Structural Basis for Cyclization Specificity of Two *Azotobacter* Type III Polyketide Synthases

A SINGLE AMINO ACID SUBSTITUTION REVERSES THEIR CYCLIZATION SPECIFICITY*

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Ryutaro Satou†, Akimasa Miyanaga‡, Hiroki Ozawa©, Nobutaka Fun‡, Yohei Katsuyama©, Ken-ichi Miyazono©, Masaru Tanokura©, Yasuo Ohnishi‡©, and Sueharu Horinouchi††

From the Departments of †Biotechnology and ‡Applied Biological Chemistry, Graduate School of Agricultural and Life Sciences, The University of Tokyo, 1-1-1 Yayoi, Bunkyo-ku, Tokyo 113-8657, Japan

**Background:** Type III polyketide synthases (PKSs) show diverse cyclization specificity.

**Results:** A single amino acid substitution in two *Azotobacter* type III PKSs reversed their cyclization specificity. Crystal structures were determined.

**Conclusion:** The volume of the active site cavity is a crucial determinant of the cyclization specificity.

**Significance:** An important insight into the cyclization specificity of type III PKSs was provided.

Type III polyketide synthases (PKSs) show diverse cyclization specificity. We previously characterized two *Azotobacter* type III PKSs (ArsB and ArsC) with different cyclization specificity. ArsB and ArsC, which share a high sequence identity (71%), produce alkylresorcinols and alklyphrones through aldol condensation and lactonization of the same polyketomethylene intermediate, respectively. Here we identified a key amino acid residue for the cyclization specificity of each enzyme by site-directed mutagenesis. Trp-281 of ArsB corresponded to Gly-284 of ArsC in the amino acid sequence alignment. The ArsB W281G mutant synthesized alkylpyrone but not alkylresorcinol. In contrast, the ArsC G284W mutant synthesized alkylresorcinol with a small amount of alklyphrone. These results indicate that this amino acid residue (Trp-281 of ArsB or Gly-284 of ArsC) should occupy a critical position for the cyclization specificity of each enzyme. We then determined crystal structures of the wild-type and G284W ArsC proteins at resolutions of 1.76 and 1.99 Å, respectively. Comparison of these two ArsC structures indicates that the G284W substitution brings a steric wall to the active site cavity, resulting in a significant reduction of the cavity volume. We postulate that the polyketomethylene intermediate can be folded to a suitable form for aldol condensation only in such a relatively narrow cavity of ArsC G284W (and presumably ArsB). This is the first report on the alteration of cyclization specificity from lactonization to aldol condensation for a type III PKS. The ArsC G284W structure is significant as it is the first reported structure of a microbrial resorcinol synthase.

Type III polyketide synthases (PKSs) are simple homodimeric ketsosynthases that are involved in the biosynthesis of various aromatic polyketides in both plants and microorganisms (1–3). Type III PKSs typically catalyze condensation of a starter acyl thioester with several acetate units derived from malonyl-CoA. Three catalytic residues, called a Cys-His-Asn catalytic triad, are responsible for this condensation. The resulting linear polyketomethylene intermediate is subjected to cyclization in the same active site cavity. Type III PKSs synthesize a wide variety of products because they differ in their preference of starter substrate selection in the number of condensation steps and in the mechanism of intramolecular cyclization of polyketomethylene intermediates. Although the starter substrate specificity and polyketide chain extension are largely understood, the cyclization step still remains elusive. Several structural and site-directed mutagenesis studies suggested that the cyclization process is defined by the architecture of the active site cavity; several amino acid residues in the active site cavity were shown to affect the cyclization specificity (1, 4–6).

Previously we characterized two type III PKSs, ArsB and ArsC, responsible for phenolic lipids biosynthesis in a Gram-negative nitrogen-fixing soil bacterium, *Azotobacter vinelandii* (7, 8). Although ArsB and ArsC share 71% amino acid sequence identity, ArsB and ArsC were shown to synthesize alkylresorcinols and alklyphrones, respectively (Fig. 1A), indicating that they are enzymatically distinct type III PKSs. Both ArsB and ArsC employ the same long-chain acyl starter substrate directly received from the type I fatty acid synthase ArsAD and subsequently catalyze three successive extensions with malonyl-CoA followed by the formation of the same tetraketide intermediate. The different catalytic properties between ArsB and ArsC depend on the cyclization specificity. ArsB catalyzes an intramolecular C2-to-C7 (numbering from the cysteine thioester) aldol condensation accompanied by thioester cleavage and

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The atomic coordinates and structure factors (codes 3VS8 and 3VS9) have been deposited in the Protein Data Bank (http://wwwpdb.org/).

† Deceased on 12th July 2009.

‡ To whom correspondence should be addressed. Tel: 81-3-5841-5123; Fax: 81-3-5841-8021; E-mail: ayasuo@mail.ecc.u-tokyo.ac.jp.

2 The abbreviations used are: PKS, polyketide synthase; LC-APCIMS, liquid chromatography-atmospheric pressure chemical ionization mass spectrometry; r.m.s.d., root mean square deviation; ORAS, 2′-oxoalkylresorcylic acid synthase; STS, stilbene synthase.
the elucidation of a key amino acid residue affecting the cyclization specificity of ArsB and ArsC. Surprisingly, only a single point mutation in both enzymes led to the swapping of their main products without a significant loss of their enzymatic activities. The crystal structures of the wild-type and G284W mutant ArsC proteins show that this single amino acid substitution drastically changes the architecture of the active site cavity. Thus our analysis reveals how type III PKSs have evolved to have different cyclization specificity.

EXPERIMENTAL PROCEDURES

Materials—Escherichia coli strains JM109, BL21 (DE3), and B834 (DE3) were purchased from Takara Biochemicals (Ohtsu, Japan). Malonyl-CoA and [2-14C]malonyl-CoA used for the enzyme assay were purchased from Sigma. Behenyl-CoA was prepared as described previously (7).

Site-directed Mutagenesis—Expression plasmids pET16b-ArsB and pET16b-ArsC (7) were used for site-directed mutagenesis. The following primers were used for site-directed mutagenesis with a QuikChange site-directed mutagenesis kit (Stratagene, La Jolla, CA): ArsB W281Y mutant, 5'-ATGACGGGGTTACCGGGGCTCGCAC-3'; ArsC G284W mutant, 5'-ATGACCTGATGACCTGCGAC-3'; ArsB W281L mutant and ArsC G284L mutants, 5'-ATGACCTGATGACCTGCGAC-3'; ArsB W281F and ArsC G284F mutants, 5'-ATGACCTTGTGACCTGCGAC-3'; ArsB W281I and ArsC G284I mutants, 5'-ATGACCTTGTGACCTGCGAC-3'; ArsB W281A and ArsC G284A mutants, 5'-ATGACCTTGTGACCTGCGAC-3'; ArsB W281I and ArsC G284I mutants, 5'-ATGACCTTGTGACCTGCGAC-3'. The following primers and their complementary primers were used for site-directed mutagenesis with a QuikChange site-directed mutagenesis kit (Stratagene, La Jolla, CA): ArsB W281Y mutant, 5'-CGACGAGATGACCTGCGAC-3'; ArsB W281L mutant, 5'-CGACGAGATGACCTGCGAC-3'; ArsB W281F and ArsC G284F mutants, 5'-CGACGAGATGACCTGCGAC-3'; ArsB W281I and ArsC G284I mutants, 5'-CGACGAGATGACCTGCGAC-3'. The mutations were confirmed by determining the nucleotide sequences.

Expression and Purification of Recombinant Enzymes—E. coli strain BL21 (DE3) harboring pET16b-ArsB or pET16b-ArsC was grown at 26 °C overnight in 2 liters of Luria-Bertani broth containing 50 μg/ml ampicillin. For the expression of selenomethionine-substituted ArsC, E. coli strain B834 (DE3) harboring pET16b-ArsC was grown at 26 °C overnight in M9 media supplemented with 50 mg/liter l-selenomethionine. The cells were collected by centrifugation and resuspended in buffer containing 50 mM Tris-HCl (pH 7.8), 300 mM sodium chloride, and 10% (v/v) glycerol. After sonication, the cellular debris was removed by centrifugation, and the supernatant was subjected to nickel-nitrilotriacetic acid superflow resin (Qiagen, Chatsworth, CA). His-tagged enzyme was eluted with a buffer containing 200 mM imidazole. After dialysis, the N-terminal His tag was removed by the treatment with Factor Xa protease. Further purification was performed by Resource Q anion exchange column (GE Healthcare). The purified enzyme was concentrated to 10 mg/ml and stored in a solution containing 10 mM Tris-HCl (pH 7.5), 150 mM sodium chloride, and 10% (v/v) glycerol.

Enzyme Assay—The standard reaction mixture contained 100 μM extender substrate (malonyl-CoA or [2-14C]malonyl-

FIGURE 1. Proposed mechanism for type III PKS reactions. A, phenolic lipid synthesis by two type III PKSs in A. vinelandii. Alkylpyrone formation via an enzyme-bound linear intermediate is depicted, but alkylpyrone can also be formed from the linear intermediate that is released from the enzyme. Two pathways have been suggested for the synthesis of alkylresorcinol by ArsB. In the "hydrolysis first" pathway, aldol cyclization occurs in parallel to decarboxylation (7, 21). In contrast, during the "aldol condensation first" pathway, the cyclization is followed by sequential β-keto decarboxylation and dehydration or by coupled decarboxylation/dehydration (32). B, alkylresorcylic acid synthesis by type III PKSs, including ORAS.

decarboxylative loss of the C1 carbon as CO₂ to yield alkylresorcinols (Fig. 1A). In contrast, ArsC catalyzes intramolecular C5 oxygen-to-C1 lactonization to yield tetraketide pyrones (Fig. 1A). ArsC also uses the triketide intermediate to yield triketide pyrones.

Given that highly homologous *arsB* and *arsC* genes are placed adjacent to each other in the phenolic lipid biosynthetic gene cluster, these genes may have descended from the same ancestor. After duplication, they possibly gain different catalytic properties with just a few amino acid substitutions. Therefore, we assumed that these two Ars enzymes could be ideal targets to obtain an insight into the structural and mechanistic basis for functional diversity of type III PKSs. Here we report
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CoA), 100 μM starter substrate (behenyl-CoA), 100 mM Tris-HCl (pH 8.0), 150 mM sodium chloride, and 6.1 μg of purified enzyme in a total volume of 100 μl. The reaction mixture was incubated at 30 °C for 30 min before being quenched with the addition of 20 μl of 6 M HCl. The products were extracted with ethyl acetate, and the organic layer was then dried out by the evaporator. The products were dissolved in 15 μl of methanol and subjected to radio-thin layer chromatography (TLC) assay, high performance liquid chromatography (HPLC), or liquid chromatography-atmospheric pressure chemical ionization mass spectrometry (LC-APCIMS) analysis. For the radio-TLC assay, Silica gel 60WF254 TLC plate (Merck, Darmstadt, Germany) with a Pegasus-B C4 reversed-phase column (4.6 × 250 mm) (Senshu Scientific, Tokyo, Japan) and 90% acetonitrile in water containing 0.1% acetic acid as an eluent at a flow rate of 1 ml/min. HPLC analysis was performed using the LaChrom Elite system (Hitachi, Tokyo, Japan) with the same column, eluent, and flow rate as for the LC-APCIMS analysis. UV spectra were detected using the Hitachi diode array detector L-2450.

Determination of Kinetic Parameters—The reactions, containing 100 μM malonyl-CoA, 50 mM Tris-HCl (pH 8.0), 150 mM sodium chloride, and 10 μg/ml purified enzyme were performed in a total volume of 900 μl. The concentration of behenyl-CoA was varied between 0.5 and 5 μM. After the preincubation of the reaction mixture at 30 °C for 5 min, the reactions were initiated by the addition of behenyl-CoA and continued for 180 s. The reactions were quenched with the addition of 180 μl of 6 M HCl, and the materials in the mixture were extracted with ethyl acetate. The organic layer was collected and then evaporated. The residual material was dissolved in 20 μl of methanol for HPLC analysis. 5-n-Hexenoylsresorcinol, 4-hydroxy-6-(2’-oxotricosyl)-2-pyryne, and 6-heneicosyl-4-hydroxy-2-pyryne were used to generate standard curves for the quantification of products. Steady-state parameters were determined by a Hanes–wolf plot.

Crystallization, Data Collection, and Structure Determination—Crystals of wild-type ArsC, selenomethionine-labeled ArsC, and the ArsC G284W mutant were grown from a 1:1 mixture of a protein solution (7 mg/ml in 10 mM Tris-HCl (pH 7.5), 150 mM sodium chloride, and 10% (v/v) glycerol) and a reservoir solution (0.1 M imidazole-HCl (pH 7.5), 0.15 M calcium acetate, and 12.5% polyethylene glycol 3350) using the hanging-drop vapor-diffusion method at 20 °C. Before x-ray data collection, the crystal was transferred into the reservoir solution supplemented with 20% (v/v) glycerol as a cryoprotectant and flash-frozen in the liquid nitrogen stream. X-ray data were collected using beamlines BL-5A and AR-NW12A (Photon Factory, Tsukuba, Japan). All data collection was indexed, integrated, and scaled using the program HKL2000 (9). Selenomethionine-labeled protein crystals were used for the single-wavelength anomalous dispersion phasing method. Selenium positions and initial phases were calculated by the program autoSHARP (10). After density modification, clearly interpretable electron density maps were obtained. The program ARP/wARP (11) was used for automatic initial protein model building and refinement. The initial model of ArsC G284W was determined by the molecular replacement method using the program MOLREP (12) with the coordinates of the refined wild-type ArsC structure as a template. Coot (13) was used for visual inspection and manual rebuilding of the model. Refmac (14) was used for refinement. The figures were prepared using PyMOL (15). Each active site volume was calculated using the program CASTp (16). The geometries of the final ArsC structures were evaluated using the program Rampage (17).

RESULTS

In Silico Analysis of Key Residues Affecting Cyclization Diversity between ArsB and ArsC—For the identification of amino acid residues contributing to cyclization diversity between ArsB and ArsC, we first compared the amino acid sequences between ArsB and ArsC on the basis of residues important for functional diversity, as proposed by Austin and Noel (1). The sequence alignment analysis revealed replacement of four amino acid residues in the active site cavity. Ser-157, Leu-219, Thr-235, and Trp-281 of ArsB were replaced with Ala-160, Met-222, Ser-238, and Gly-284 in ArsC (Fig. 2A). To narrow down key residue candidates, we also aligned the sequences with other type III PKS sequences. Besides ArsB and ArsC, several microbial type III PKSs have been identified as alkyresorcylic acid synthases, alkyresorcinol synthases, and alkylpyrone synthases. For example, Mycobacterium tuberculosis PKS18 and Bacillus subtilis BpsA were reported to produce alkylpyrone via lactonization (18, 19). In contrast, Actinoplanes missouriensis AgaA was reported to produce both alkyresorcinol and alkyresorcylic acid via decarboxylative aldol condensation and aldol condensation, respectively (20). Neurospora crassa ORAS (2’-oxoalkylresorcylic acid synthase), Streptomyces griseus SrsaA, and Myxococcus xanthus FtpA were also reported to produce alkyresorcylic acid via aldol condensation (Fig. 1B) (6, 21–24). Trp-281 of ArsB is a conserved aromatic residue among these alkyresorcinol- and/or alkyresorcylic acid-producing type III PKSs although other residues, Ser-157, Leu-219, and Thr-235, show no significant conservation. This Trp residue is replaced by a smaller residue in alkylpyrone-producing enzymes such as PKS18 and ArsC. However, BpsA, which was identified as a pyrone synthase according to in vitro and heterologous expression studies (19), has a Trp residue in this position just like alkyresorcinol- and/or alkyresorcylic acid-producing enzymes. We assume that BpsA is a potential alkyresorcinol- and/or alkyresorcylic acid-producing enzyme. In fact, BpsA was also reported to produce alkyresorcinol, although its main product is alkylpyrone (19). Recently, Rubin-Pitel et al. (6) focused on this aromatic residue of ORAS and showed that the substitution of the aromatic residue Phe by Gly in ORAS altered product profiles from alkyresorcylic acids to alkylpyrones. Therefore, we hypothesized that this residue diversifies cyclization specificity between aldol condensation and lactonization in microbial type III PKSs.

Identification of Trp-281 of ArsB and Gly-284 of ArsC as Key Residues for Their Cyclization Specificity—To prove this hypothesis, we constructed ArsB W281G and ArsC G284W mutants and...
analyzed their reaction products. Radio-TLC analysis showed that G284W substitution of ArsC resulted in the considerable reduction of alkylpyrone production and the concomitant increase of alkylresorcinol production (Fig. 2B). Although alkylpyrones appeared to be produced in a somewhat larger amount in the ArsC G284W reaction than in the ArsB reaction, the ArsC G284W mutant showed a comparable $k_{cat}/K_m$ value for alkylresorcinol synthesis to that for alkylpyrone synthesis catalyzed by wild-type ArsC (Table 1). This result showed that G284W mutation altered the cyclization specificity of ArsC from lactic acid.
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Cyclization to aldol condensation without a significant loss of the polyketide chain elongation activity. In contrast, W281G substitution of ArsB resulted in the complete loss of alkylresorcinol production and the simultaneous increase of alkylpyrone production (Fig. 2B). We note that triketide pyrone was produced in a larger amount than tetraketide pyrone in the ArsB W281G reaction, whereas triketide and tetraketide pyrones were produced in a similar amount in the ArsC reaction. This result is in agreement with the steady-state kinetic parameters. Our "reciprocal" mutational analysis of ArsB and ArsC (which produced ArsB-W281G and ArsC-G284W) clearly demonstrates that Trp-281 of ArsB or Gly-284 of ArsC occupies a critical position for cyclization of polyketomethylene intermediates.

Next, we substituted Trp-281 of ArsB and Gly-284 of ArsC to other aromatic and aliphatic residues (Phe, Tyr, Val, Ile, Leu, and Ala) to explore the effect of steric and aromatic properties of the amino acid residue at this position (Fig. 3). The ArsB W281F and W281Y mutants retained alkylresorcinol-producing activity and did not produce alkylpyrone. The ArsB W281L mutant produced alkylpyrone as well as alkylresorcinol. Other ArsB mutants such as W281V, W281I, and W281A severely decreased alkylresorcinol-producing activity, and their major products were alkylpyrone, as observed for the ArsB W281G mutant. These results indicate that alkylresorcinol/alkylpyrone production roughly depends on the size of the amino acid residue at position 281 in ArsB. The presence of a bulky amino acid residue seems to allow the generation of alkylresorcinol and exclude the production of alkylpyrone in ArsB. In contrast, in the case of ArsC, mutants other than G284W, which was substituted to Phe, Tyr, Val, Ile, Leu, and Ala, did not acquire the alkylresorcinol-producing activity. Thus, some substitution pairs such as ArsB W281Y and ArsC G284Y mutants did not share product profiles. This inconsistency may be explained by the other three amino acid variations in the active site cavities of ArsB and ArsC, as described above (see “Discussion”).

Overall Structures of ArsC and ArsC G284W—We have determined the crystal structure of the wild-type and G284W mutant ArsC proteins at 1.76 and 1.99 Å resolution, respectively (Table 2). The overall structure of the ArsC G284W mutant is almost identical to that of the wild-type enzyme (root mean square deviation (r.m.s.d.) of Ca atoms between the A chain of the wild-type protein and the A chain of the G284W mutant is 0.21 Å). Each of the ArsC structures has four dimers per crystallographic asymmetric unit. The ArsC structure adopts the conserved homodimeric αβαβα thiolase fold observed in other crystal structures of type III PKSs (Fig. 4A). A search for structurally related proteins using the program Dali (25) revealed that the closest protein is Dictyostelium discoideum Steely, a hybrid enzyme composed of a type I fatty acid synthase and a type III PKS (26) (Z score = 43.0, r.m.s.d. of 2.2 Å, sequence identity of 21%). We note that ArsC also exhibits structural similarity with PKS18 (18) (Z score = 41.8, r.m.s.d. of 2.3 Å, sequence identity of 20%) and ORAS (6, 21) (Z score = 40.7, r.m.s.d. of 2.4 Å, sequence identity of 17%). Thus ArsC is structurally similar to other type III PKSs; however, the N-terminal 120-residue region of ArsC shows structural deviations. The
unique structural element is the loop-insertion region (84–110) between helix α3 and helix α5 (Fig. 4A). This loop is a disordered region in five of the eight molecules in the crystallographic asymmetric unit, indicating that this region is flexible.

Structure of the Substrate-binding Site of ArsC—Previous structural studies on plant type III PKSs, alfalfa chalcone synthase (27) and daisy 2-pyrone synthase (4), revealed that several residues of the helix α3 are involved in interactions with two phosphates of CoA at the entrance of the CoA binding tunnel, suggesting that helix α3 plays an important role in CoA substrate binding. In microbial type III PKSs using acyl-CoA substrates, helix α3 occupies a similar position (Fig. 5). Even Steely, which accepts acyl-ACP substrates, has helix α3 at the equivalent position (Fig. 5). However, in the ArsC structure, helix α3 shifts away from the CoA-binding site by 8 Å (Fig. 5), resulting in the rearrangement of the CoA binding region. Although ArsB and ArsC were shown to react with the acyl-CoA substrate in vitro, they are supposed to interact with the ACP domain of ArsA in vitro. For example, the acyl binding tunnel of PKS18 extends out to the surface of the protein (18). In the ORAS structure, the tunnel does not extend to solvent and is inwardly directed into the protein (6). ArsC has a straight acyl binding tunnel extending toward the surface of the protein, as observed for PKS18. However, the unique flexible loop between helix α3 and helix α5 forms the base of this acyl binding tunnel (Fig. 6). The side chains of Phe-95 and Ile-97 in this flexible loop block solvent access to the acyl binding tunnel. This flexible loop may function as a lid during substrate binding.

Structural Basis for the Cyclization Specificity—The active site cavity containing the catalytic triad Cys-186–His-334–Asn-367 is placed inside the ArsC molecule, similar to those of other type III PKSs. Gly-284, the key residue for cyclization specificity, is located at the center of the active site cavity in the ArsC structure (Fig. 7A). The active site cavity extends beyond the Gly-284 position and reaches to the dimeric interface. Ala-160 of ArsC participates in the formation of the active site cavity of the neighboring monomer. Plant type III PKSs have a similar cross-subunit interaction. Met-137 of chalcone synthase corresponding to Ala-160 of ArsC is highly conserved in plant type III PKSs and makes cross-subunit interactions by protruding into the second monomer and forming the base of the active site cavity (27). Chalcone synthase also has a small residue, Gly-256, at the equivalent position as Gly-284 of ArsC, enabling the opening of the active site cavity to the dimeric interface.

In the structure of the ArsC G284W mutant, the bulky side chain of the mutated residue protrudes toward the space of the active site cavity and blocks the connection to the dimeric interface (Fig. 7A). Thus, this substitution causes a significant reduction of cavity volume from 592 Å3 in wild-type ArsC to 293 Å3 in the ArsC G284W mutant (Fig. 8, A and B). This change of

### Table 2
Data collection, phasing, and refinement statistics

| Datasets | Wild type | G284W mutant | SeMet* |
|----------|-----------|---------------|--------|
| Data collection statistics | | | |
| Beamline | PF BL-5A | PF AR-NW12A | PF BL-5A |
| Wavelength (Å) | 1.00000 | 1.00000 | 0.97915 |
| Space group | P2₁ | P2₁ | P2₁ |
| Unit-cell parameters | | | |
| a (Å) | 102.32 | 102.89 | 102.53 |
| b (Å) | 143.39 | 142.76 | 143.55 |
| c (Å) | 129.62 | 129.77 | 129.92 |
| β (degree) | 110.33 | 110.45 | 110.20 |
| Resolution (Å) | 50.00-1.76 | 50.00-1.99 | 50.00-2.53 |
| Unique reflections | 345,492 (37,894) | 244,208 (15,807) | 115,095 (9,308) |
| Redundancy | 3.8 (2.7) | 5.4 (4.8) | 7.4 (7.2) |
| Completeness (%) | 97.2 (80.3) | 96.9 (61.6) | 99.3 (96.8) |
| Rmerge (%) | 0.120 (0.546) | 0.140 (0.821) | 0.086 (0.114) |
| Mean (I/σ(I)) | 21.3 (2.2) | 19.9 (1.9) | 39.4 (22.5) |
| Refinement statistics | | | |
| Rfree (%) | 18.3 | 20.5 |
| Rfree (%) | 23.0 | 27.3 |
| No. of protein atoms | 24,691 | 24,661 |
| No. of waters | 2,676 | 2,369 |
| No. of sodium ions | 8 | 8 |
| r.m.s.d. | | | |
| Bond lengths (Å) | 0.027 | 0.023 |
| Bond angles (°) | 2.110 | 1.872 |
| Ramachandran plot | | | |
| Favored region (%) | 96.6 | 96.1 |
| Allowed region (%) | 3.3 | 3.5 |
| Outer region (%) | 0.1 | 0.4 |

* ArsC selenomethionine (SeMet) derivative data were used only for initial phase determination.
cavity volume does not affect the elongation reaction with malonyl-CoA but strikingly affects the cyclization reaction. This substitution also gives rise to the following three minor structural arrangements. First, the polypeptide backbone between Trp-284 and Pro-286 moves 0.8 Å toward the catalytic triad. Second, the presence of the bulky side chain of Trp-284 causes a conformational movement of Met-293 (Fig. 7B). The side chain of Met-293 swings by 40° toward the acyl binding tunnel, thereby narrowing the entrance of the acyl binding tunnel. Third, the presence of the aromatic side chain affects the positioning of water molecules. In the structure of wild-type ArsC, a water molecule network is present in the active site cavity, and Ser-369 holds a water molecule with a hydrogen bond (Fig. 9A). Judging from the short distance (3.7 Å) between this water molecule and the catalytic cysteine, this water molecule possibly plays an important role in catalysis. However, in the structure of the ArsC G284W mutant, the water molecule network is disrupted by the bulky side chain of Trp-284, and Ser-369 loses the ability to retain the water molecule (Fig. 9B).

Although the overall sequence identity is relatively low between the ArsC G284W mutant and ORAS (sequence identity, 17%) or between wild-type ArsC and PKS18 (sequence identity, 20%), their cyclization manners are similar to each other (7, 21). To find out whether alkylresorcinol synthases and alkylpyrone synthases have a common structural property, we compared the active site cavities of ArsC with other type III PKSs. The structural comparison with ORAS revealed that ORAS has a slightly larger cavity volume (392 Å³) than the ArsC
G284W mutant (293 Å³). This may enable ORAS to catalyze an additional fourth polyketide extension step for the synthesis of pentaketide compounds. ORAS has an aromatic residue, Phe-252, corresponding to Trp-284 in the ArsC G284W mutant (Figs. 7C and 8, A and C). The position and orientation of Trp-284 in the structure of the ArsC G284W mutant are well superimposed with those of the ArsC G284W mutant structure. Residues of wild-type ArsC and the G284W mutant are shown in cyan and green sticks, respectively. B, the superimposed structures of wild-type ArsC and the ArsC G284W mutant. The whole A chain atoms of the wild-type ArsC structure were superimposed onto those of the ArsC G284W mutant structure. Residues of wild-type ArsC and the G284W mutant are shown in cyan and green sticks, respectively. C, the superimposed structures of ArsC G284W and ORAS. Residues of the ArsC G284W mutant and ORAS (PDB code 3EUO) are shown in green and yellow sticks, respectively. D, the superimposed structures of wild-type ArsC with PKS18. Residues of wild-type ArsC and PKS18 (PDB code 1TED) are shown in cyan and magenta sticks, respectively.

DISCUSSION

Crystal structures of many type III PKSs have been determined. As for microbial type III PKSs, structures of PKS18 and ORAS were determined as a pyrone synthase and a resorcylic acid synthase, respectively (6, 18, 22). However, no structure of a microbial resorcinol synthase has been available. In this study we succeeded in the determination of structures of wild-type ArsC (pyrone synthase) and an ArsC G284W mutant ("artifi-
Because the amino acid sequence of ArsB (resorcinol synthase) is highly homologous to that of the ArsC G284W mutant, the structure of the ArsC G284W mutant could provide important insights into the catalytic mechanism of microbial resorcinol-producing type III PKSs, as described below.

Both lactonization and aldol condensation reactions should start from the elimination of an active methylene proton.
between two carbonyl groups to form an enolate anion. In the lactonization reaction, the resulting anion at the oxygen directly attacks a carbonyl group. In the case of aldol condensation, the active anion reacts with a carbonyl group via the olefinic carbon to form a C–C bond. Our mutagenic and structural studies clearly showed that Trp-281 of ArsB and Gly-284 of ArsC should occupy a critical position for the cyclization specificity of each enzyme. Trp-281 appears to be important in providing a steric wall for the proper folding of the polyketomethylene intermediate to aldol condensation in ArsB. The existence of an aromatic group may restrict the conformation of the growing polyketide chain and probably excludes spontaneous lactonization by preventing the oxygen of enolate from getting close enough to react with a carbonyl group. Thus, Trp-281 in ArsB properly controls the number of condensation reactions during the polyketide extension and the conformation of the intermediates during the cyclization so that ArsB produces only tetraketide alkylresorcinol. In contrast, ArsC, which has Gly-284 at the equivalent position, produces triketide and tetraketide alkylpyrones via lactonization. The large cavity of ArsC may give flexibility to the polyketomethylene intermediate chain, giving a derailment product. Therefore, ArsC cannot exclude the formation of a triketide product by spontaneous cyclization caused from the flexibility of the triketide intermediate, allowing the formation of both triketide and tetraketide pyrones. Thus the condensation and cyclization steps seem not to be strictly controlled in the ArsC reaction. We assume that *arsB* and *arsC* genes may have descended from the same ancestor and that ArsB and ArsC probably acquired different catalytic properties by amino acid substitutions. Replacement of Gly by Trp (at position 281) in ArsB or replacement of Trp by Gly (at position 284) in ArsC would be critical evolutionarily events to generate a different specificity in cyclization (aldol condensation or lactonization). It is difficult to predict, however, the original cyclization specificity of the ancestral enzyme.

We showed that substitution of Trp-281 by Gly in ArsB altered its cyclization specificity from aldol condensation to lactonization to produce triketide and tetraketide pyrones, which are common derailment products of PKSs. Several similar results in which pyrone-producing activity was conferred on type III PKSs by amino acid substitution have been previously reported (4, 6, 29, 30). To date, however, there has been no report on the alteration of cyclization specificity of a type III PKS from lactonization to aldol condensation. In this context, there is only one report to confer a resorcinol-producing activity on a plant type III PKS; mutation of eight amino acid residues was required for altering the cyclization specificity of alfalfa chalcone synthase from a chalcone-producing Claisen condensation to a stilbene (resorcinol)-producing aldol condensation (5). In the present study we showed for the first time that substitution of Gly-284 by Trp in ArsC altered its cyclization specificity from lactonization to aldol condensation. It should be emphasized that only one amino acid residue is a critical determinant of different cyclization specificity between ArsB and ArsC.

As described above, our results clearly showed that Trp-281 of ArsB plays a critical role in aldol condensation. There are other microbial type III PKSs that catalyze aldol condensations that also have aromatic residues such as Trp and Phe at the corresponding position. Thus our results contribute to prediction of cyclization specificity based on amino acid sequence of microbial type III PKSs. However, our results also suggest that differences in three other amino acid residues in the active site cavity (Ser-157, Leu-219, and Thr-235 in ArsB and Ala-160, Met-222, and Ser-238 in ArsC) caused several differences in catalytic properties between recombinant (wild-type or mutant) ArsB and ArsC enzymes that have the same amino acid residue at the critical position (281 of ArsB and 284 of ArsC). First, the enzymatic properties of ArsC G284W differed from those of ArsB; ArsC G284W produced resorcinol as the major product with a small, but significant amount of pyrone, whereas ArsB produced resorcinol with trace amounts of pyrone. This result suggests that ArsC G284W is somewhat inferior to ArsB as a resorcinol synthase and indicates that the shape of the active site cavity of ArsC G284W is less suited for aldol condensation than that of ArsB. Second, the enzymatic properties of ArsB W281G also differed from those of ArsC; ArsB W281G produced ~4-fold greater amounts of triketide pyrone than tetraketide pyrone, whereas ArsC produced both triketide and tetraketide pyrones in comparable amounts. This indicates that the shape of the active site cavity of ArsC G284W is not as ideally suited to the three rounds of chain-elongation compared to the active site of ArsC. Third, some of the substitution pairs between ArsB and ArsC, such as ArsB W281Y and ArsC G284Y mutants, did not share product profiles; ArsB W281Y and W281F produced alkylresorcinol, but ArsC G284Y and G284F produced alkylpyrone but not alkylresorcinol. In other words, when the amino acid residue at position 281 is an aromatic residue, the resorcinol synthase activity is retained in the alkylresorcinol synthase ArsB. In contrast, in the case of substitution of Gly-284 in the alkylpyrone synthase ArsC, only Trp, but not Tyr or Phe, can confer the resorcinol synthase activity to ArsC. This result also implies the entire shape of the active site cavity of ArsC G284W is less suited for the aldol condensation than the active site cavity of ArsB. All three results suggest that not only is one amino acid residue (for example, Trp-281 of ArsB and Gly-284 of ArsC) critical but also other amino acid residues in the active site cavity are important and can affect the enzymatic properties including cyclization specificity of microbial type III PKSs. This notion should be considered in the sequence-based prediction of cyclization specificity of microbial type III PKSs. In fact, BpsA mainly produces alkylpyrone, not alkylresorcinol, even though it has a Trp residue at the critical position (19). This indicates the shape of the active site cavity of BpsA is also not likely ideal for aldol condensation.

As for the plant type III PKS, Austin et al. (5) determined the structure of resorcinol-producing stilbene synthase (STS) and proposed the reaction mechanism. STS is proposed to employ an “aldol switch” mechanism with the thioesterase-like hydro- gen bond network composed of Thr-132, Glu-192, and Ser-338, allowing the activation of a water molecule for the hydrolysis of the C1-cysteine thioester bond. The resulting free carboxylic acid intermediate is cyclized via aldol condensation and then decarboxylated to form a resorcinol product. However, the
structure of the ArsC G284W mutant suggests that ArsB does not use such a reaction mechanism. ArsB contains Cys-155 at the position corresponding to Thr-132 in STS. The presence of this Cys residue hinders the formation of the hydrogen bond network in the ArsC G284W mutant. ORAS also has Cys-120 at the corresponding position (6). Moreover, in the ArsC G284W mutant, no water molecule for the hydrolysis of the thioester bond is present adjacent to Ser-369, which is equivalent to Ser-338 in STS. Thus, the aldol switch network is not conserved in microbial resorcinol- and resorcylic acid-producing type III PKSs. Another difference is that STS does not have a steric wall, as observed in ArsB. Trp-281 of ArsB is replaced by Gly in STS.

In other plant alkyresorcylic acid synthases, *Oryza sativa* ARAS1 and ARAS2 (31), this aromatic residue is replaced by Met. Consequently, several amino acid variations and structural differences are observed between microbial and plant aldol condensation-catalyzing type III PKSs, indicating that different reaction mechanisms are employed. In the reaction catalyzed by plant type III PKSs, the aldol switch triggers the formation of an acidic intermediate by cleavage of the thioester bond of the Cys-bound linear intermediate. The acidic intermediate is then cyclized via aldol condensation (5). This study and that of the ORAS structure (6) indicate, however, that microbial type III PKSs do not make use of the aldol switch mechanism but instead provide a steric wall so that polyketomyelene intermediates may be folded into suitable forms for cyclization via aldol condensation. Interestingly, Posehn et al. (32) suggested that aldol cyclization catalyzed by ArsB occurs before hydrolysis of the thioester bond. The identification of amino acid residues directly responsible for aldol condensation in ArsB and the elucidation of the molecular mechanism for the reaction are our future challenges.

In conclusion, we showed that substitution of a single amino acid resulted in the alteration of the cyclization specificity of two *Azotobacter* type III PKSs, ArsB and ArsC; Trp-281 of ArsB and Gly-284 of ArsC are crucial residues for the specificity of cyclization. Structural analysis revealed that the volume of the active site cavity was a crucial determinant of the cyclization specificity of each enzyme. Thus this study provides a vital element for predicting cyclization specificity of microbial type III PKSs on the basis of their amino acid sequence. It is noteworthy that we succeeded for the first time in altering the cyclization specificity of a type III PKS from lantonization to aldol condensation by replacement of only a single amino acid residue.

REFERENCES

1. Austin, M. B., and Noel, J. P. (2003) The chalcone synthase superfamily of type III polyketide synthases. *Nat. Prod. Rep.* **20**, 79–110

2. Katsuyama, Y., and Horinouchi, S. (2010) *Comprehensive Natural Products II: Chemistry and Biology*, Vol. 1, pp. 147–170, Elsevier Science Publishing Co., Inc., New York

3. Morita, H., Abe, I., and Noguchi, H. (2010) *Comprehensive Natural Products II: Chemistry and Biology*, Vol. 1, pp. 171–225, Elsevier Science Publishing Co., Inc., New York

4. Jez, J. M., Austin, M. B., Ferrer, J., Bowman, M. E., Schröder, J., and Noel, J. P. (2000) Structural control of polyketide formation in plant-specific polyketide synthases. *Chem. Biol.* **7**, 919–930

5. Austin, M. B., Bowman, M. E., Ferrer, J. L., Schröder, J., and Noel, J. P. (2004) An aldol switch discovered in stilbene synthases mediates cyclization specificity of type III polyketide synthases. *Chem. Biol.* **11**, 1179–1194

6. Rubin-Pitel, S. B., Zhang, H., Vu, T., Brunzelle, J. S., Zhao, H., and Nair, S. K. (2008) Distinct structural elements dictate the specificity of the type III pentaketide synthase from *Neurospora crassa*. *Chem. Biol.* **15**, 1079–1090

7. Funa, N., Ozawa, H., Hirata, A., and Horinouchi, S. (2006) Phenolic lipid synthesis by type III polyketide synthases is essential for cyst formation in *Azotobacter vinelandii*. *Proc. Natl. Acad. Sci. U.S.A.* **103**, 6536–6531

8. Miyanaga, A., Funa, N., Awakawa, T., and Horinouchi, S. (2008) Direct transfer of starter substrates from type I fatty acid synthase to type III polyketide synthases in phenolic lipid synthesis. *Proc. Natl. Acad. Sci. U.S.A.* **105**, 871–876

9. Vonotinos, Z., and Minor, W. (1997) Processing of x-ray diffraction data collected in oscillation mode. *Methods Enzymol.* **276**, 307–326

10. Morris, R. J., Perrakis, A., and Lamzin, V. S. (2002) ARP/wARP’s model-building algorithms. I. The main chain. *Acta Crystallogr. B. Biol. Crystallogr.* **58**, 960–975

11. Morris, R. J., Perrakis, A., and Lamzin, V. S. (2002) ARP/wARP’s model-building algorithms. II. The side chain. *Acta Crystallogr. B. Biol. Crystallogr.* **58**, 976–986

12. Vagin, A., and Teplyakov, A. (1997) MOLREP: an automated program for molecular replacement. *J. Appl. Crystallogr.* **30**, 1022–1025

13. Emsley, P., and Cowtan, K. (2004) Coot: Model-building tools for molecular graphics. *Acta Crystallogr. D. Biol. Crystallogr.* **60**, 2126–2132

14. Murshudov, G. N., Vagin, A. A., and Dodson, E. J. (1997) Refinement of macromolecular structures by the maximum-likelihood method. *Acta Crystallogr. D. Biol. Crystallogr.* **53**, 240–255

15. Dundas, J., Ouyang, Z., Tseng, J., Binkowski, A., Turpaz, Y., and Liang J. (2006) CASTp: Computed atlas of surface topography of proteins with structural and topographical mapping of functionally annotated residues. *Nucleic Acids Res.* **34**, W116–W118

16. Lovell, S. C., Davis, I. W., Arendall, W. B., 3rd, de Bakker, P. I., Word, J. M., Prisant, M. G., Richardson, J. S., and Richardson, D. C. (2003) Structure validation by Ca geometry. φ, ψ, and CB deviation. *Proteins* **50**, 437–450

17. Sankaranarayanan, R., Saxena, P., Marathe, U. B., Gokhale, R. S., Shanmugam, V. M., and Rukmini, R. (2004) A novel tunnel in mycobacterial type III polyketide synthase reveals the structural basis for generating diverse metabolites. *Nat. Struct. Mol. Biol.* **11**, 894–900

18. Nakano, C., Ozawa, H., Akanuma, G., Funa, N., and Horinouchi, S. (2009) Biosynthesis of aliphatic polyketides by type III polyketide synthase and methyltransferase in *Bacillus subtilis*. *J. Bacteriol.* **191**, 4916–4923

19. Awakawa, T., Fujita, N., Hayakawa, M., Ohnishi, Y., and Horinouchi, S. (2011) Characterization of the biosynthesis gene cluster for alkyl-O-dihydrogeranyl-methoxyhydroquinones in *Actinoplanes missouriensis*. *ChemBioChem* **12**, 439–448

20. Funa, N., Awakawa, T., and Horinouchi, S. (2007) Pentaketide resorcylic acid synthesis by type III polyketide synthase from *Neurospora crassa*. *J. Biol. Chem.* **282**, 14476–14481

21. Goyal, A., Saxena, P., Rahman, A., Singh, P. K., Kasbekar, D. P., Gokhale, R. S., and Sankaranarayanan, R. (2008) Structural insights into biosynthesis of resorcinolic lipids by a type III polyketide synthase in *Neurospora crassa*. *J. Struct. Biol.* **162**, 411–421

22. Funabashi, M., Funa, N., and Horinouchi, S. (2008) Phenolic lipids synthesized by type III polyketide synthase confer penicillin resistance on *Streptomyces griseus*. *J. Biol. Chem.* **283**, 13983–13991

23. Hayashi, T., Kitamura, Y., Funa, N., Ohnishi, Y., and Horinouchi S. (2011) Fatty acyl-AMP ligase involvement in the production of alkylresorcylic acid by a *Mycococcus sunthus* type III polyketide synthase. *ChemBioChem* **12**, 2166–2176

24. Holm, L., and Sander, C. (1995) Dali. A network tool for protein structure comparison. *Trends Biochem. Sci.* **20**, 478–480

25. Austin, M. B., Saito, T., Bowman, M. E., Haydock, S., Kato, A., Moore, B. S., Kay, R. R., and Noel, J. P. (2006) Biosynthesis of *Dictyostelium discoideum* differentiation-inducing factor by a hybrid type I fatty acid-type III polyketide synthase. *Nat. Chem. Biol.* **2**, 494–502

26. Ferrer, J. L., Jez, J. M., Bowman, M. E., Dixon, R. A., and Noel, J. P. (1999) Structure of chalcone synthase and the molecular basis of plant polyketide biosynthesis. *Nat. Struct. Biol.* **6**, 775–784
28. Austin, M. B., Izumikawa, M., Bowman, M. E., Udwary, D. W., Ferrer, J. L., Moore, B. S., and Noel, J. P. (2004) Crystal structure of a bacterial type III polyketide synthase and enzymatic control of reactive polyketide intermediates. *J. Biol. Chem.* **279**, 45162–45174

29. Funa, N., Ohnishi, Y., Ebizuka, Y., and Horinouchi, S. (2002) Alteration of reaction and substrate specificity of a bacterial type III polyketide synthase by site-directed mutagenesis. *Biochem. J.* **367**, 781–789

30. Shimokawa, Y., Morita, H., and Abe, I. (2010) Structure-based engineering of benzalacetone synthase. *Bioorg. Med. Chem. Lett.* **20**, 5099–5103

31. Matsuzawa, M., Katsuyama, Y., Funa, N., and Horinouchi, S. (2010) Alkylresorcylic acid synthesis by type III polyketide synthases from rice *Oryza sativa*. *Phytochemistry* **71**, 1059–1067

32. Posehn, S. E., Kim, S. Y., Wee, A. G. H., and Suh, D.-Y. (2012) Mapping the mechanism of the resorcinol ring formation catalyzed by ArsB, a type III polyketide synthase from *Azotobacter vinelandii*. *ChemBioChem* **13**, 2212–2217