TITLE:
High-resolution structure of the recombinant sweet-tasting protein thaumatin I.

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CITATION:

ISSUE DATE:
2011-06

URL:
http://hdl.handle.net/2433/142313

RIGHT:
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High-resolution structure of the recombinant sweet-tasting protein thaumatin I

Thaumatin, an intensely sweet-tasting plant protein, elicits a sweet taste at a concentration of 50 nM. The crystal structure of a recombinant form of thaumatin I produced in the yeast *Pichia pastoris* has been determined to a resolution of 1.1 Å. The model was refined with anisotropic B parameters and riding H atoms. A comparison of the diffraction data and refinement statistics for recombinant thaumatin I with those for plant thaumatin I revealed no significant differences in the diffraction data. The R values for recombinant thaumatin I and plant thaumatin I ($F_o > 4\sigma$) were 9.11% and 9.91%, respectively, indicating the final model to be of good quality. Notably, the electron-density maps around Asn46 and Ser63, which differ between thaumatin variants, were significantly improved. Furthermore, a number of H atoms became visible in an OMIT map and could be assigned. The high-quality structure of recombinant thaumatin with H atoms should provide details about sweetness determinants in thaumatin and provide valuable insights into the mechanism of its interaction with taste receptors.

1. Introduction

Thaumatin is a sweet-tasting protein that elicits a sweet taste sensation at 50 nM. Since it is nearly 100 000 times sweeter than sucrose on a molar basis (Van der Wel & Loeve, 1972), thaumatin has potential use as a low-calorie sweetener for industrial applications and could be useful in clarifying the mechanisms of the perception of sweet taste (Temussi, 2002, 2006; Masuda & Kitabatake, 2006).

Thaumatin is isolated from the arils of *Thaumatococcus daniellii* Benth, a plant native to tropical West Africa. Plant thaumatin consists of five intensely sweet forms, with two major components (thaumatin I and II) and three minor components (thaumatin a, b and c). The amino-acid sequences of the major components of thaumatin have been determined by Edman degradation and the nucleotide sequences have also been determined (Iyengar et al., 1979; Lee et al., 1988; Edens et al., 1982; Ide, Masuda et al., 2007). The results showed that four or five positions (N46K, S63R, K67R, R76Q and D113N) differ between the two major components of thaumatin.

Thaumatin is frequently used in crystallization studies and 34 structures have been deposited in the Protein Data Bank to date. Although plant-sourced thaumatin contains several variants, mainly thaumatin I and thaumatin II, no further purification tends to be performed for crystallization. For this reason, some parts of the electron-density maps, especially at residues 46 and 63, are often unclear. To clarify these ambiguous regions, the determination of an atomic resolution structure by analyses using homogeneous samples would provide a number of insights (Sauter et al., 2001; Howard et al., 2004; Biadene et al., 2007; Wang et al., 2007; Sherawat et al., 2007).

Although much research has been devoted to producing thaumatin using microorganisms (Faus, 2000; Masuda & Kitabatake, 2006), the production of an active sweet form is still difficult since thaumatin contains eight disulfide bonds. While some studies have achieved acceptable yields of recombinant thaumatin, there were still a few amino acids attached at the N-terminus owing to a deficient protease...
Table 1
Differences in amino-acid sequence among thaumatin variants.

<table>
<thead>
<tr>
<th>Thaumatin</th>
<th>Amino-acid position</th>
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</thead>
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<tr>
<td></td>
<td>46</td>
</tr>
<tr>
<td>Amino-acid sequence</td>
<td></td>
</tr>
<tr>
<td>I (Iyengar et al., 1979)</td>
<td>Asn</td>
</tr>
<tr>
<td>A (Lee et al., 1988)</td>
<td>Asn</td>
</tr>
<tr>
<td>B (Lee et al., 1988)</td>
<td>Lys</td>
</tr>
<tr>
<td>I (Kaneko &amp; Kitabatake, 2001)</td>
<td></td>
</tr>
<tr>
<td>Deduced amino-acid sequence from cDNA</td>
<td></td>
</tr>
<tr>
<td>I (Ily, Masuda et al., 2007), AF355098</td>
<td>Asn</td>
</tr>
<tr>
<td>II (Edens et al., 1982), J01209</td>
<td>Lys</td>
</tr>
<tr>
<td>II (Verrips et al., 1982), A15660</td>
<td>Lys</td>
</tr>
<tr>
<td>II (Masuda et al., 2004)</td>
<td>Lys</td>
</tr>
</tbody>
</table>

(Weickmann et al., 2004; Lombraña et al., 2004; Masuda et al., 2004; Ide, Kaneko et al., 2007). More recently, homogeneous recombinant thaumatin was successfully obtained from the yeast *Pichia pastoris* using a pre-sequence of thaumatin as a secretion signal (Ide, Masuda et al., 2007; Masuda et al., 2010).

In the present study, the high-resolution structure of recombinant thaumatin I produced by *P. pastoris* was determined at 1.1 Å resolution. The similarities and differences in the refined structures of recombinant thaumatin I as well as plant thaumatin I were investigated in detail. A high-resolution structural analysis with H atoms showed an improvement of the maps and could discriminate between thaumatin variants.

2. Materials and methods

2.1. Amino-acid sequence of thaumatin I

The amino-acid sequence of thaumatin I was determined by Iyengar and coworkers by Edman degradation (Iyengar et al., 1979). Later, Lee and coworkers re-examined the two major components of natural thaumatin and designated them thaumatin A and thaumatin B (Lee et al., 1988). The amino-acid sequences of thaumatin A and thaumatin B differed from that of thaumatin I determined by Iyengar and coworkers by one amino acid (Asp113 instead of Asn) and two amino acids (Lys46 instead of Asn and Asp113 instead of Asn), respectively.

The nucleotide sequence of thaumatin II was determined by Edens et al. (1982) and the deduced amino-acid sequence showed that thaumatin II differs from thaumatin I at five positions (N46K, S63R, K67R, R76Q and N113D) and from thaumatin A at four positions (N46K, S63R, K67R and R76Q). Recently, the nucleotide sequence of thaumatin II from cloned cDNA has been determined in detail (Ide, Masuda et al., 2007). The deduced amino-acid sequence from this cDNA differed from that of thaumatin I reported by Iyengar and coworkers at residue 113 (Asp113 instead of Asn) and was identical to that of thaumatin A (Table 1).

2.2. Materials

Plant thaumatin was purchased from Wako Pure Chemical Industries Ltd, Osaka, Japan. Potassium sodium tartrate was obtained from Nacalai Tesque Inc., Kyoto, Japan. N-(2-Acetamido)-iminodiacetic acid (ADA) was obtained from Dojindo, Kumamoto, Japan. All other chemicals were of guaranteed reagent grade for biochemical use.

2.3. Expression and purification of recombinant thaumatin I

The expression of recombinant thaumatin was performed in *P. pastoris* as described previously (Masuda et al., 2005, 2010). A 7 L fermenter (TS-M7L, Takasugi Seisakusho Co., Tokyo, Japan) was used to secrete recombinant thaumatin I into the culture medium. After centrifugation of the culture medium, the supernatant was applied onto an SP-Sephadex C-25 cation-exchange column and the bound proteins were eluted with 5 mM sodium phosphate buffer pH 7.0 containing 500 mM NaCl. The fractions containing thaumatin were combined and precipitated with 75% ammonium sulfate and purified by HW-50F gel-filtration chromatography (Toyosh Co, Tokyo, Japan). The concentrations of thaumatin were determined spectrophotometrically using a molar extinction coefficient ε278 of 17 000 M cm⁻¹ (van der Wel & Loeve, 1972).

2.4. Purification of plant thaumatin I

The purification of plant thaumatin I was performed as described by Kaneko & Kitabatake (1999). Briefly, thaumatin solution was applied onto an SP-Sephadex C-25 cation-exchange column and the bound proteins were eluted with a linear gradient of 20–120 mM NaCl in 5 mM sodium phosphate buffer pH 7.0. Fractions containing thaumatin I were further purified by HW-50F gel-filtration chromatography. The purity of the proteins was checked by SDS-PAGE and native PAGE.

2.5. Crystallization and data collection

The purified recombinant thaumatin and plant thaumatin were concentrated using a Centricon-10 (Millipore, Bedford, Massachusetts, USA) and the protein concentration was measured using a NanoDrop ND-1000 spectrophotometer (NanoDrop Technologies Inc., Rockland, Delaware, USA). Crystallization was performed using the hanging-drop vapour-diffusion method. The hanging drops were prepared by mixing 5 µl 10–100 mg ml⁻¹ protein solution and 5 µl reservoir solution. The reservoir solution consisted of 0.1 M ADA, 0.5–1.0 M potassium sodium tartrate pH 6.5–6.8 and 0, 10 or 25% glycerol. All solutions were prepared with sterile water and filtered with a MILLEX-GV membrane (Millipore). Diffraction data were obtained to 1.1 Å resolution. The crystal was placed in a cold nitrogen-gas stream and X-ray diffraction images were collected using an R-AXIS V area detector (Rigaku, Tokyo, Japan) with synchrotron radiation of wavelength 0.7 Å at the BL-38B1 station of SPring-8 (Hyogo, Japan). The data obtained were processed, merged and scaled using the HKL-2000 program package (Otwinowski & Minor, 1997). Data-collection and structure-solution statistics are shown in Tables 2 and 3.
2.6. Structure refinement and validation

The structures of recombinant thaumatin and plant thaumatin were determined by molecular replacement using the program MOLREP (Vagin & Teplyakov, 2010) in the CCP4 suite (Winn et al., 2011) with the previously reported thaumatin structure as a reference (PDB code 1rqw; Q. Ma & G. M. Sheldrick, unpublished work). Modelling was performed using TURBO-FRODO on a Silicon Graphics computer and Coot (Emsley & Cowtan, 2004). \(|F_o| - |F_c|\) and \(2|F_o| - |F_c|\) maps were used to locate the correct model. Several rounds of refinement were carried out to improve the model by increasing the data resolution to 1.5 Å. Water molecules were incorporated where the difference density exhibited a value of greater than 3σ above the mean and the \(2|F_o| - |F_c|\) map showed a density of greater than 1.0σ. Refinement was performed using SHELXL97 (Sheldrick, 2008) with stereochemical restraints based on those of Engh & Huber (1991). All reflections were included with no σ cutoff; 5% of the data were randomly selected and omitted during refinement for cross-validation by means of the free R factor (Brünger, 1992). The occupancy of the major conformation was refined first; the second or third conformation was then assigned and refined based on its \(|F_o| - |F_c|\) map and finally atoms of all conformations were refined. Disordered residues were taken into consideration and possible side-chain conformations were generated. The occupancies of the disordered residues were treated as free variables and were refined using SUMP and FVAR restraints. Refinement of the model with isotropic B factors against data to a resolution of 1.1 Å using SHELXL resulted in an \(R_{work}\) of 17.55% and an \(R_{free}\) of 19.54%. Subsequent anisotropic B-factor refinement against data to 1.1 Å resolution lowered \(R_{work}\) and \(R_{free}\) to 11.97% and 15.01%, respectively (corresponding to a fall in \(R_{free}\) of 4.53%). Next, H atoms were modelled using HFIX and \(R_{work}\) and \(R_{free}\) fell to 10.80% and 13.54%, respectively. H atoms were only included for protein atoms, not for tartrate/glycerol/solvent atoms. To finalize the refinement, the model including H atoms was refined against all data for 30 cycles using conjugate-gradient least-squares minimization, leading to an R factor of 10.83% for 102 055 reflections and 9.11% for 80 465 reflections with \(F_o > 4\sigma\) in the resolution range 10–1.1 Å. The quality of the structure was assessed using PROCHECK (Laskowski et al., 1993) and WHAT IF (Hekkelman et al., 2010). Standard uncertainties were estimated with SHELXPRO through a block-diagonal calculation after removal of all restraints (Sheldrick, 2008). The electron-density maps and structural images were generated using PyMOL (DeLano, 2002).

The coordinates and observed intensities of recombinant thaumatin I and plant thaumatin I have been deposited in the PDB (accession codes 3al7 for recombinant thaumatin and 3ald for plant thaumatin).

3. Results and discussion

3.1. Crystallization of recombinant thaumatin I

In the presence of sodium potassium tartrate, thaumatin crystallizes in a tetragonal system. We purified recombinant thaumatin I from the culture medium of the yeast P. pastoris and attempted its crystallization. Numerous pyramid- and bipyramid-shaped crystals were obtained in a few days. We also attempted crystallization in the presence of 25% (v/v) glycerol for a high-resolution structural analysis. The first crystal appeared in only two weeks in the presence of 25% (v/v) glycerol, which was much faster than for purified plant thaumatin I and unpurified thaumatin, crystals of which appeared in three to four weeks (Charron et al., 2002). It was suggested that the rate of equilibration of the protein solution by vapour diffusion was significantly reduced in the presence of glycerol and it took at least 30 d to form pyramid-shaped crystals when unpurified plant thaumatin was used (Charron et al., 2002). It appeared that the homogeneous recombinant thaumatin affected the nucleation of the

![Figure 1](https://repository.kulib.kyoto-u.ac.jp)

**Figure 1**
Overall structure of recombinant thaumatin I. Tartrate ions are shown in red and glycerol molecules are shown in blue.

![Figure 2](https://repository.kulib.kyoto-u.ac.jp)

**Figure 2**
Ramachandran plot for recombinant thaumatin. Glycine residues are represented by triangles. 90.5% of the residues are located in most favoured regions, 8.9% in additional allowed regions and 0.6% (Asp25) in generously allowed regions.
crystals. The crystals obtained in this condition belonged to space group $P4_1212$. The unit-cell parameters are listed in Table 2.

### 3.2. Overall structure of recombinant thaumatin I

The final model of recombinant thaumatin I consisted of 207 residues with a total of 3386 protein atoms including 1633 H atoms, two tartrate ions, four glycerol molecules and 476 water molecules (Fig. 1). A single thaumatin molecule is contained in the asymmetric unit. The \(R\) factor (\(R_{\text{cryst}}\)) for the final structure is 9.11\% for 80 465 reflections with \(F_o > 4\sigma\) and 10.83\% for all data (102 055 reflections) in the resolution range 10–1.1 Å, suggesting that the final refined model is of high quality. A Ramachandran plot calculated for the final model showed that 90.5\% of the residues are in most favoured regions, 8.9\% are in additional allowed regions and 0.6\% are in generously allowed regions. No residues are in disallowed regions (Fig. 2, Table 3). The residue located in the generously allowed region is Asp25, which is part of the thaumatin loop (Asp21-Ala22-Ala23-Leu24-Asp25-Ala26); this was not observed in other non-sweet thaumatin-like proteins (Koiwa et al., 1999). The average $B$ factors for $C^\alpha$, side-chain and all protein atoms were 9.82, 16.39 and 17.02 Å\(^2\), respectively. Positional uncertainties were plotted against thermal parameters and 73\% of them were located in the region below the 0.03 Å positional e.s.d. and 30 Å\(^2\) equivalent $B$ value (Fig. 3). We used 22 204.63 Da as the molecular mass of recombinant thaumatin and plant thaumatin to calculate the Matthews coefficient $V_M$ (Matthews, 1968). The Matthews coefficient $V_M$ and solvent content were calculated as 2.81 Å\(^3\) Da\(^{-1}\) and 56.21\%, respectively (Table 3). OMIT maps of Phe90, Ile102 and Tyr199 clearly showed positive density (green) around these residues, suggesting the existence of H atoms (Fig. 4). Furthermore, the OMIT map of Asp113 is quite different from that of Asn104 (Fig. 5). So far, it has been quite difficult to determine whether the residue at 113 is Asp or Asn in plant thaumatin directly from the density maps. This might be because the resolution around the residue is still poor and unpurified plant thaumatin contains some variants. Since the residue in the recombinant structural communications

#### Table 3

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<tr>
<th>Final model statistics.</th>
<th>Recombinant</th>
<th>Plant</th>
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<td>Resolution (Å)</td>
<td>10–1.10</td>
<td>10–1.10</td>
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<tr>
<td>Unique reflections</td>
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<td>102886 [(F_o &gt; 4\sigma; 89235)]</td>
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<td>Isotropic $R_{\text{work}}/R_{\text{free}}$ (%)</td>
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<td>17.82/19.91 (16.38/18.53)</td>
</tr>
<tr>
<td>Anisotropic $R_{\text{work}}/R_{\text{free}}$ (%)</td>
<td>11.97/15.01 (10.19/13.16)</td>
<td>12.19/15.37 (11.07/14.27)</td>
</tr>
<tr>
<td>Anisotropic $R_{\text{work}}/R_{\text{free}}$ (including protein H atoms) (%)</td>
<td>10.80/13.54 (9.07/11.73)</td>
<td>10.96/13.71 (9.87/12.64)</td>
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<tr>
<td>$R_{\text{cryst}}$ (%)</td>
<td>10.83 (9.11)</td>
<td>10.99 (9.91)</td>
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<tr>
<td>Protein atoms (non-H)</td>
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<td>Protein atoms (H)</td>
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<td>Glycerol atoms</td>
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<td>$B$ factors (Å(^2))</td>
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<tr>
<td>Average</td>
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<tr>
<td>Solvent content (%)</td>
<td>56.21</td>
<td>56.23</td>
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</table>

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Figure 3

E.s.d. analysis of recombinant thaumatin I. Plot of positional uncertainty versus the thermal parameter for C atoms (black), N atoms (cyan) and O atoms (red) from the structure of recombinant thaumatin at a resolution of 1.1 Å.

Figure 4

The model and electron density for recombinant thaumatin I around Phe90, Ile102 and Tyr199. (a) Phe90. (b) Ile102. (c) Tyr199. $\sigma$-weighted $2mF_o - DF_c$ maps omitting H atoms contoured at 3.0σ are shown in green. The difference in density clearly shows numerous H atoms.
nant thaumatin is Asp, the results of the present study should help to clarify the residue at position 113 in plant thaumatin.

3.3. Amino-acid residues in multiple conformations

The quality of the 1.1 Å resolution electron-density map of recombinant thaumatin I allowed the side chains of 20 residues (Glu4, Arg8, Leu31, Glu35, Asp55, Tyr57, Asp70, Arg79, Arg82, Met112, Arg119, Arg122, Val124, Ser155, Cys159, Lys163, Glu168, Arg171, Asp179 and Arg200) to be modelled in two conformations; however, we could not identify alternative structures of Cys66, Cys126 and Cys193 in the electron-density map of recombinant thaumatin I (Fig. 6). It is known that the electron density of the disulfide bridges and Cys residues as having alternative side-chain conformations; how-

3.4. Comparison of the structure of plant thaumatin

The differences in r.m.s. deviation on Cα atoms between the final refined structure of recombinant thaumatin (PDB entry 3al7) and those of purified plant thaumatin [PDB entries 2vhk (Asherie et al., 2009) and 3ald] and unpurified thaumatin [PDB entries 1rqw (Q. Ma & G. M. Sheldonr, unpublished work), 1kwn (Sauter et al., 2002) and 1lxz (Charron et al., 2002)] were investigated. The results showed that the r.m.s.d. values against purified thaumatin (2vhk and 3ald) were 0.067 and 0.073 Å, respectively, whereas they ranged from 0.19 to 0.24 Å when compared with unpurified thaumatin (1rqw, 1kwn and 1lxz). These results indicated that recombinant thaumatin I is similar in overall structure to purified plant thaumatin I (2vhk and 3ald) but significantly different from unpurified thaumatin.

We also attempted to crystallize purified plant thaumatin I under the same conditions as used for recombinant thaumatin I. The data set for plant thaumatin had an Rmerge of 7.0% and was 99.7% complete (Table 2). There were no significant differences in data collection between recombinant thaumatin and plant thaumatin.

The structure of purified thaumatin I was refined to a final R factor of 10.99% for 102 886 reflections and of 9.91% for reflections with Fo > 4σ (89 235 reflections) in the resolution range 10.0–1.1 Å. The average B factors for plant thaumatin were 10.53 Å² for Cα atoms, 16.39 Å² for side-chain atoms and 17.33 Å² for all atoms (Table 3). Compared with the results for recombinant thaumatin I, the final R

Figure 5
Comparison of the models of recombinant thaumatin I and plant thaumatin I around Asn104 and Asp113. The OMIT maps of recombinant thaumatin are shown in (a) for Asp113 (B factors: OD1, 7.74 Å²; OD2, 9.28 Å²) and (c) for Asp104 (B factors: ND2, 6.78 Å²; OD1, 7.21 Å²). In contrast, the OMIT maps of plant thaumatin I are shown in (b) for Asp113 (B factors: OD1, 8.39 Å²; OD2, 10.19 Å²) and (d) for Asp104 (B factors: ND2, 7.40 Å²; OD1, 7.93 Å²). The σA-weighted 2mFo − DFo maps contoured at 1.0σ are drawn in blue and the mFo − DFo maps omitting H atoms contoured at 3.0σ are shown in green.

Figure 6
Models of Cys residues in recombinant thaumatin I. Four of the eight disulfide bonds are indicated: (a) Cys56–Cys66, (b) Cys121–Cys135, (c) Cys126–Cys177 and (d) Cys159–Cys164. Disulfide linkages are shown in orange. σA-weighted 2mFo − DFo maps contoured at 1.0σ are drawn in blue and mFo − DFo maps contoured at 3.0σ and −3.0σ are shown in green and red, respectively.
factor of plant thaumatin is slightly higher than that of recombinant thaumatin. The maps of plant thaumatin showed no obvious alternative conformations of the 16 Cys residues apart from Cys159 (data not shown). In contrast, the electron-density maps around Asn46 and Ser63, maps contoured at 1.0σ, −DF, maps contoured at 3.0σ and −3.0σ are shown in blue and mFo − DFo, maps contoured at 3.0σ and −3.0σ are shown in green and red, respectively.

The atomic structural features of recombinant thaumatin with H atoms obtained in this study will provide important information on the perception of the sweet taste of thaumatin.

This work was supported by Grants-in-Aid for Young Scientists (B) (TM: No. 19780074) and Scientific Research (C) (TM: No. 22580105) from The Japan Society for the Promotion of Science and by the Japan Food Chemical Research Foundation. The synchrotron-radiation experiments were performed at BL35B1 and BL26B1 at SPring-8 with the approval of the Japan Synchrotron Radiation Research Institute (JASRI; proposal Nos. 2009A1096 and 2009B1379).

References


