead of the cation–π interaction.

**Broad substrate specificity**

Because the catalytic and substrate recognition residues of ARMAN-2 EndA are conserved in all types of archaenal EndAs as described above, these residues are probably not involved in the broad substrate specificity of the ARMAN-2 EndA. We searched for the specific regions responsible for the specificity of ARMAN-2 EndA based on the structure-based sequence alignment. As a result, two specific regions were found (Supplementary Figure S2B and S2C highlighted in cyan and green). To confirm the findings, we performed a structural comparison of the specific regions of ARMAN-2 EndA and counterparts, namely, the 5β2 type Aeropyrum pernix (APE)-EndA and Nanoarchaeum equitans (NEQ)-EndAs (Figure 5A and B). Shown in Figure 5A is the specific region (158–168 residues) of ARMAN-2 EndA which forms a loop structure on the enzyme surface close to the catalytic triad. The ARMAN-2 type-specific loop (ASL) is positioned at a similar location to the CSL in APE-EndA, which plays a significant role in the broad substrate specificity. The conformation of ASL resembles that of CSL, although amino acid similarity and identity are not found with the exception of one positively-charged residue (K161 in ARMAN-2 EndA and K44 in APE-EndA in Figure 5A right panel). Notably, the configurations of K161 in ASL and K44 in CSL are such that they are positioned in the same direction. The K44 residue in APE-EndA has been shown to be essential for the enzymatic activity and broad substrate specificity (23). Accordingly, we hypothesized that the K161 residue in the ASL is probably involved in the splicing activity and broad substrate specificity of ARMAN-2 EndA. When we superimposed the structure of ARMAN-2 EndA onto that of NEQ-EndA (Supplementary Figure S3D and S3E), another specific loop (240–247 residues) of ARMAN-2 EndA was found to be positioned in the same way as the specific loop (90–98 residues) of NEQ-EndA (Figure 5B). These loops are close to the catalytic triad, and the catalytic His residue is located in both of these specific loops. Furthermore, it was expected that some positively-charged residues (K93, K94, K96 and R97) in the specific loop of NEQ-EndA would be important for the broad-specificity (32,34). We hypothesized that the K241 and K244 residues of ARMAN-2 EndA may correspond to the positively-charged residues of NEQ-EndA, although there is no sequence similarity or identity between the ARMAN-2 and NEQ-EndAs in this specific region.

To examine whether these residues (K161, K241 and K244) are implicated in the enzymatic activity and broad substrate specificity of ARMAN-2 EndA, we constructed three alanine mutants (K161A, K241A and K244A) and subsequently conducted an intron cleavage assay of the mutants (Figure 6A and B). As shown in Figure 6A and B, the wild-type ARMAN-2 EndA, K241A and K244A mutants could remove introns from both the pre-tRNA\textsuperscript{His} and pre-tRNA\textsuperscript{Cys}. In contrast, the K161A mutant did not cleave the introns. These results demonstrate that the K161 residue is essential for the cleavage of introns with strict and relaxed BHB motifs. To understand the importance of the K161 residue structurally, we constructed a docking model of ARMAN-2 EndA complex with RNA based on the reported α2 type AFU-EndA and RNA complex structure (Figure 7A) (6). The RNA in the reported complex contains a BHB motif. The docking model shows that the K161 residue is situated near the 3-phosphate group adjacent to the bulge structure of the RNA, suggesting that the K161 residue captures this 3’-phosphate group (or 3’-phosphate of the third nucleotide in the loop structure), fixes the substrate, and thereby is essential for cleavage activity. To clarify whether the K161 residue in the ASL plays a key role in determining the substrate specificity, we initially created an AFU EndA mutant protein (AFU-ASL) in which Lys175 was replaced by the ASL sequence (GTYKVKSEH) of ARMAN-2 EndA (Figure 7B and C). We also made one additional mutant (ASL-K178A mutant), in which the K178 residue of the AFU-ASL mutant, corresponding to the K161 residue of AMRNA-2 EndA, was replaced with alanine. We then analyzed the substrate specificity of these two mutants. As shown in Figure 7D, the AFU-ASL and ASL-K178A cleaved the intron with a strict BHB motif from the anticodon loop in a similar manner as wild-type ARMAN-2 and AFU-EndAs. The wild-type AFU-EndA and ASL-K178A mutant, however, barely cleaved the intron with a relaxed BHB motif from the T-loop of the pre-tRNA\textsuperscript{Cys} (Figure 7E). In contrast, the AFU-ASL mutant effectively cleaved the intron from the pre-tRNA\textsuperscript{Cys} just as well as the wild-type ARMAN-2 EndA did although the cleavage fragment of 3’-half with intron was shown. Thus, these results clearly demonstrate that the insertion of the ASL conferred ARMAN-2 EndA-like broad substrate specificity to AFU-EndA, which otherwise has narrow substrate specificity. Furthermore, it was demonstrated that the K161 residue in the ARMAN-2 EndA plays a key role in the broad substrate specificity acting as the RNA recognition site, in a similar way to the function of the K44 residue in the CSL of APE-EndA (23).

**Evolution of the fourth type of archaenal EndA**

With respect to the evolution of three archaenal EndA families (\(\alpha^2, \alpha^4\) and \(\alpha_2\beta_2\)), it has been proposed that the \(\alpha\) subunit gene of \(\alpha_4\) type EndA was first duplicated and then one was subfunctionalized to encode the \(\beta\) subunit, thereby suggesting that the \(\alpha\) and \(\beta\) subunits are evolutionarily derived from the same origin (9). Because of the striking structural and sequential similarities between the ARMAN-2 EndA and the three other types of archaenal EndAs (Figure 2, Supplementary Figure S2 and S3), the \(\varepsilon\) protomer of the ARMAN-2 EndA is likely to also share the common evolutionary origin of the \(\alpha\) and \(\beta\) subunits.

The uncultured acidophilic archaean ARMAN-2 and its lineages were predominantly found in a chemosynthetic biofilm and grown in acidic and metal-rich environments, suggesting a mutualistic relationship with deep-sea hydrothermal vents.
In the biofilm, several eubacteria and archaea including the order of *Thermoplasmatales* are found. Intriguingly, the ARMAN lineages were shown to physically connect to the *Thermoplasmatales* using a 3D cryo-electron tomographic reconstruction (35). Furthermore, a virus was found to be on the cell wall of the ARMAN lineages, indicating an infection of the ARMAN lineages with the virus. Therefore, genetic diversity may occur in the biofilm community via horizontal and/or lateral gene transfer. In fact, ARMAN-2 has many genes homologous to those of Crenarchaea and eubacteria despite the phylogenetic affiliation to the deeply branched Euryarchaea. It is noteworthy that our previous (24) and current studies demonstrate the recombination of the EndA gene in ARMAN-2 cells. Our structure-based sequence alignment shows that the $\alpha^N$ unit is homologous to the N-terminal subdomain of the $\alpha$ subunit from Euryarchaeal EndAs, and that the $\alpha$ and $\beta^C$ units share homology with the $\alpha$ subunit and C-terminal subdomain of the $\beta$ subunit from Crenarchaeal EndAs, respectively (Supplementary Figure S2). Accordingly, ARMAN-2 EndA appears to have undergone a genetic recombination of the three subunits. As a result, the $\alpha^N$, $\alpha$ and $\beta^C$ units are currently found as the structural and functional element of ARMAN-2 EndA. At position 384, a tryptophan residue responsible for RNA recognition site is only found at the end of the $\alpha$ subunit of the EndAs from Crenarchaea and Nanoarchaea. Given these findings, it is likely that the C-terminal subdomain of the Crenarchaeal $\beta$ subunit is...
incorporated into the end of the Crenarchaeal β subunit, resulting in the formation of the α-ββ′ unit of ARMAN-2 EndA. Thus, this could be understood as an example of so-called ‘domain shuffling’ occurring naturally.

The two specific loop regions of ARMAN-2 EndA were proposed as candidates responsible for the broad substrate specificity from our structural comparison (Figure 5). One of these, the ASL, has been shown to be involved in the broad substrate specificity of ARMAN-2 EndA (Figures 6 and 7). The ASL probably has the same function as the CSL from Crenarchaeal EndA. However, no significant sequence similarity is found except for the conserved Lys residue that functions as the substrate recognition site. Therefore, this suggests that the ASL was acquired by a distinctly independent evolutionary pathway to the CSL, so-called ‘convergent evolution’. In contrast, another specific loop is positioned at a location similar to the specific loop of NEQ-EndA (Figure 5B), but has not been shown to be involved in the enzymatic activity and substrate specificity (Figure 6A and B). If the loop had continuous positively-charged residues like the NEQ-EndA, ARMAN-2 EndA may possess the broad substrate specificity even if the ASL is missing. In any case, there seems to be two different structural strategies to obtain the broader specificity as observed in the αββ′ and ε2 types of archaean EndAs gained by convergent evolution. First, the conserved Lys residue on either the ASL or CSL adjacent to the active site functions as the substrate recognition site. Second, the continuous positively-charged residues on the specific loop including the catalytic His residue function as the substrate recognition site. In the case of the ARMAN-2 EndA, the second strategy may have been lost because of an earlier acquisition of the ASL.

A bona fide role for tRNA introns remains unclear except for the methylation on the 2′-O-ribose of G32 and G34 in tRNA1<sup>1tp</sup> from *Haloflexa volcanii* (36). Randou and Söll have argued that the gain of tRNA introns provides protection against integration of mobile genetic elements, such as conjugative plasmids and viruses (37). A total of 56% of tRNA genes are interrupted with both the strict and relaxed BHB motif introns in the ARMAN-2 (23). In contrast, the lineages, ARMAN-4 and ARMAN-5 have only the strict BHB motif introns in 15% of tRNA genes, consistent with possession of the α4 type EndA (24). Because the prototype of archaean EndA was proposed to be an α4 type (9), transition from the α4 to the ε2 type might allow an increase in the number and diversity of tRNA introns at non-canonical positions in ARMAN-2 cells. Furthermore, the CRISPR immune system that protects from virus (38) is absent from the genomes of all three ARMAN groups. Therefore, inclusion of the ASL in the ε2 type of ARMAN-2 EndA may expand the disrupted tRNA genes for defense against the integration of mobile genetic elements as previously proposed in the case of inclusion of the CSL in the Crenarchaeal EndA (23). Moreover, given the report demonstrating that the gain of tRNA introns occurred relatively recently (39,40), incorporation of the ASL region into the ARMAN-2 EndA might have been a dominant advantage for the survival of ARMAN-2 cells in the biofilm community.

In conclusion, our structural study of the ARMAN-2 EndA, which is the fourth type of archaean EndA, has shown the precise architecture of the ε protomer that consists of three units (αε<sup>4</sup>, α and β<sup>4</sup>). The three units form the εε<sup>2</sup> homodimer. There is striking structural and functional similarity among all four types (αε<sup>2</sup>, αεβ<sup>2</sup> and εε<sup>2</sup>) of archaean EndAs, suggesting that the four types of archaean EndAs are derived from a common ancestor. However, the two linker loops connecting the three-unit and the ASL are distinct in the ARMAN-2 EndA. The two linkers play an important role to facilitate the three-unit formation, and the ASL confers the broad substrate specificity on ARMAN-2 EndA. Furthermore, our structure based sequence alignment of the ARMAN-2 EndA exhibits a trace of gene recombination of the α and β subunits from Eurarchaeal and Crenarchaeal EndAs. These results broaden understanding of the mechanism underlying gain of function in protein architecture. In the ASL, the K161 residue functions as a RNA recognition site and thereby broadens the specificity of ARMAN-2 EndA. The ASL has arisen from convergent evolution to play a similar role to the CSL of Crenarchaeal EndA. Inclusion of the ASL in ARMAN-2 EndA may have allowed increasing number and diversity of tRNA introns for the protection from mobile genetic elements. Thus, our findings further enhance the possibility of coevolution of the archaean EndA architecture and disrupted tRNA genes.

**Figure 6.** Intron cleavage activities and specificities of the wild-type ARMAN-2 EndA and its mutants (K161A, K241A and K244A): (A) ARMAN-2 pre-tRNA<sup>1tp</sup>(UGU); (B) ARMAN-2 pre-tRNA<sup>Cys</sup>(GCA). The cleaved products are shown using arrows at the right side of the gel.
ACCESSION NUMBERS
The structure factor and coordinates have been deposited in the Protein Data Bank (PDB code 4FZ2).

SUPPLEMENTARY DATA
Supplementary Data are available at NAR Online: Supplementary Figures 1–3 and Supplementary References [41,42].

ACKNOWLEDGEMENTS
The authors thank the staff members of the beam-line facility at SPring-8 (Hyogo, Japan) for their technical support during data collection. The synchrotron radiation experiments were performed at the BL38B1 in the SPring-8 with the approval of the Japan Synchrotron Radiation Research Institute (JASRI) (Proposal No. 2012A1098).

FUNDING
Funding for open access charge: The Grant-in-Aid for Young Scientists (B) [No. 24770125 to A.H.]; Grant-in-Aid for Science Research (B) [No. 23350081 to H.H.]; Japan Society for the Promotion of Science.

Conflict of interest statement. None declared.
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