Research Article

Structural and functional studies of SAV0551 from Staphylococcus aureus as a chaperone and glyoxalase III

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The DJ-1/ThiJ/PfpI superfamily of proteins is highly conserved across all biological kingdoms showing divergent multifunctions, such as chaperone, catalase, protease, and kinase. The common theme of these functions is responding to and managing various cellular stresses. DJ-1/ThiJ/PfpI superfamily members are classified into three subfamilies according to their quaternary structure (DJ-1-, YhbO-, and Hsp-types). The Hsp-type subfamily includes Hsp31, a chaperone and glyoxalase III. SAV0551, an Hsp-type subfamily member from Staphylococcus aureus, is a hypothetical protein that is predicted as Hsp31. Thus, to reveal the function and reaction mechanism of SAV0551, the crystal structure of SAV0551 was determined. The overall folds in SAV0551 are similar to other members of the Hsp-type subfamily. We have shown that SAV0551 functions as a chaperone and that the surface structure is crucial for holding unfolded substrates. As many DJ-1/ThiJ/PfpI superfamily proteins have been characterized as glyoxalase III, our study also demonstrates SAV0551 as a glyoxalase III that is independent of any cofactors. The reaction mechanism was evaluated via a glyoxylate-bound structure that mimics the hemithioacetal reaction intermediate. We have confirmed that the components required for reaction are present in the structure, including a catalytic triad for a catalytic action, His78 as a base, and a water molecule for hydrolysis. Our functional studies based on the crystal structures of native and glyoxylate-bound SAV0551 will provide a better understanding of the reaction mechanism of a chaperone and glyoxalase III.

Introduction

The DJ-1/ThiJ/PfpI superfamily (DJ-1 superfamily hereinafter) is a group of proteins that are universal to all the kingdoms of life [1]. The members of this superfamily share structural and functional similarity to human DJ-1, which is known to be associated with Parkinson disease [2,3]. The crystal structures of several members of the DJ-1 superfamily have been solved. Most of them are oligomers, with each monomer consisting of center-aligned β-strands surrounded by α-helices, which is called an α/β sandwich fold [4]. A completely conserved cysteine is located at a sharp turn between a β-strand and an α-helix. The sharp turn motif is referred to as the ‘nucleophilic elbow’ and is known to allow easy access to the cysteine by substrates [5]. DJ-1 superfamily proteins are divided into the following distinct subfamilies based on their quaternary structure: (i) DJ-1-type, (ii) YhbO-type, and (iii) Hsp-type. Each of the DJ-1 subfamilies also differs in the architecture of the conserved cysteine. DJ-1-type subfamily proteins (DJ-1 and YajL) form dimeric interfaces consisting of α-helices, β-strands, and loops. The conserved cysteine does not form a catalytic triad in human DJ-1 and bacterial YajL [6,7]. Many functions of DJ-1 have been revealed including chaperone, neurone protection, and glyoxalase III [8-10]. The YhbO-type subfamily includes...
the representative YhbO from *Escherichia coli*, SAV1875 from *Staphylococcus aureus*, *PfpI* from *Pyrococcus horikoshii*, and Ton1285 from *Thermococcus onnurineus*. YhbO-type subfamily members form dimers with three α-helices. The catalytic triad in this subfamily consists of a cysteine, histidine, and an acidic residue from the adjacent subunit. Ton1285 and *PfpI* from the YhbO-type subfamily are known to cleave peptide bonds [4,11]. Hsp-type subfamily members have a unique dimerization surface consisting of N-terminal β-strands and loops [12]. As heat shock proteins (HSPs) are considered molecular chaperones, most members of the Hsp-type subfamily function as chaperones, specifically holding chaperones [13]. Holding chaperones capture unfolded proteins with their surface structures for proper folding. The discriminating surface structures, canyons, and bowls of Hsp31 from *E. coli* and *Vibrio cholerae* distinguish them from other subfamily members [12,14]. Hsp-type DJ-1 subfamily proteins have a catalytic triad consisting of a cysteine, histidine, and an acidic residue from the cap domain of the same subunit. With the catalytic cysteine in the catalytic triad, most proteins in the Hsp-type subfamily function as glyoxalase III [15-18].

The distinct subclass classification correlates with structural characteristics, but known functions of DJ-1 superfamily proteins cannot be classified into subclasses. From our previous study, we determined the surface structure of the YhbO-type protein SAV1875 differs from the YhbO-type subfamily proteins but is similar to Hsp-type subfamily proteins. SAV1875 shows chaperone activity similar to other Hsp-type subfamily proteins [19]. However, the Hsp-type protein Glx3 from *Candida albicans* and YDR533C from *Saccharomyces cerevisiae* do not show chaperone activity. The differences in the chaperone activity within the Hsp-type subfamily are shown to be based on the importance of their surface structure, as these two proteins lack the 45 amino acid N-terminal region [16]. The glyoxalase III function was first revealed in Hsp-type subfamily proteins, though YajL and DJ-1 from the DJ-1-type and YhbO from the YhbO-type subfamily also show the same function [9,20]. Despite the structural and functional differences in the classes, many DJ-1 superfamily proteins perform chaperone and glyoxalase III functions. Under stress conditions, chaperone proteins protect cells by stabilizing unfolded proteins, giving the cell time to repair or resynthesize damaged proteins, which is important for protecting cells from severe stress conditions [21]. Glyoxalase III directly converts the toxic methylglyoxal substrate to non-toxic β-lactate without glutathione. Methylglyoxal is a toxic endogenous electrophile component that damages proteins, nucleic acids, and lipids [22]. Therefore, studying the DJ-1 superfamily of proteins is important for understanding cellular protection mechanisms from various stress conditions.

In the present study, the structural and functional identification of hypothetical SAV0551 from *S. aureus* Mu50 was conducted. *S. aureus* Mu50 (ATCC 700699) is a vancomycin-intermediate *S. aureus* that displays robust virulence properties. It causes severe infectious diseases ranging from mild infections, such as skin infections and food poisoning, to life-threatening infections, such as sepsis, endocarditis, and toxic shock syndrome [23]. However, because of the bacterial resistance, the prognosis for *S. aureus* infection is still poor despite early diagnosis and appropriate treatment [24]. Although there are various theories regarding how *S. aureus* acquires antibiotic resistance, very little is known. In this regard, understanding the bacterial defense mechanisms in *S. aureus* against antibiotics is crucial. Furthermore, the DJ-1 superfamily of proteins is predicted to have stress-response functions under multiple stress conditions [25]. Therefore, we designed structural and functional experiments with SAV0551, a DJ-1 superfamily protein from *S. aureus* Mu50, as an extension of our previous work on SAV1875. We elucidated the structure and function of SAV0551 as a chaperone and glyoxalase III. The catalytic triad mutants (C190A, H191A, and D221A) and absolutely conserved glutamate mutant (E81A) were designed to identify structure-related functional differences. The mutants lost glyoxalase III activity but maintained chaperone activity. We show a co-crystal structure of SAV0551 with the known glyoxalase III inhibitor glyoxylate, and it shows the intermediate structure in the glyoxalase III reaction. Our structural and functional research on previously unknown SAV0551 presents new residues that are involved in glyoxalase III reaction.

**Experimental**

**Cloning, protein expression, and purification**

The predicted ORF of SAV0551 was amplified from *S. aureus* Mu50 genomic DNA using standard PCR methods. The forward and reverse oligonucleotide primers were designed using the published genome sequence, 5′-GACCTGCATATGTCACAAAGATGTAATGAATTAAG and 5′-GTCACCTGAGTTATTTTGTATTGCATTAAAATCAT, respectively, where the bases underlined represent the Ndel and Xhol restriction enzyme cleavage sites. The amplified DNA was inserted into the Ndel/Xhol-digested expression vector pET-21a (+) (Novagen). The resulting construct contains eight non-native residues at the C-terminus (LEHHTHHH) that facilitate protein purification. The accuracy of the cloning was confirmed by DNA sequencing. The resulting expression plasmid was then transformed into *E. coli* BL21(DE3) cells (Novagen).
To prepare mutants, the EZchange Site-directed Mutagenesis kit (Enzymonics) was used to generate point mutations in the SAV0551 recombinant pET-21a(+) plasmid. The point mutations resulted in separate multiple recombinant plasmids, specifically E81A, C190A, H191A, and D221A. The sequences of the reconstructed mutants were confirmed by DNA sequencing (results not shown).

The wild-type and SAV0551 mutants’ (E81A, C190A, H191A, and D221A) cells were grown at 37°C until the OD600 reached 0.6 and expression was induced by the addition of IPTG to a final concentration of 0.5 mM. After an additional 4 h of growth at 37°C, cells were harvested by centrifugation and resuspended in 50 mM Tris, pH 7.5, 0.5 M NaCl, and 20 mM imidazole buffer. Cells were lysed by sonication at 4°C and the supernatant was loaded onto Ni2+-NTA (Ni2+-nitrilotriaceticate) affinity column (Qiagen; 3 ml of resin per liter of cell culture) previously equilibrated with the same buffer. The column was washed extensively with wash buffer (50 mM Tris, pH 7.5, 0.5 M NaCl, and 50 mM imidazole); then the bound protein was eluted with elution buffer (50 mM Tris, pH 7.5, 0.5 M NaCl, and 500 mM imidazole) until there was no detectable absorbance at 280 nm in the eluant. Fractions containing protein were concentrated to ~2 ml and applied to a Superdex 75 (10/300 GL) column (GE Healthcare Life Sciences) that had been equilibrated with the final buffer (50 mM Tris, pH 7.5 and 0.15 M NaCl). The purities of SAV0551 were judged to be over 95% by SDS/PAGE. The protein solution was concentrated using 10000 Da molecular-mass cut-off spin columns (Millipore). The protein concentration was estimated by measuring the absorbance at 280 nm, employing the calculated extinction coefficient of 35535 M⁻¹·cm⁻¹ (Swiss-Prot; http://www.expasy.org).

### Crystallization, data collection, structure determination, and refinement

Crystallization was performed at 293 K by the hanging-drop vapor diffusion method using 24-well VDX plates (Hampton Research). Initial crystallization conditions were established using screening kits from Hampton Research (Crystal Screens I and II, Index, PEG/Ion, and MembFac) and from Emerald BioSystems (Wizard I, II, III, and IV). For the optimal growth of the SAV0551 crystals, each hanging drop was prepared on a siliconized cover slip by mixing 1 μl of protein solution (15 mg/ml) and 1 μl of precipitant solution (23% (w/v) PEG3350, 100 mM BisTris, pH 6.0), and this drop was equilibrated against a 1-ml reservoir of precipitant solution. These conditions yielded needle-shaped crystals for each protein that grew to dimensions of 1.0 × 0.3 × 0.2 mm in two days. The co-crystal with glyoxylate were prepared by addition of 50 mM glyoxylate, 0.2–1 μl of protein solution and 1 μl of precipitant solution (30% (w/v) PEG3350, 100 mM BisTris, pH 6.5). All crystals belonged to space group P2₁2₁2₁ and contained eight molecules per asymmetric unit.

For crystal freezing, the crystals were transferred to a cryoprotectant solution with 30% (v/v) ethylene glycol in the crystallization condition for several minutes before being flash-frozen in a stream of nitrogen gas at 100 K. Diffraction data were collected on beamline 5C at the Pohang Light Source, South Korea. The raw data were processed and scaled using the HKL2000 program suite [26]. Further data analysis was carried out using the CCP4 suite [27]. Data collection statistics are summarized in Table 1.

To determine the structure of the wild-type SAV0551, molecular replacement was used with the program Molrep [28] within the CCP4 suite [27] using the homologous structure of Hsp31 from *V. cholerae* (PDB code: 414N) as a search model. The SAV0551 and Hsp31 from *V. cholerae* shows 49% sequence identity. To determine the structure of glyoxylate-bound SAV0551, molecular replacement was performed using the wild-type SAV0551 as the search model. Refinement of each crystal structure was done through iterative cycles of model building using COOT [29], followed by refinement of the models with Refmac5 and phenix.refine [30,31]. A 5% portion of the data were set aside prior to refinement for the *R* <sub>free</sub> calculations for each dataset [32]. Solvent and glyoxylate molecules became apparent in the later stages of refinement and were added into the model. Further refinement was pursued until no further decrease in *R* <sub>free</sub> was observed. Structural alignments were carried out using the program PyMOL (http://www.pymol.org) and UCSF Chimera (http://www.cgl.ucsf.edu/chimera) [33], which were then used for the construction and generation of all the figures. Protein interfaces, surfaces, and assemblies were calculated using the PISA server at the European Bioinformatics Institute (http://www.ebi.ac.uk/pdbe/prot_int/pistart.html) [34].

### Determination of chaperone activity

To monitor the chaperone activity of the wild-type and SAV0551 mutants, citrate synthase was employed as a substrate [35,36]. Initially, to identify chaperone activity, 75 μg of citrate synthase (Sigma–Aldrich) was mixed with a solution of 100 mM Tris, pH 8.0, 20 mM DTT, 6 M guanidine chloride (GnCl). The citrate synthase mixture (75 μg of citrate synthase, 100 mM Tris, pH 8.0, 6 M GnCl, 20 mM DTT) was incubated for 1 h at 25°C; consequently, the citrate synthase in this solution was denatured. After incubation, refolding of citrate synthase was achieved by 100-fold dilution with a solution of 100 mM Tris (pH 8.0) containing 5 μM wild-type and SAV0551 mutants. The diluted
Table 1 Crystallographic data collection and refinement statistics

| Data collection                        | SAV0551       | SAV0551 with glyoxylate |
|----------------------------------------|---------------|-------------------------|
| Beamline 1                            | PAL-5C        | PAL-5C                  |
| Wavelength (Å)                         | 0.98          | 0.98                    |
| Resolution range (Å)                  | 40.00-2.60    | 40.00-3.01              |
| Space group                           | P2\text{1}2\text{1}2\text{1} | P2\text{1}2\text{1}2\text{1}| |
| Unit cell parameters (Å)              | a = 96.28     | a = 97.20               |
|                                        | b = 129.45    | b = 130.12              |
|                                        | c = 187.85    | c = 187.95              |
| Observations (total/unique)           | 472, 116/72, 192 | 590, 230/47, 559        |
| Completeness (%)                      | 99.3 (97.4)   | 99.8 (99.2)             |
| Rwork (%)                             | 18.3          | 18.8                    |
| Rfree (%)                             | 26.5          | 26.6                    |
| Protein atoms                         | 17,906        | 17,836                  |
| Water molecules                       | 292           | 7                       |
| Average B value (Å²)                  | 40.4          | 44.0                    |
| r.m.s.d. bond (Å)                     | 0.021         | 0.009                   |
| r.m.s.d. angle (°)                    | 1.020         | 1.114                   |
| Ramachandran analysis (%)             |               |                         |
| Favored region                        | 94.3          | 92.9                    |
| Allowed region                        | 5.4           | 6.7                     |
| Outliers                              | 0.3           | 0.4                     |

4Numbers in parentheses indicate the statistics for the last resolution shell.
5Rwork = ΣI-<hkl>-ΣI<hkl> ; where I<kl> is the mean value of intensity of all reflections measured for all data sets.
6Rfree = ΣFobs-Fcalc-ΣFobs/Fcalc ; where Fobs is the observed structure factor amplitude, and Fcalc is the structure factor calculated from the model.

solution was mixed with acetyl-CoA, oxaloacetate, MnCl2 and 5,5′-dithiobis-(2-nitrobenzoic acid) (DTNB) to detect the activity of citrate synthase (100 mM Tris, pH 8.0, 1 mM DTNB, 0.2 mM MnCl2, 0.4 mM oxaloacetic acid, 0.3 mM acetyl-CoA). After mixing, only the active, refolded enzyme will break acetyl-CoA into the acetyl group and CoA. The CoA reacts with DTNB, which acts as a coloring agent, and this produces a yellow TNB-CoA-SH compound that is detectable at 412 nm using a Multi-Mode microplate reader (SpectraMax M5e) [37]. After a 25-min reaction period, the highest specific activity value obtained for TNB-CoA-SH was considered 100% and the lowest was 0%. The calculated specific activities of the wild-type and SAV0551 mutants were expressed as a percentage of this value.

Determination of glutathione-independent glyoxylase (glyoxylase III) activity

The glyoxylase activity assay in the wild-type and SAV0551 mutants was performed using the methylglyoxal (Sigma–Aldrich, 40% solution) in a 1D NMR experiment with absorption spectroscopy. For NMR analysis of the SAV0551 enzymatic reactions, 60 mM methylglyoxal was mixed with 50 μM SAV0551 or the cysteine mutant (C190A). The reaction mixture was incubated at 37°C for 5 min in 20 mM HEPES buffer (pH 7.5) and included 50 mM KCl and 2 mM DTT. The NMR samples contained 10% D2O for deuterium locking. All the 1D NMR experiments were performed at 298 K on an 800 MHz Bruker Ascend NMR spectrometer equipped with a cryogenic probe. All spectra were acquired and processed using the Topspin 3.2 program. On-resonance presaturation pulses were used to suppress the water signal. 1H chemical shifts were internally referenced to 4,4-dimethyl-4-silapentane-1-sulphonic acid (DSS). The number of scans (NS) was 128 and the relaxation delay was 2 s, giving a total experimental time of ~9 min.

For absorption spectroscopy, the reactions were initiated by the addition of 6 mM methylglyoxal to the enzyme in the reaction buffer (20 mM HEPES, pH 7.5, 50 mM KCl, and 2 mM DTT). The methylglyoxal and protein (wild-type or SAV0551 mutants) reaction mixtures were incubated at 37°C. The final protein concentration was 10 μM. The
remaining methylglyoxal at each time point was determined by reaction with 2,4-dinitrophenyldihydrazone (DNPH) to generate the purple chromophore methylglyoxal-bis, 2,4-dinitrophenylhydrazone after alkali treatment [38]. The assay was performed by removing a 50 μl sample from the reaction at fixed time points after enzyme addition (0, 15, 30, 45, 60, 75, 90, 105, and 120 s) and rapidly mixing the sample with 0.9 ml of distilled water. To this solution, 0.33 ml of a freshly prepared stock of DNPH reagent (0.2% DNPH dissolved in 2 N HCl) was immediately added and incubated at 37°C for 15 min. This highly acidic solution stops the enzymatic reaction, whose rate was already greatly diminished by the initial 19-fold dilution into water. The purple color of the hydrazine is developed by the addition of 1.67 ml of 3.8 M NaOH and incubation for 10 min at room temperature followed by measurement of the absorbance at 550 nm in a SpectraMax M5e. All the measurements were made in triplicate.

PDB codes
Protein co-ordinates and structure factors have been deposited in the RCSB PDB under codes 5XR1 for the native SAV0551 and 5XR3 for SAV0551 with glyoxylate.

Results
Crystal structure of SAV0551
The crystal structure of SAV0551 was determined at 2.6 Å resolution including the additional C-terminal histidine tag. The asymmetric unit contains eight molecules and each of them consists of two domains: a core domain whose similarity to DJ-1 superfamily proteins had been expected and a cap domain, which is also called the ‘P’ region. The core domain consists of an α/β sandwich fold with nine α-helices and six β-strands. Six β-strands, β4 (residues 95–100), β1 (residues 53–58), β5 (residues 151–156), β6 (residues 184–189), β8 (residues 265–268), and β7 (residues 259–262), are aligned in the center, wherein the latter β7 strand is antiparallel to the central β-strands. Nine helices surround the core of β-strands. α-helices α5 (residues 137–143), α6 (residues 160–163), α7 (residues 166–168), α8 (residues 170–181), α9 (residues 192–198), α12 (residues 239–245), α13 (residues 271–291), α2 (residues 79–91), and α4 (residues 118–132) are around the core in a clockwise direction. Additionally, four short α-helices (α1 (residues 26–32), α3 (residues 110–112), α10 (residues 218–222), and α11 (residues 224–227)) and two β-strands (β2 (residues 64–66) and β3 (residues 72–74)) form the cap domain. The core and cap domains are linked via a 22-residue long-linker (residues 33–53). Cys190 and His191 from the core domain and Asp221 from cap domain constitute a catalytic triad. The Cys190 σ1 interacts with the His191 N3, while the Asp221 carboxylate hydrogen bonds with His191 N2. This catalytic triad is well conserved in Hsp-type DJ-1 superfamily proteins (Figure 1A).

The crystal structure shows that SAV0551 is a compact homodimer. The loops (residues 1–25 and 67–71) and β-strands (β2 (residues 64–66) and β3 (residues 72–74)) from cap domain are responsible for dimer formation. Additionally, loops located in the core domain participate at the dimerization interface. The dimeric interface buries ~1110 Å2 per subunit. There are three charge–charge interaction pairs between Arg63 and Glu118, Lys107 and Glu63, and Lys132 and Asp19 on the dimeric interface of the cap domain. Hydrogen bonding interactions are observed at the dimeric interface of the core and cap domains and include Glu18, Asp19, Glu61, Glu62, Arg63, Ser102, Tyr104, and Lys236. The dimer crystal structure of SAV0551 is consistent with the results from size-exclusion chromatography, which showed a single band at ~64 kDa in solution (the molecular weight of a SAV0551 monomer is ~32 kDa) (Figure 1B).

The surface structure and electrostatic distribution of SAV0551 further display distinctive characteristics of the Hsp-type subfamily. The overall structure of the dimer is a trapezoidal formation with extensive negative charge along the core domain’s concave surface. This concave surface structure, also known as a canyon, is well conserved in the Hsp-type subfamily [1,4]. The dimer surface structure of the cap domain results in the formation of a groove that measures ~20 Å in diameter. This bowl structure does not display any charge. Therefore, it is called a hydrophobic bowl in the Hsp-type subfamily (Figure 1C). These surface structural features are considered essential factors for the binding of unstructured proteins [12]. Based on its quaternary structure, SAV0551 is classified as a definite member of the Hsp-type subfamily with chaperone function.

Comparison of SAV0551 with DJ-1 superfamily proteins
The overall fold of SAV0551 monomer shows structural similarity to other Hsp-type subfamily proteins as predicted from the sequence homology, including E. coli Hsp31 (54% sequence identity), V. cholera Hsp31 (49% sequence identity), S. cerevisiae YDR533c (12% sequence identity), and C. albicans Glx3 (14% sequence identity). A structure-based sequence alignment was acquired using the Clustal Omega web server tool [39,40] and viewed using the ESPript web server tool [41,42]. The overall fold of SAV0551 is similar to other Hsp-type subfamily members.
Figure 1. Crystal structure of SAV0551

(A) Ribbon representation of the SAV0551 monomer with two domains. SAV0551 shows a sandwich-folded main domain and a cap domain. Each domain is connected by a linker (green). SAV0551 has an α1-linker-β1-β2-α2-β4-α4-β5-α5-β6-α7-β6-α9-α10-α11-α12-β7-β8-α13 topology. The Cys190 at a sharp turn between β6 and α9 forms a catalytic triad with His191 from the main domain and Asp221 from the cap domain. (B) The SAV0551 dimer is shown in ribbon representation from a side view. Chain A is colored in pink and chain B is colored in blue. The cap domain of each subunit is shown in a darker color. The β-strands and loop from the cap domain are mainly involved at the dimeric interface. (C) The potential surface charge of the SAV0551 crystal structure calculated by UCSF Chimera, where the surfaces are colored between −10 kcal/mol·e (red) and +10 kcal/mol·e (blue) (1 kcal = 4.184 kJ). Most of the surface is negatively charged. The canyon and bowl are indicated.

The SAV0551 structure was submitted to the DALI server (http://ekhidna.biocenter.helsinki.fi/dali_server/) to identify structural homologs. The DALI algorithm reveals that SAV0551 is structurally similar to Hsp31 proteins with high Z scores. Most of the structural matches were DJ-1 family members, including Hsp31 from V. cholerae (Z score = 46.5, 0.88 r.m.s.d., 277 equivalent Co, 49% sequence identity), Hsp31 from E. coli (Z score = 45.2, 0.73 r.m.s.d., 226...
equivalent Ca, 54% sequence identity), YDR533c from *S. cerevisiae* (Z score =24.2, 1.85 r.m.s.d., 196 equivalent Ca, 12% sequence identity), and Glx3 from *C. albicans* (Z score =23.8, 2.12 r.m.s.d., 200 equivalent Ca, 14% sequence identity). The other matches, such as Hsp33 from *S. cerevisiae* (Z score =23.5, 1.97 r.m.s.d., 189 equivalent Ca, 27% sequence identity), DJ-1 from human (Z score =19.6, 2.32 r.m.s.d., 139 equivalent Ca, 27% sequenced identity) and protease Pfpl from *P. horikoshii* (Z score =18.9, 1.70 r.m.s.d., 158 equivalent Ca, 23% sequence identity), were also identified. Sequence alignment and superposition of the SAV0551 structure with the Hsp31 proteins from *E. coli* and *V. cholerae*, YDR533c from *S. cerevisiae*, and Glx3 from *C. albicans* are shown in Figures 2A,B, respectively.

Although the tertiary structures of DJ-1 superfamily members are similar, quaternary structures vary according to the DJ-1 subfamilies (DJ-1-, YhbO-, and Hsp-types). DJ-1-type subfamily proteins form dimer using α-helices, β-strands, and loops. YhbO-type subfamily proteins use three α-helices to form a dimer, and Hsp-type subfamily members interact with other subunit by N-terminal β-strands and loops [4]. In addition to the difference in the quaternary structure, another obvious difference is the architecture of the catalytic triad. Although all DJ-1 superfamily proteins have a reactive cysteine at a sharp turn between a β-strand and an α-helix, this cysteine forms different active site constellations depending on the subfamilies. The catalytic triad is not detected in DJ-1-type subfamily proteins. The DJ-1 type DJ-1 forms a catalytic dyad with a nearby histidine [43]. The DJ-1-type protein YajL does not form a catalytic dyad/triad [7]. YhbO-type proteins constitute a catalytic triad with cysteine, the histidine next to the cysteine, and an acidic residue from the other subunit. The Hsp-type proteins form a catalytic triad using the cysteine, the histidine next to the cysteine, and an acidic residue from an intrasubunit from the cap domain [4]. YhbO-type and Hsp-type subfamily proteins have an analogous catalytic triad, though the triads differ in the orientation of the acidic residue (Figure 3). Because the constellation of the three residues in the catalytic triad is identical between YhbO-type and Hsp-type subfamily proteins, the bond angles and distances between the atoms are similar. Although it is known that the catalytic triad may contribute to the protease and glyoxalase III mechanisms in the YhbO-type and Hsp-type subfamily, respectively, the exact function of the catalytic triad in the DJ-1 superfamily has not yet been revealed. In that sense, this structural difference may implicate the function of the catalytic triad and type of potential substrates.

### Chaperone activity of SAV0551 and its mutants

Hsp-type DJ-1 subfamily proteins are highly conserved in many bacterial species and function as molecular chaperones that facilitate protein folding. Under stress conditions, such as heat shock, pH shift, or oxidation, the increased expression of chaperone proteins protects cells by stabilizing unfolded proteins. Hsp31 proteins are known as chaperones, which holding molecular chaperones. The surface structure of Hsp31 is crucial for holding unfolded substrates. Hsp31 from *E. coli* work as a dimer by forming a deep acidic canyon and bowl at its dimeric interfaces [44]. When the surfaces of Hsp31 from *E. coli* were viewed, hydrophobic patches were detected around the canyon for the binding of unstructured proteins. SAV0551 expresses similar surface patterns to Hsp31 from *E. coli*, including a canyon and a bowl. SAV0551 has a deep acidic canyon that winds from the dimeric interface to each side of the subunit, and bowl structure is located on the bottom side (Figure 1C). This characteristic surface structure is well conserved in Hsp-type DJ-1 subfamily members. Other types of DJ-1 subfamily proteins have also shown chaperone activity including the DJ-1-type DJ-1 and the YhbO-type SAV1875 [8,19]. Even though the whole cap domain is absent from the DJ-1-type or YhbO-type subfamily members, the overall surface structure are conserved. Considering that the mechanism of chaperone activity is dependent upon the surface structure, the catalytic triad is not shown to be relevant to its action. To determine the chaperone activity of SAV0551 and any correlation between the catalytic triad and chaperone activity, catalytic triad mutants C190A, H191A, and D221A were designed to identify the structural and functional differences of SAV0551. The observed data showed a chaperone-facilitated renaturation of citrate synthase with the wild-type and SAV0551 mutants (Figure 4). There were no significant differences in the chaperone activity of the wild-type and SAV0551 mutants. From the present study, we identified that SAV0551 is an *S. aureus* chaperone protein and that the presence of cysteine is not a key element for the chaperone function.

### Glyoxalase III activity of SAV0551 and its mutants

Endogenous methylglyoxal is generated as an unavoidable consequence of glycolysis. It is also formed by lipid peroxidation systems, acetone metabolism, and DNA degradation [22,45,46]. Methylglyoxal is a known endogenous and environmental mutagen that can modify both DNA and proteins. To detoxify methylglyoxal, there is a glyoxalase system that converts methylglyoxal to non-toxic D-lactate. The traditional glyoxalase system is accomplished by the sequential action of the two thiol-dependent enzymes glyoxalase I and II in the presence of glutathione (GSH) [47]. Recently, a GSH-independent glyoxalase system that utilizes Hsp31 was identified in *E. coli*. In this system, the *E. coli* Hsp31 directly converts methylglyoxal to D-lactate in a single step, independent of GSH [15].
Figure 2. Comparison of SAV0551 with Hsp31 proteins from different origins

(A) Sequence alignment of SAV0551 with other Hsp31s from different origins. Sequence alignment of SAV0551 with E. coli Hsp31 (54% sequence identity), V. cholerae Hsp31 (49% sequence identity), S. cerevisiae Hsp31 (12% sequence identity), and C. albicans Hsp31 (14% sequence identity) proteins. Identical residues are colored white on a red background and similar residues are red on a white background. Secondary-structure elements (springs are α-helices and arrows are β-strands) are represented above the sequences and are numbered. The figure was constructed using ESPript [41,42].

(B) The superposition of SAV0551 (sea green) on the E. coli Hsp31 (pink, PDB code: 1PV2), V. cholerae Hsp31 (blue, PDB code: 4I4N), S. cerevisiae Hsp31 (green, PDB code: 1QVV), and C. albicans Hsp31 (orange, PDB code: 4LRU) structures. The overall fold of SAV0551 is similar to the other Hsp31 proteins. The figure was constructed using UCSF Chimera [33].

To determine the glyoxalase III activity of SAV0551, 1D NMR experiments were used. The enzymatic reaction could be analyzed by identifying newly generated product peaks. Due to the different monomeric and polymeric structures formed by methylglyoxal in H$_2$O, all the corresponding proton peaks appeared in the conventional 1D NMR spectrum of methylglyoxal (Figure 5A). The assignment of these peaks was performed on the basis of the
previous resonance assignment of methylglyoxal and its derivatives as well as with the ACD/NMR program predictor [48,49]. The spectrum predominantly showed three peaks for monohydrated and dihydrated monomeric forms of methylglyoxal; the methyl protons (CH₃) of the monohydrate and dihydrate correspond to resonances at 2.292 and 1.363 ppm, respectively, and the alkyl proton (CH) of the monohydrate corresponds to a resonance at 5.275 ppm. The peak for the dihydrate alkyl proton was not observed because its signal at ~4.8 ppm was suppressed by the water signal. Many low-intensity peaks for the polymeric forms mainly appeared in a range from 1.0 to 1.8 ppm. The reference spectrum of d-lactate in the same buffer was acquired to assign its chemical shifts (Figure 5B). The two main proton peaks for d-lactate appeared at ~4.12 ppm (CH) and 1.337 ppm (CH₃). After the enzymatic reaction of methylglyoxal with SAV0551, the 1D spectrum of this solution was obtained as shown in Figure 5C. We observed new peaks that correspond to the d-lactate produced by SAV0551 as well as a significant reduction in the peak intensities of methylglyoxal monohydrate and dihydrate. This result demonstrated that the two hydrated forms of methylglyoxal can be used as a substrate for SAV0551 to produce d-lactate, which is consistent with our data obtained by absorption spectroscopy. Additionally, the peak for Tris buffer, in which the proteins were dissolved, appeared at 3.73 ppm; the peaks for HEPES buffer were slightly changed, possibly because the production of d-lactate caused a decrease in the pH. An unassigned peak at 1.91 ppm, which might come from a methylglyoxal derivative, increased to 2.06 ppm.

In addition to the 1D NMR spectroscopy, the glyoxalase III activities of SAV0551 and its mutants were assessed by absorption spectroscopy. The amounts of methylglyoxal were detected at an absorbance of 550 nm after the reaction with DNPH to generate the purple chromophore methylglyoxal-bis-2,4-dinitrophenylhydrazone after alkali treatment [16]. The wild-type and SAV0551 mutants were employed as the enzymes. The levels of methylglyoxal were substantially reduced by the time when it was mixed with wild-type SAV0551. However, the glyoxalase III activities of the SAV0551 mutants were diminished (Figure 6). The E81A, C190A, and H191A mutants showed little or no activities, while the D221A mutant displayed marginal activity. From our results, SAV0551 is determined to be a
glyoxalase III and the key residues involved in its function are Glu81 and catalytic triad residues Cys190, His191, and Asp221.

Components required for glyoxalase III function

The mechanism of the traditional glyoxalase system, which includes glyoxalase I and II, is well known, and cofactors such as GSH and metal ions are necessary for the reaction [50-52]. However, even though the substrate and product are the same, glyoxalase III does not require any cofactors. According to the known glyoxalase III mechanism via *Arabidopsis thaliana* DJ-1, the SAV0551 glyoxalase III mechanism is expected to be performed by the residues in the active site, including the catalytic triad (Cys190, His191, and Asp221), an absolutely conserved glutamate amongst the DJ-1 superfamily (Glu81), and a base residue histidine residue that is positioned differently compared with the structure of *A. thaliana* DJ-1 [52]. The characteristic aspect of the glyoxalase III mechanism is that the enzyme itself has all the required components within the active site; therefore, it functions without any cofactors.

To produce D-lactate from methylglyoxal, a ketone carbonyl group (the C2 carbonyl group) is reduced to a hydroxyl group, while the aldehyde carbonyl group is oxidized to an alcohol. When the glyoxalase III mechanism is compared with the traditional glyoxalase system (glyoxalases I and II), the initial nucleophilic reaction is completed by the catalytic cysteine instead of GSH, which is an essential cofactor in the traditional glyoxalase system. Once the hemithioacetal forms, glyoxalase I moves the proton from C1 to C2 and generates S-D-lactoylglutathione. The Zn2+ ions stabilize the enediol intermediate to lower the free energy of proton transfer, which finally leads to the production of S-D-lactoylglutathione [53,54]. In the *A. thaliana* DJ-1 glyoxalase III mechanism, the backbone glycine stabilizes O1C11 and an absolutely conserved glutamate stabilizes O2C2 so that the proton can transfer from C1 to C2.
Figure 5. NMR-based assessment of SAV0551 glyoxalase III activity

(A) 1D $^1$H NMR spectrum for methylglyoxal. (B) 1D $^1$H NMR spectrum for D-lactate. (C) 1D $^1$H NMR spectrum for D-lactate produced by SAV0551. Newly generated and decreased peaks are represented by a ‘*’ and ‘#’, respectively. The peak at 3.73 ppm is derived from the Tris buffer in which the protein stock is dissolved. An expanded view of the peaks for MGO dihydrate (CH$_3$) and -lactate (CH$_3$) are shown in the rectangular box. The reaction was performed at 37 °C in 20 mM HEPES (pH 7.5) 50 mM KCl, and 2 mM DTT. The reaction concentrations of SAV0551 and its substrate methylglyoxal were 50 μM and 60 mM, respectively.

via the glutamate residue, similar to the glyoxalase I mechanism. The histidine residues in the glyoxalase III active site substitute for the action of the Zn$^{2+}$ ions. The final step is hydrolysis, which is the function of glyoxalase II, and metal ions are also required for the introduction of the water ions. Once again, the histidine residue holds the water molecule to aid in the hydrolysis.

In light of the structural comparison between A. thaliana DJ-1 and SAV0551, the active site residues are well conserved, except for the base histidine. Although the A. thaliana DJ-1 histidine residue (His$^{52}$) is proven to be crucial for the glyoxalase III function, the geometry of the histidine residues is different in SAV0551 crystal structure compared with A. thaliana DJ-1. Two histidine residues (His$^{78}$ and His$^{160}$) are positioned in the active site of SAV0551 (Figures 7A,B).
Figure 6. Glyoxalase III activity of the wild-type and SAV0551 mutants
SAV0551 degrades methylglyoxal. Thus, 6 mM methylglyoxal was used initially, and the remaining amount of methylglyoxal was calculated after incubation with the wild-type and E81A, C190A, H191A, and D221A mutant proteins over time.

Figure 7. Catalytic triad and surrounding residues around the Cys active site
The glyoxylate-bound co-crystal structures of SAV0551, A. thaliana DJ-1, and human DJ-1. Distances are given in Angstroms. Hydrogen bonds are shown in dark dotted lines. (A) Chain A in the glyoxylate-bound structure of SAV0551. Unlike the O1C1 and O2C2 oxygen atoms, the glyoxylate C2 oxygen atom (O3C2) does not form stable hydrogen bonds. The 2Fo-Fc electron density map contoured at a level of 1.0σ is shown in gray. (B) The A. thaliana DJ-1 co-crystal structure with glyoxylate (cyan, PDB code: 4OGG) and the corresponding region in SAV0551 (light green). All three glyoxylate oxygen atoms form stable hydrogen bonds. (C) The glyoxylate co-ordination in human DJ-1 (magenta, PDB code: 4OGF) and the corresponding region in SAV0551 (light green). The co-crystal structure with glyoxylate is stabilized by hydrogen bonds through the introduction of water molecules.

Structural study of the SAV0551 active site as a glyoxalase III
To secure the SAV0551 glyoxalase III reaction intermediate and to determine the difference in the histidine location between A. thaliana DJ-1 and SAV0551, SAV0551 was crystallized with the DJ-1 glyoxalase III inhibitor glyoxylate. The glyoxylate is stably co-ordinated in the active site of A. thaliana DJ-1 and Homo sapiens DJ-1. In the A. thaliana DJ-1 co-crystal structure, the glyoxylate C1 carbon–oxygen atom (O1c1) forms stable hydrogen bonds with His121 (2.64 Å) and Gly90 (2.64 Å). The other oxygen (O2c2) atom at the C2 carbon is stabilized by Glu19 (2.82 Å) and a water molecule (2.91 Å). The third oxygen atom (O3c2), which is a methyl group in the substrate methyglyoxal structure, also shows hydrogen bonds with His52 (2.73 Å) and a water molecule (3.12 Å) (Figure 7B). In the human DJ-1 co-crystal structure, the oxygen atom at C1 (O1c1) forms stable hydrogen bonds with Glu18 (3.02 Å) and Gly75 (2.61 Å). The two oxygen atoms at C2 show hydrogen bonds with water molecules (2.71 and 2.71 Å, respectively) (Figure 7C) [52]. However, the glyoxylate in the SAV0551 crystal structure is not perfectly co-ordinated, potentially because of the relatively low resolution. Still, the SAV0551 co-crystal structure shows the clear density of the glyoxylate C1 and the interaction with Glu18 and Gly159. With this information, the rigid glyoxylate structure is well positioned with respect to the active site. Amongst the three glyoxylate oxygen atoms, O3c2 does not form hydrogen bonds, while O1c1 and O2c2 show stable hydrogen bonds. O1c1 forms a hydrogen bond with His191 (2.93 Å) and Gly159.
(2.48 Å). The C2 carbon–oxygen atom (O2C2) forms a hydrogen bond with His78 (2.83 Å) and a water molecule (2.71 Å). The third oxygen atom (O3C2) is 3.35 Å from a water molecule and 3.65 Å from His191, which demonstrate weak electrostatic bonds. Because of the different constellation of the residues in the SAV0551 and A. thaliana DJ-1 active sites, the glyoxylate co-ordination in SAV0551 can be limited because the O3C2 is not stabilized by residues or water molecules. The glyoxylate-bound crystal structure in SAV0551 is similar to the glyoxalase III reaction intermediate (hemithioacetal) because the glyoxalase III substrate methylglyoxal has a single oxygen atom on C2. The O3C2 is a methyl group in methylglyoxal, and the methyl group does not form any hydrogen bonds. From the glyoxylate-bound structure, we determined that His78 provides a hydrogen atom to the O2C2 and that a water molecule attacks the C1 atom to hydrolyze the n-lactoyl cysteine structure to produce n-lactate. Therefore, we can conclude that SAV0551 is a glyoxalase III that possesses all the required components in its active site, including His78, which acts as a base in the glyoxalase III mechanism. Additionally, in the SAV0551 co-crystal structure, excess amounts of glyoxylate are also located in the aspartate-rich regions within the adjacent cleft of two irrelevant chains, which are not dimers.

Discussion

The structural studies herein confirmed SAV0551 from S. aureus Mu50 as a member of the DJ-1 superfamily and that it has all the pertinent traits of this group, including the sandwich fold and nucleophilic arm. It specifically belongs to the Hsp-type subfamily because of the presence of cap domain and aspects of its dimeric interface. Simultaneously, Hsp-type DJ-1 subfamily proteins are categorized as Class 20 (Hsp31) chaperones by the recent classification of small HSPs (SHSP) [55]. Hsp31 from E. coli functions as a holding chaperone and displays chaperone activity by holding unfolded protein with its canyon, bowl, and hydrophobic surface structure [12]. SAV0551 has a similar canyon, bowl, and hydrophobic surface structure as E. coli Hsp31, suggesting that SAV0551 may work as a chaperone. We discovered that the wild-type and SAV0551 mutants assisted in the folding of citrate synthase, indicating the importance of overall its structure rather than specific residues.

Accumulation of methylglyoxal causes endogenous damage by reacting with proteins and nucleic acids. The DJ-1 superfamily, especially Hsp-type DJ-1 subfamily proteins, is well known for its role in the cellular defense system toward methylglyoxal toxicity. SAV0551 is also predicted to perform similar biological functions as members of the Hsp-type subfamily. To determine the glyoxalase activity, 1D NMR and absorption spectroscopy experiments were performed. The co-crystal structure with the known inhibitor glyoxylate was also determined to elucidate the structure of the intermediate. The results demonstrate that SAV0551 is a glyoxalase III that produces non-toxic n-lactate from toxic methylglyoxal independent of any cofactors. We discovered that residues in the catalytic triad (Cys190, H191, and D221), an absolutely conserved Glu81, and His78 are responsible for the glyoxalase III function. The catalytic cysteine (Cys190) attacks the methylglyoxal aldehyde group (C1) and forms a hemithioacetal intermediate. Then, Glu81 moves a proton from C1 to C2 via stabilization of the enediol through the neighboring residues His78, Gly159, and His191 and a water molecule. After the formation of n-lactoyl cysteine, a water molecule attacks C1 and hydrolyzes it to produce Δ-lactate. In the present study, we have shown the significance of the catalytic triad residues (Cys190, H191, and D221) and the absolutely conserved glutamate residue (Glu81) in the DJ-1 superfamily and revealed the important base residue His78 via a co-crystal structure.

In summary, the crystal structure of the Hsp-type protein SAV0551 and a co-crystal structure with glyoxylate were determined. SAV0551 has the distinctive characteristic structure of a holding chaperone, including the oligomeric state, surface canyon, and bowl structure and is similar to E. coli Hsp31. SAV0551 shows chaperone activity regardless of active site mutations, whereas the catalytic cysteine and active site residues are crucial for its glyoxalase III function. Because DJ-1 superfamily proteins have multiple stress defense functions including chaperone and glyoxalase III, our study will promote a better understanding of how bacteria can defend against stress conditions. By focussing on DJ-1 superfamily proteins such as SAV0551, we can gain insights into the essential cellular stress control systems against stress condition. Future research on DJ-1 family proteins and in vivo studies are required to fully understand the fundamentals of stress response functions.

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Competing interests

The authors declare that there are no competing interests associated with the manuscript.
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Author contribution
H.J.K. designed and performed the experiments, analyzed the data, and wrote the paper under the supervision of A.-R.K. and B.-J.L. H.J.K. and K.-Y.L. performed the NMR measurements and analyzed the NMR data.

Abbreviations
DTT, dithiothreitol; DNPH, 2,4-dinitrophenylhydrazine; DTNB, 5,5′-dithiobis-(2-nitrobenzoic acid); GSH, glutathione; HSP, heat shock protein.

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