Non-Structured Amino-Acid Impact on GH11 Differs from GH10 Xylanase

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Abstract
The Aspergillus niger xylanase (Xyn) was used as a model to investigate impacts of un-structured residues on GH11 family enzyme, because the β-jelly roll structure has five residues (Ser1Ala2Gly3Ile4Asn5) at N-terminus and two residues (Ser183Ser184) at C-terminus that do not form to helix or strand. The N- or/and C-terminal residues were respectively deleted to construct three mutants. The optimal temperatures of XynΔN, XynΔC, and XynΔNC were 46, 50, and 46°C, and the thermostabilities were 15.7, 73.9, 15.5 min at 50°C, respectively, compared to 48°C and 33.9 min for the Xyn. After kinetic analysis, the substrate-binding affinities for birch-wood xylan decreased in the order XynΔC>XynΔNC>XynΔN, while the Kcat values increased in the order XynΔC<XynΔNC<Xyn<XynΔN. The C-terminal deletion increased the GH11 xylanase thermostability and Topt, while the N- and NC-terminal deletions decreased its thermostability and optimal temperature. The C-terminal residues created more impact on enzyme thermal property, while the N-terminal residues created more impact on its catalytic efficiency and substrate-binding affinity. The impact of non-structured residues on GH11 xylanase was different from that of similar residues on GH10 xylanase, and the difference is attributed to structural difference between GH11 jelly-roll and GH10 (ββ)8.

Introduction
Enzyme is widely used in industrial biocatalysis, such as fermentation, bio-energy production, food, beverage, paper and pulp, etc. However, the higher temperatures demand for robust enzymes. Many strategies have been used to improve enzyme thermostability. For example, disulphide-bonds were introduced to the Thermomonospora fusca xylanase [3]. Homologous segments were recombined between two Streptomyces lividans xylanases [4]. The N-terminus of mesophilic S. olivaceoviridis xylanase was substituted with the homologous region of thermophilic Thermomonospora fusca xylanase [5]. Disulphide-bond was introduced to the T. reesei xylanase internal region and the Thermomyces lanuginosus xylanase N-terminus [6,7].

From structural view point, enzyme is composed of irregular segment, α-helix, and β-strand. The later two are known as regular secondary structural elements. Unlike regular secondary structures, irregular segments are composed of amino-acids that are referred to as non-structured residues. Compared with stabilizing impact of regular secondary structures on enzyme thermostability [8,9], impact of non-structured residue is seldom investigated. Recently, the non-structured residues were shown to disturb the Aspergillus niger GH10 xylanase thermostability and catalytic activity [10]. However, the GH10 xylanase exhibits a (β/α)8 structure, which represents only one family of enzymes, and the (β/α)8 structure consists mainly of α-helices and β-strands [11]. The issue remains unsolved whether the non-structured residues have a general role or not in other family of enzymes. GH11 xylanase is a big family, and its β-jelly roll structure consists of 6% enzymes [12]. As a model of GH11 family enzyme, an xylanase (Xyn) was cloned from A. niger (GenBank: EU375728) [13]. The 185 amino-acid enzyme is the smallest GH11 xylanase, and exhibits the typical β-jelly roll structure (PDB: 1UKR) [14,15], unlike (ββ)8 structure of GH10 xylanase. The GH11 Xyn structure resembles a partially closed right hand, which is composed of two large β-pleated sheets and one helix [12]. The sheets are composed of anti- and parallel strands connected by irregular segments, and the only one helix is regarded to have stabilizing contribution to enzyme thermostability [8,9].

Results
Construction of the Mutants
The XynΔN, XynΔC, and XynΔNC genes were amplified and cloned to the pET20b (Fig. 1). After sequence accuracies of the extracted recombinant plasmids had been confirmed by DNA
sequencing, the transformed cells containing accurate recombinant plasmids, pET20b-xynΔn, pET20b-xynΔc, and pET20b-xynΔnc, were induced to express xylanases. The proteins, XynΔN, XynΔC, and XynΔNC, appeared on the SDS-PAGE gel as discrete bands at 29 kDa (Fig 1, Table 1). The larger apparent molecular masses are attributed to the xylanase having acidic property and a C-terminal His6 tag. The xylanase pI is 4.42, and its optimal reaction pH, pHopt, is 3.8. Acidic protein was found to bind less SDS, and therefore had a larger apparent molecular mass [17–19].

Enzyme Properties

The impacts of non-structured residues on enzyme properties were determined by assaying the mutants in parallel with the wild xylanase. Each data point was assayed for three independent reactions. The pH opt values of three mutants were 3.6 in imidazole-biphthalate buffer, 0.2 units lower than the Xyn (Fig 2, Table 1). The optimal reaction temperatures, T opt values, of XynΔN, XynΔC, and XynΔNC were calculated to be 13.7, 15.5, and 15.5 min, respectively. Both XynΔN and XynΔNC were ~50% thermostable as the Xyn; whereas, the XynΔC was 2.2-times more thermostable than the Xyn (Fig 3, Table 1). The C-terminal residue deletion increased enzyme thermostability, while the N- and NC-terminal residue deletions decreased enzyme thermostability. The increasing impact of C-terminal deletion on enzyme T opt and thermostability was counteracted by the decreasing impact of N-terminal deletion, showing that the impact of N-terminal residues on enzyme thermal properties was bigger than that of the C-terminal residues.

When enzyme kinetics were determined at pHopt and T opt conditions, the substrate-binding affinities for birch-wood xylan were determined and fitted with the Arrhenius equation. The thermal in-activation half-lives (t1/2), thermostabilities, of XynΔN, XynΔC, and XynΔNC and Xyn, were amplified with the Arrhenius equation. The thermal in-activation half-lives (t1/2), thermostabilities, of XynΔN, XynΔC, and XynΔNC and Xyn, were amplified with the Arrhenius equation. The thermal in-activation half-lives (t1/2), thermostabilities, of XynΔN, XynΔC, and XynΔNC and Xyn, were amplified with the Arrhenius equation. The thermal in-activation half-lives (t1/2), thermostabilities, of XynΔN, XynΔC, and XynΔNC and Xyn, were amplified with the Arrhenius equation. The thermal in-activation half-lives (t1/2), thermostabilities, of XynΔN, XynΔC, and XynΔNC and Xyn, were amplified with the Arrhenius equation. The thermal in-activation half-lives (t1/2), thermostabilities, of XynΔN, XynΔC, and XynΔNC and Xyn, were amplified with the Arrhenius equation. The thermal in-activation half-lives (t1/2), thermostabilities, of XynΔN, XynΔC, and XynΔNC and Xyn, were amplified with the Arrhenius equation. The thermal in-activation half-lives (t1/2), thermostabilities, of XynΔN, XynΔC, and XynΔNC and Xyn, were amplified with the Arrhenius equation. The thermal in-activation half-lives (t1/2), thermostabilities, of XynΔN, XynΔC, and XynΔNC and Xyn, were amplified with the Arrhenius equation.
terminal deletion created intermediate impact on enzyme catalytic efficiency. Thus, the increasing impact of N-terminal deletion on enzyme catalytic activity was counterbalanced by the decreasing impact of C-terminal deletion. The N-terminal deletion produced more impact on enzyme kinetic parameters.

Structural Analysis

The five N-terminal non-structured residues created more impact on the enzyme substrate-binding affinity and catalytic activity, while the two C-terminal ones created more impact on its thermostability. To explain the different impacts of N- and C-terminal residues on enzyme thermal and catalytic properties, the *A. niger* xylanase structure was drawn according to the PDB data (1UKR, Fig 4) [14,15]. Two active-site residues of the enzyme are Glu79 and Glu170. The nucleophile residue, Glu79, is held in place by interactions to Gln129 and Tyr70. The acid-base catalyst residue, Glu170, is hydrogen-bonded with Tyr81 and Asp37. According to the analysis of *A. niger* xylanase structure [14], substrate-binding sites of the Xyn are Gln129, Tyr70, Tyr72, Tyr81, and Asp37. The N-terminal non-structured residues, Ser1Ala2Gly3Ile4Asn5, are closer to the substrate-binding residues, therefore, have more impact on enzyme kinetics. The C-terminal non-structured residues, Ser183Ser184, are more distal to the substrate-binding residues, but closer to the C-terminal strand, therefore, have less impact on enzyme activity, but more impact on thermal properties.

Discussion

Rational Deletion of Terminal Non-structured Residues

Previously, either the entire domains were truncated completely, or the terminal residues were deleted successively [20–26]. Unlike that, the GH11 xylanase seven terminal non-structured residues were rationally deleted based on structural analysis, because the residues do not form to regular secondary structural elements. The N-terminal deletion decreased enzyme $T_{\text{opt}}$ for 2°C and thermostability for 50%, while the C-terminal deletion increased enzyme $T_{\text{opt}}$ for 2°C and thermostability for 2.2-times. This is consistent with the homologous recombination analysis of *S. lividans* xylanase [4]. The deletion of N-terminal helix also increased the N-glycanase deglycosylation activity [22]. The successive truncation analysis showed that the seven N-terminal and the three C-terminal residues were unnecessary for the *Clostridium thermocellum* lichenase activity [23]. The five amino-acid mutations at N-terminus might confer structural stability, and hence, prevented the overall thermal unfolding of mesophilic *S.*

| Xylanase | pH$_{\text{opt/pI}}$ | $T_{\text{opt}}$ (°C) | $t_{1/2}$ (50°C) (min) | $V_{\text{max}}$(umol/l/s)/$K_{\text{m}}$(mg/ml)/$K_{\text{cat}}$(1/s) | MM(kDa) | Apparent/ Theoretical | Number of amino acids |
|----------|-------------------|---------------------|-----------------------|---------------------------------------------------------------|---------|---------------------|----------------------|
| Xyn      | 3.8 ±0.3/4.42     | 48±0.1              | 33.9                  | 14.6±3.2/18.6±4.4±200                                         | 29/21.1 | 184                 |
| XynAN    | 3.6 ±0.3/4.42     | 46±0.2              | 15.7                  | 16.1±3.9/38.8±11.7/222.7                                      | 28/20.6 | 179 (sagin)         |
| XynAC    | 3.6 ±0.2/4.42     | 50±0.3              | 73.9                  | 10.6±6.6/6.6±6.6±158.9                                      | 28/20.9 | 182 (sagin)         |
| XynNC    | 3.6 ±0.2/4.42     | 46±0.2              | 15.5                  | 12.2±6.0/25.2±6.0±182.0                                      | 28/20.4 | 177 (sagin)(ss)     |

Note: $T_{\text{opt}}$: optimal reaction temperature for activity, $t_{1/2}$: thermal in-activation half-life, after incubation at 50°C for different intervals, residual activity was determined and expressed as a ratio to the un-incubated enzyme, and data were fitted with Arrhenius function. The $t_{1/2}$ was calculated according to the decay function to indicate enzyme thermostability. Using the Student’s t-Test ($p<0.05$), the XynAC is more thermostable than the Xyn, and it is more thermostable than the XynAN and the XynNC. The latter two enzymes are statistically similar thermostable. Kinetics was determined for birch-wood xylan at each enzyme pH$_{\text{opt}}$ and $T_{\text{opt}}$ conditions, and the data were fitted with Hill function to calculate $V_{\text{max}}$ and $K_{\text{m}}$. doi:10.1371/journal.pone.0045762.t001

**Figure 2.** Xylanase optimal pH and optimal temperature. The pH$_{\text{opt}}$ was determined from pH 3.2 to 4.2 at 0.2 unit interval in 50 mM imidazole-biphthalate buffer (left). The $T_{\text{opt}}$ was determined from 42 to 54°C at 2°C interval (right).

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Figure 3. Xylanase thermostability and kinetics. After incubation at 50°C at 10-min interval from 10 to 30 min, the residual activity was assayed and expressed as a ratio relative to the un-incubated xylanase activity. The data were fitted with the equation \( y = A \ e^{-kt} \) (Origin, version 8.0), and thermostability \( (\ln A/\ln 2) \) was calculated according to the decay function. The kinetics were assayed at \( T_{opt} \) and \( pH_{opt} \) conditions using birch-wood xylan at concentrations from 0 to 13 mg/ml. The data were fitted with the Hill function to calculate maximal activity \( (V_{max}) \) and \( K_m \) (Origin, version 8.0).

Different Impacts of Non-structured Residues between GH11 and GH10 Xylanase

Very recently, we investigated the impacts of similar non-structured residues on the GH10 xylanase. The N- or/and C-terminal deletions increased enzyme thermostability for \( \sim 2 - 4 \)-fold and activity for 4 - 7-fold [10]. The N-terminal deletion decreased the GH10 xylanase \( T_{opt} \) for 6°C, but the C-terminal deletion increased its \( T_{opt} \) for 6°C. The impacts of N- and C-terminal deletions were opposite on the enzyme \( T_{opt} \), but additive on its thermostability. The impact of five N-terminal residues was more on the GH10 enzyme kinetics, but less on its thermostability than that of the one C-terminal residue. Compared with the two investigations, C-terminal deletion increased both GH10 and GH11 enzyme \( T_{opt} \) values and thermostabilities, and N-terminal deletion decreased both enzyme \( T_{opt} \) values. Whereas, N-terminal deletion decreased the GH11 enzyme thermostability, but increased the GH10 enzyme thermostability. The impacts of N- and C-terminal deletion were additive on GH10 enzyme thermostability, but opposite on GH11 enzyme thermostability. The inconsistence was attributed to the structural difference between GH10 (\( \beta/\alpha \)) and GH11 \( \beta \)-jelly roll. The \( \beta/\alpha \) structure is mainly composed of helices and strands, with one \( \beta/\alpha \) as a repeat unit; while the \( \beta \)-jelly roll structure is composed of many strands and only one helix, with one strand as a repeat unit. Unlike \( \beta/\alpha \), \( \beta \)-jelly roll does not easily fold to active conformation. Moreover, the \( \beta/\alpha \) structure is relatively more thermostable than the \( \beta \)-jelly roll. Probably, N-terminal deletion seriously disturbed the GH11 folding, but slightly disturbed the GH10 enzyme, because helix folds more easily than strand. The N-terminal non-structured residues are seated at upstream of strand, and probably, act as leading element for its folding. The C-terminal non-structured residues are seated at downstream of strand, therefore, scarcely disturb its folding.

At present, more and more structures were crystallized and deposited in PDB. As to those un-crystallized enzymes, structures can be modeled through comparative homologous modeling. All these progresses provide more structural information for enzyme rational engineering. According to the investigations of non-structured residue impact on GH11 and GH10 xylanases, N- and C-terminal non-structured residues are unnecessary for activities of both \( \beta/\alpha \) and \( \beta \)-jelly roll enzymes. Strategy for rational enzyme engineering is proposed as that thermostability can be improved by deleting C-terminal non-structured residues, and catalytic efficiency can be increased by deleting N-terminal non-structured residues.

Materials and Methods

Materials and Reagents

The \( A. \ niger \) Xyn gene (GenBank: EU375728), which encodes a 185-residue mature xylanase, was cloned into pET20b(+) (Novagen, Shanghai, China). Molecular biology reagents, including \( \Phi \) polymerase, restriction endo-nucleases NdeI and XhoI, T4 DNA ligase, DNA marker, protein marker, were purchased from Takara Inc (Dalian, China).

Construction of the Mutants

Based on structural analysis, the non-structured residues, Ser1Ala2Gly3Ile4Asn5 and Ser183Ser184, were respectively deleted to construct three mutants, Xyn\( \Delta \)N, Xyn\( \Delta \)C, and Xyn\( \Delta \)NC. PCR was carried out using 16.5 \( \mu \)g of pET20b-xyn, 1.0 \( \mu \)mol of each of two related primers, 5 U of \( \Phi \) polymerase, 4.0 \( \mu \)mol of dNTPs, and 1× polymerase buffer with the following thermal cycling: 4 min denaturation at 94°C, followed by 30 cycles of 1 min denaturation at 94°C, 1 min annealing at temperature given below, and 1 min extension at 72°C. The reaction was extended for 5 min at 72°C and cooled subsequently to 4°C.

olivoacarii xylanase [5]. The introduction of disulphide-bond showed that the flexible N-terminus disturbed enzyme thermostabilities [1–3,6,7]. The N- and C-terminal deletions created opposite effects on the enzyme thermal properties, thereby, the Xyn\( \Delta \)N\( C \) and thermostability decreased for 2°C and for 50%. However, the N-terminal deletion increased enzyme catalytic activity. Structural analysis shows that the N-terminal residues are closer to the substrate-binding residues, therefore, probably involve in substrate-binding.

Non-structured residues on the GH10 \( \beta \)-jelly roll structure is relatively more thermostable than the GH11 \( \beta/\alpha \) structure. Probably, N-terminal deletion seriously disturbed the GH11 folding, but slightly disturbed the GH10 enzyme, because helix folds more easily than strand. The N-terminal non-structured residues are seated at upstream of strand, and probably, act as leading element for its folding. The C-terminal non-structured residues are seated at downstream of strand, therefore, scarcely disturb its folding.

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The XynD gene was amplified using p2/p3 and annealing at 33.6°C. The XynDC and XynDNC genes were amplified using p1/4 and p2/p4 and annealing at 35.8°C. The wild Xyn was amplified using p1/p3 and annealing at 37°C. The primer sequences used are as follows, with the underlined letters showing NdeI/XhoI restriction sites: p1: ggaattc catatg agtgccggtatca, p2: ggaattc catatg -tacgtgcaaaactac, p3: ctaaatta ctcgag agaggagatcgtga; p4:ccaaatta ctcgag gatcgtgacactgg.

Following PCR amplification, the genes were cloned into pET20b(+) plasmids that had been pre-digested with NdeI/XhoI to delete redundant endo-nuclease sites. The recombinant plasmids were transformed into E. coli BL21(DE3) competent cells, then extracted and sequenced with an ABI 3730 automated sequencer to confirm gene accuracy (Invitrogen Biotechnology, Shanghai, China). The transformed cell containing accurate recombinant plasmids were grown and induced to produce xylanases according to standard protocols [16]. A C-terminal His tag was included in the xylanase sequences to allow the proteins to be purified with Co²⁺-binding resin (Amersham Bioscience). Active fractions were pooled and further purified using sephadex G-25. The xylanases were detected using 12.5% polyacrylamide SDS-PAGE, stained with Coomassie brilliant G-250. Protein concentration was measured by a Spectrophotometer.

Figure 4. The Aspergillus niger Xyn structure. The N-terminal non-structured residues are Ser1Ala2Gly3Ile4Asn5, and the C-terminal non-structured residues are Ser183Ser184. Only the Gly3Ile4Asn5 and Ser183 were shown, because the deposited structure (PDB: 1UKR) did not contain the Ser1Ala2 at N-terminus and the Ser184 at C-terminus. The active-site residues are Glu79 and Glu170. The substrate-binding residues are Asp37, Trp72, Tyr79, Tyr81, and Gln129. The N-terminal residues are closer to the substrate-binding residues, therefore, have more impact on enzyme catalytic activity and substrate-binding affinity. The C-terminal residues are more distal to the substrate-binding residues but closer to the C-terminus, therefore, have less impact on its catalytic properties but more impact on its thermal properties.

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was calculated. The kinetics were assayed at \( T_{\text{opt}} \) and \( pH_{\text{opt}} \)
for xylanase activity. The data were fitted with the equation
\[
y = A e^{-kt}
\]
where \( A \) is the maximum activity and \( k \) is a kinetic constant. The optimal reaction temperature, \( T_{\text{opt}} \), was determined from 42 to 50 mM imidazole-biphthalate buffer at 0.2 unit interval. The enzymes were pre-incubated at 50°C, and the residual activity was assayed at 10-min interval from 10 to 30 min and expressed as a ratio relative to the un-incubated xylanase activity. The data were fitted with the equation \( y = A s^{-bt} \) (Origin, version 8.0), and the thermal inactivation half-life \( (1/2) \) was calculated. The kinetics were assayed at \( T_{\text{opt}} \) and \( pH_{\text{opt}} \) conditions for 5 min using birch-wood xylan at concentrations from 0 to 13 mg/ml (Sigma-Aldrich, Shanghai, China). The data were fitted with the Hill function to calculate maximal activity (\( V_{\text{max}} \)) and \( Km \) (Origin, version 8.0).

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Author Contributions
Conceived and designed the experiments: LL. Performed the experiments: XS. Analyzed the data: PY LW HC. Wrote the paper: LL.

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The xylanase standard activity was determined on birch-wood xylan using dinitrosalicylic acid method (DNS) [16]. Enzyme reaction was carried out for 16 min using 100 \( \mu \)L enzyme, 100 \( \mu \)L 1% birch-wood xylan, and 600 \( \mu \)L imidazole-biphthalate buffer, thereafter, 600 \( \mu \)L DNS was added and boiled. The reaction system was added to 5 mL with water, and absorbance was determined using UV2000 ultraviolet spectrophotometer at 550 nm.

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