# Serratia marcescens B4A chitinase thermostability enhancement by S390I QuikChange site directed mutagenesis

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#### **Abstract**

Thermostable chitinases are useful for industrial and biotechnological applications. This paper reports the stabilization of chitinase from Serratia marcescens B4A through rational mutagenesis. Changing of Ser 390 to Ile in S. marcescens. The stabilization was enhanced through entropic stabilization by reduction of the loop length and also by increasing of the beta chain length. With this replacement, polar uncharged residue changed to non-polar one and increased the hydrophobic interactions. Furthermore Isoleucine has branched β-carbon that restricts the backbone conformation more than non-branched residues. Finally all of these factors lead to entropic stabilization and thermal stabilization. The results exhibited that the optimal temperature and pH for enzyme activity of native chitinase were not changed by mutagenesis which showed that mutation didn't affect the original characteristics of the enzyme, the K<sub>m</sub> values of native and mutant chitinase were different very little, showing that the affinity of enzyme towards the substrate and also the natural flexibility of chitinase did not change by mutation. Besides the V<sub>max</sub> value of the mutant chitinase was decreased, while its pH stability was increased briefly, but its thermal stability was increased remarkably. Mutation made chitinase to tolerate high temperatures up to 90°C. In addition its activity was increased at 50°C, 60°C for 120 min and up to 2 hours of incubation period and the mutant chitinase demonstrated a high level of activity at 60°C. These results show that entropic stabilization works well for chitinase and this approach may be generally applicable for stabilization of other proteins.

**Keywords:** Chitinase, Thermal stability, *Serratia marcescens* B4A, Loop,  $\beta$  chain, unfolding

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#### Introduction

Chitin, a polysaccharide made of β-(1,4) linked N-acetyl D-glucosamine units, is the second most plentiful biopolymer and a source of renewable raw materials (Patil et al., 2000; Annamalai et al., 2011; Wang et al., 2015). Chitin has applications in industrial, biotechnology, agriculture, material and medical, food and nutrition sciences (Xia et al., 2011). Chitinases (EC 3.2.1.14) are widespread enzymes in nature with ability to degrade chitin smaller chito-oligosaccharide into segments (Bhattacharya et al., 2007; Tsuji et al., 2010). These enzymes are efficient biomarkers in cancer disease (Wang et al., 2015) and are used for treatment of CF-associated fungal disease (Hector et al., 2016). Some chitinases such as chitinase 3-like 2 and chitotriosidase are used in early MS diagnosis (Elmonem et al., 2016; Møllgaard et al., 2016).

Thermostable chitinases various demand in biotechnology sectors (Bjørk et al., 2003; Aliabadi et al., 2016). The need, similar to the other interesting enzymes, has been addressed through protein engineering. However, the use of a rational design for protein stability improvement seems to be protein specific and any of such attempts would require to be directed by the detailed structural information (Giver et al., 1998; Gåseidnes et al., 2003; Khurana et al., 2011; Matroodi et al., 2013) and understanding the correlations between the sequence, function and the structure (Eijsink *et al.*, 2004; Khurana *et al.*, 2011).

Several reasons have been ascribed superior stability of thermophilic proteins including greater hydrophobicity (Mozhaev Martinek, 1984; Haney et al., 1997; Kumar et al., 2000) better packing, shorter loops (Russell et al., 1997), smaller and less numerous cavities (Salminen et al., 1996; Kumar et al., 2000), less uncharged polar residues (Haney et al., 1999; Fields, 2001), increased occurrence of proline residues (Bogin et al., 1998; Kumar et al., 2000), and increased hydrogen bonding (Vogt and Argos, 1997).

Serratia marcescens is one of the pioneer bacterium in chitin degradation (Watanabe et al., 1997; Bjørk et al., 2003). S. marcescens B4A chitinase (Babashpour et al., 2012) plays an important role in its plant antiproperties. 3D-Structure pathogenic analysis of this enzyme suggests that Serine 390 substitution of with could Isoleucine improve its thermostability by shortening the loop size and increasing the  $\beta$ -sheet length. To examine the assumption, Quik Change Site Directed Mutagenesis (QCSDM) was employed to modify S. marcescens B4A chitinase structure and its thermostability. The outcome of the research has been presented in this paper.

#### Materials and methods

Materials

The plasmid pOE60 chitinase (previous work) (Babashpour et al., 2012; Zarei et al., 2012) was used for amplification of chitinase gene. Escherichia coli strain DH5α (Invitrogen, Carlsbad, CA, USA) was used as the host for molecular cloning of recombinant pTZ57R/T (Fermentas. Waltham, Massachusetts, USA). pET26b (Fermentas. Waltham, USA) Massachusetts, plasmids harboring native and mutant chitinase respectively. BL21 gene, Escherichia coli strain (Invitrogen. Carlsbad, CA, USA) was also used for protein expression. Flake crab shell chitin and 3, 5-dinitrosalicylic acid (DNS) were purchased from Sigma (St. Louis, MO, USA). Colloidal chitin was prepared according to the modified method of Roberts and Selitrennikoff and use of commercial chitin (Takiguchi, 1991).

In silico studies

Homology modeling and 3D-structure analysis

The amino acid sequence of chitinase retrieved from NCBI database was used as a template in BLASTP search against PDB to find appropriate templates for homology modeling. Amino acid sequence of *Serratia marcescens* B4A chitinase (accession number HM473183) was retrieved from the GenBank database (http://www.ncbi.nlm.nih.gov).

BLASTP search (Altschul *et al.*, 1997) of a query (chitinase sequence) was

performed against Protein Data Bank (PDB). Homology modeling for the native and the mutant chitinase of Serratia marcescens B4A was performed by MODELLER v9.10 (http://www.salilab.org/modeller/) (Webb and Sali, 2014) using opened form of chitinase (1edq A) downloaded from PDB server as the template. The MODELLER generated structure of the mutant chitinase was further analyzed by Ramachandran analysis generated by Procheck (Laskowski et al., 1993), Qmean server (Benkert et al., 2008), ProSA-web (Wiederstein and Sippl, 2007), and RMSD value calculation (Humphrey et al., 1996). Omean is a parameter (between 0 and 1) for assessment of model credibility and ProSA shows quality score of the predicted model in the context of all known protein structures. Furthermore, the structural similarity between the native and mutant chitinase was checked by the **RMSD** value calculation using **VMD** software (Humphrey et al., 1996).

# Experimental studies

Mutagenesis

The Quik Change Site Directed Mutagenesis strategy was used for mutagenesis. The plasmid pOE60 harboring the chitinase gene (previous work) (Babashpour et al., 2012; Zarei et al., 2012) was used as the template for producing the mutant chitinase. The forward (5'-CACATCTTCCTGATGATCTACGAC TTCTACG-3') (5' and reverse

# CGTAGAAGTCGTAGATCATCAGG

AAGATGTG-3') mutant primers were used in polymerase chain reaction. PCR product was digested with 1-2 unit of *Dpn*I (Jena. science. Germany) at 37°C for 30 min. The correctness of mutation in the chitinase gene was verified by sequencing of the chitinase gene.

# Cloning and expression

For reinforcement of the mutant chitinase gene and insertion of restriction sites in two sides of gene were designed two oligonucleotide primers, a forward primer (Emr-F-chit: 5'-

ATAATCATATGGGGCGCAAATTT AATAAACC-3', NdeI site with ATG as the start codon) and reverse primer 5'-(Emr-R-chit: ATATACTCGAGATCTTTGAACGC CGGCGC-3', XhoI site). Following, the PCR product (1,701 bp) was cloned pTZ57R/T then recombinant into plasmid was digested with NdeI and XhoI and the restricted segments were isolated and ligated into the pET26b expression vector which has been digested with the same enzymes. A single colony of recombinant plasmid carrying the mutant chitinase which had been transformed into BL21 (DE3) Escherichia coli strain was cultured in LB medium. Therefore protein expression was induced by addition of 1 isopropyl-cD-thiogalactoside mM (IPTG) and was verified by SDS-PAGE analysis. The protein content of the determined extracts was by the Bradford method (Bradford, 1976).

#### Enzyme activity

Chitinase activity was assayed by the modified method of Roberts Selitrennikoff (Takiguchi, 1991) using colloidal chitin (1% w/v chitin in 20 mM phosphate buffer, pH 7.5) as the substrate. The enzymatic reaction was run for 60 min at 50°C. To stop the reaction, the mixture was boiled. Then, DNS (1% w/v) was added and the resulting mixture was maintained at 100°C for 10 min. The extent of the enzymatic hydrolysis was evaluated spectrophotometrically 540 at (Miller, 1959).

Stability and kinetics of the native and mutant recombinant chitinases

To determine the optimum pH, the chitinases activity was measured at various pH (s) ranging from 3–12 using 50 mM glycine–acetate–phosphate buffer. The enzyme stability at various pH(s), was determined by preincubation of the enzyme at the desired pH in the absence of the substrate for 90 min, followed by the enzyme assay at optimum pH.

To determine the optimum temperature, the enzymatic reaction was run in the presence of the substrate at different temperatures between 10–90°C in 20 mM phosphate buffer (pH 6) for 45 min, followed by the enzyme assay at pH and °C.

Thermal stability was measured by two methods: (1) measurement of the thermal stability at different temperatures. To do so, the enzyme was incubated at different temperatures ranging from 50 to 90°C for 90 min, then colloidal chitin was added, and the enzyme assay was done at 50°C (optimum temperature). Measurement of the thermal stability at different times at two temperatures of 50 and 60°C. Every 20 min (up to 120 min), the reaction mixture was sampled and the samples were placed on ice for 30 min, then colloidal chitin was added to assay the enzyme at 50°C (optimum temperature).

Kinetics parameters of the chtinases were calculated by double-reciprocal analysis of the corresponding kinetics curves obtained from the corresponding enzymatic reactions in the presence of various concentrations of the substrate ranging from 0 to 50 mg mL<sup>-1</sup> at pH and °C (Lineweaver and Burk, 1934).

#### Results

Homology modeling and structure validation

Using Serratia marcescens B<sub>4</sub>A chitinases (PDB: HM473183) as the reference structure in the computational studies, it was found that Ser390 could be more important to the structural stability of HM473183 as compared with the other suggestions. Therefore, substitution of Ser390 with Ile was further studied. This substitution not only decreases the number of amino acids with polar residues, but it also increases the chance of the beta sheet extension in the protein skeleton since Ile, as a branched amino acid, has conformational preference for the beta sheet structure. As a result,

suggested substitution was anticipated to shorten the loop section (Fig. XX), where ser390 was placed, in order to enhance the structural stability through the reduction of the entropy of the whole molecule. To appraise the change in protein stability after having the substitution. the thermodynamic stability ( $\Delta\Delta G$ ) using flexible and fixed backbone method was measured by Eris server (Yin et al., 2007). This resulted in  $\Delta\Delta \mathbf{G}$  of 2.18 kcal mol<sup>-1</sup> for the chitinase. Moreover. mutant appropriateness of the mutation site was verified by an automated web site: http://bioinformatics.org/primerx

(Zheng et al., 2004).

Homology modeling was performed to predict the 3D structure (opened form) of the mutant. Based on the sequence identity between the query and the template protein sequence, the PDB number of 1edq\_A with 99% identity was recognized for prognostication of the opened form of both chitinases. Then, the 3D structures of native and mutant chitinase were optimized using MODELLER v9.10 (Fig. 1). ProSA and QMEAN scores were found to be -9.75 and 0.837 (Z-score=0.88) for the mutant chitinase and -9.56 and 0.84 (Zscore=0.91) chitinase. for native respectively.

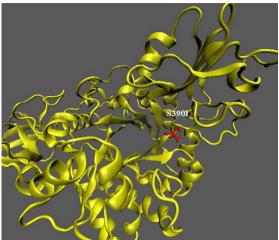


Figure 1: Optimized 3D structure of the opened form of mutant chitinase generated by MODELLER v9.10 using 1edq\_A as the template. The protein structures are shown as new cartoon representation. Visualization was performed using VMD software.

Ramachandran plot demonstrated that 93.7 % of the amino acid residues of the mutant chitinase were in the most favored region and 6.3 % in the additional allowed region. Superimposition of native and the mutant resulted in RMSD value of 2.9 Å for the  $C\alpha$  atoms. These results indicated that the structures of the mutant and template are similar and the homology model is reliable.

## Mutagenesis and expression

After mutagenesis with the QCSDM strategy, the accuracy of mutation was verified by DNA sequencing. Then, the mutant chitinase gene was sub cloned in pET26b expression vector which resulted in successful expression of the enzyme in BL21 (DE3) *Escherichia coli* strain. The protein produced a 62 kD band on SDS-PAGE (Fig. 2).

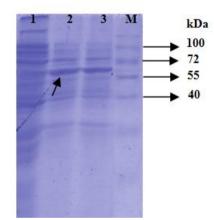


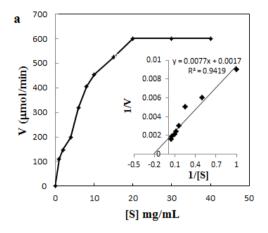
Figure 2: SDS-PAGE analysis of native and mutant chitinase. Lanes 1 total recombinant before induction, 2-3 native and mutant chitinase after induction (filled arrow is 62 kDa band refer to chitinase), M protein size marker.

# Kinetics parameters

Measurement of the Michaelis constant was showed that the  $K_m$  values of native and mutant chitinases were 4/5 mg mL (Fig. 3a) and 4/7 mg mL<sup>-1</sup> (Fig. 3b), respectively. Very little different of the  $K_m$  values of native and mutant

chitinase showing that the affinity of enzyme towards the substrate has not been changed. The  $V_{max}$  value of native and mutant chitinase were observed to

be 588/23 Unit ( $\mu$ M min<sup>-1</sup>) of protein (Fig. 3a) and 153/84 Unit ( $\mu$ M/min) of protein (Fig. 3b), respectively.



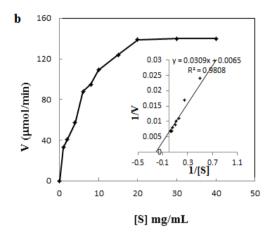


Figure 3: Michaelis—Menten curve of native and mutant chitinase activity. *Inset* Lineweaver—Burk plot. a) The Km values and the Vmax value of native chitinase were found to be 4/5 mg mL<sup>-1</sup> and 588/23 Unit (µmol min<sup>-1</sup>), respectively. b) The Km values and the Vmax value of mutant chitinase were found to be 4/7 mg/ml and 153/84 Unit (µmol min<sup>-1</sup>) of protein, respectively.

#### Effect of pH and Temperature

The optimal pH(s) for native and the mutant chitinases were found to be 6 and 5, (Fig. 4a), respectively. However, the enzymes were stable between pH range of 6-10 for native and 5-10 for the mutant chitinase (Fig. 4b). The optimal temperatures of 50°C and 45°C were obtained for native and the mutant, (Fig. 4c), respectively.

#### Thermal stability

Thermal stability of chitinases was measured at different temperatures.

Although the activities of both chitinases were reduced with the rise of temperature, native chitinase activity was reduced faster and reached zero at 70°C, while the mutant chitinase could maintain about 20 percent of its activity (Fig. 5a). Measurement of temperature stability during the time showed that the native enzyme activity at 50 and 60°C was decreased, but the activity of mutant chitinase increased over the time (Fig. 5b).

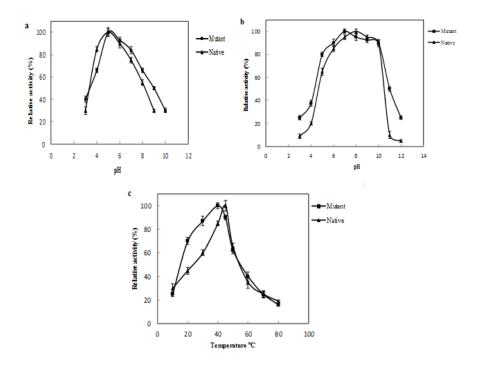


Figure 4: Effects of pH and temperature on the activity and stability of the native and mutant chitinase. a) Effect of pH on enzymes activity, enzymes assays performed over a range of pH values (3–10) using colloidal chitin as substrate under standard assay conditions. The optimal pH for the chitinase activity of native and mutant chitinases was found to be 6 and 5 respectively. b) Effect of pH on enzyme stability, enzymes preincubated at various levels of pH (3–12) without the substrate for 90 min. Following adjustment of the enzymes to pH 6, colloidal chitin was added and then enzymes assays performed under standard assay conditions. The enzymes were stable between pH range of 6-10 for native and 5-10 for mutant chitinase. c) Effect of temperature on enzymes activity, enzymes assays performed in the range of 10–90°C in pH 6 using colloidal chitin as substrate under standard assay conditions. The optimal temperature for enzyme activity of native and mutant was found to be 50°C and 45°C, respectively.

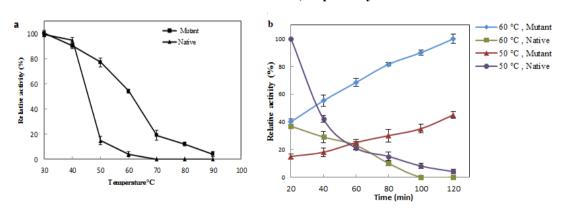


Figure 5: Thermal stability of chitinase. a) Effect of different temperatures on the activity and stability of native and mutant chitinase after incubation for 90 min. b) Effect of temperature on the stability of native and mutant chitinase after incubation at 50, 60°C in various time intervals.

#### Discussion

Research on the thermostable chitinases attractive for finding efficient enzymes with more usage in industrial process (Roodi et al., 2017). Recent studies clearly exhibit that the kinetic stability of proteins might be better by rational design, but that it may be difficult to exactly determine which region and amino acid of the protein should be considered. There are a commonplace strategy for increasing thermal protein stability through shortening of exposed loop regions which leads to decreasing the entropy and elevating the free energy of protein unfolding (Thompson and Eisenberg, 1999). Furthermore other researchers showed that the proportion of beta sheet amino acids is meaningfully high in thermophiles. This result might be obtained through enhancing hydrogen bonding which leads to increased stability (Chakravarty and thermal Varadarajan, 2000). In this study we wanted to stabilize the chitinase by reduction of the loop length and increasing of beta sheet length altogether. For this purpose determined amino acids that locate in the border of the loop and beta sheet. and considered that candidate sites did not have unfavorable steric interactions.

Also researchers showed that for increasing thermal stability, declining uncharged polar amino acids is very significant. The elimination of these residues may also reduce deamination of asparagine and glutamine by serine and threonine at high temperatures

(Vogt and Argos, 1997; Haney et al., 1999). Serine due to its short side chain mostly involve in local interactions, but large side chains amino acids generally contribute in short and long range interactions (Hornby, 1993; Kumar et al., 2000). Thus we selected Serine 390 resides in the end of the loop and replaced it with Isoleucine residue because of several reasons: Isoleucine residue tends to place in beta sheet and this replacement causes the reduction of the loop length and enhancement of the beta sheet length. Furthermore, increase of protein hydrophobicity stabilizez thermophile proteins (Mozhaev and Martinek. 1984: Hanev et al., 1999). Also entropy of unfolding dependent on the kind of the side chain, for example in this case flexibility of the selected loop is decreased by the presence of long or bulky side chains (Némethy et al., 1966). Also studies showed that polar uncharged residue in comparison of one non-polar is less favorable in thermostable protein (Mozhaev 1984; Chakravarty Martinek, and Varadarajan, 2000). Our studies demonstrated that mutant chitinase has been compacted and required time and temperature for its activity. Native and mutant chitinase kinetic studies showed that mutant chitinase had more activity than the native type because the native chitinase could not tolerate the assay this condition condition. but favorable for the mutant chitinase to increase its flexibility and exposed catalytic site for substrate. Because the enzyme was partially purified, in

chitinase activity assay two negative controls were used. The first negative control was cocktail without chitinase; second negative control cocktail with BL21 containing pET26b bacterial cell extract. Any enzymatic activity was observed in two negative controls. Alteration in enzyme binding site may lead to increase in K<sub>m</sub> (Fields and Somero, 1998; Fields, 2001). In our study the brief change of mutant chitinase K<sub>m</sub> showed that the enzyme substrate affinity was similar to the native enzyme. The  $V_{\text{max}}$  of mutant smaller chitinase is than native chitinase because the calculation of the V<sub>max</sub> value was performed in routine condition while mutant chitinase was rigided and at this condition showed low activity. Reduction of mutant chitinase V<sub>max</sub> without change of K<sub>m</sub> was similar to the noncompetitive inhibition. Studies have shown that there is correlation between thermal stability and the optimum temperature (Danson et al., 1996; Daniel et al., 2001). **Optimum** temperature increased in several engineered thermostabilized enzymes and doesn't change in the others (Thomas and Scopes, 1998; Bjørk et al., 2003). In our research the optimum temperature and pH didn't change remarkably for both the mutant and native chitinases, indicating that mutation did not affect the main characteristic of the enzyme. pH stability of the mutant and native chitinases were the same, but thermal stability was increased remarkably. Mutation caused the chitinase to tolerate high temperatures up to 90°C and it remained stable at 50°C, 60°C for up to 120 minutes. This characteristic is very important for industrial and biotechnological applications.

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