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K.M.G.
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my)Protein engineering of selected residues
from conserved sequence regions of a
novel *Anoxybacillus* α -amylaseVelayudhan Ranjani¹, Štefan Janeček^{2,3}, Kian Piaw Chai¹, Shafinaz Shahir¹,
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The α -amylases from *Anoxybacillus* species (ASKA and ADTA), *Bacillus aquimaris* (BaQA) and *Geobacillus thermoleovorans* (GTA, Pizzo and GtamyII) were proposed as a novel group of the α -amylase family GH13. An ASKA yielding a high percentage of maltose upon its reaction on starch was chosen as a model to study the residues responsible for the biochemical properties. Four residues from conserved sequence regions (CSRs) were thus selected, and the mutants F113V (CSR-I), Y187F and L189I (CSR-II) and A161D (CSR-V) were characterised. Few changes in the optimum reaction temperature and pH were observed for all mutants. Whereas the Y187F ($t_{1/2}$ 43 h) and L189I ($t_{1/2}$ 36 h) mutants had a lower thermostability at 65 °C than the native ASKA ($t_{1/2}$ 48 h), the mutants F113V and A161D exhibited an improved $t_{1/2}$ of 51 h and 53 h, respectively. Among the mutants, only the A161D had a specific activity, k_{cat} and k_{cat}/K_m higher (1.23-, 1.17- and 2.88-times, respectively) than the values determined for the ASKA. The replacement of the Ala-161 in the CSR-V with an aspartic acid also caused a significant reduction in the ratio of maltose formed. This finding suggests the Ala-161 may contribute to the high maltose production of the ASKA.

Most α -amylases (EC 3.2.1.1) belong to family 13 of glycoside hydrolases (GH13)^{1,2}. The polyspecific family GH13 covering more than 30 different amyolytic and related enzyme specificities with nearly 16,000 sequences ranks among the largest of the GH families². It is also the main α -amylase family of the α -amylase CAZy clan GH-H that is formed by the families GH13, GH70 and GH77³⁻⁷. The family GH13 has been divided into curator-based subfamilies (currently 40)⁸, although some of them were previously established^{9,10}, and others have not yet been established¹¹. α -Amylase is a very common enzyme produced by most living organisms³. Because it is necessary for the digestion of complex starchy substrates in living systems¹², it is also important in various branches of industrial production^{13,14}. In addition to other bacterial producers, several *Anoxybacillus* species have been shown to produce α -amylases with potentially industry-desired properties including the ability to digest raw starch¹⁵⁻¹⁹.

Recently, we described the biochemical properties of two *Anoxybacillus* α -amylases produced by the closely related *Anoxybacillus* sp. SK3-4 (ASKA) and *Anoxybacillus* sp. DT3-1 (ADTA)¹⁸. Interestingly, although both these α -amylases belong to the main α -amylase family GH13, the phylogenetic tree clearly demonstrates that their cluster is independent of all existing GH13 subfamilies¹⁸. A few other α -amylases exhibiting close relatedness to ASKA and ADTA were reported, i.e., those from *B. aquimaris*¹¹ and *G. thermoleovorans*²⁰⁻²². This novel, currently undefined GH13 subfamily of α -amylases may be most closely related to subfamily GH13_1 of extracellular fungal α -amylases². It, however, can be distinguished from other GH13 subfamilies with the α -amylase specificity by the exclusive presence of two consecutive tryptophan residues (Trp-201 and Trp-202; *B. aquimaris* α -amylase numbering) positioned at helix α 3 of the catalytic TIM-barrel domain¹¹. Of the three α -amylases from *G. thermoleovorans*, two of them are identical (amyA from the strain Pizzo⁷ and GTA from the strain CCB US3 UF5)^{21,22} and share 94% sequence identity with the third one (Gt-amyII from the strain MTCC 4220)²⁰.

A preliminary inspection of the CSRs revealed that several positions are not conserved in ASKA when compared with α -amylase counterparts from other GH13 subfamilies. The main goal of the present study was to determine the function of some distinct amino acid residues in the ASKA CSRs using site-directed mutagen-



Table 1 | Sequences of the GH13 α -amylase subfamily represented in this study by ASKA. The length shown is the total number of amino acids

No.	Organism	GenBank	UniProt	Length	CAZy
1	<i>Anoxybacillus</i> sp. SK3-4 ¹⁸	AFI49455.1	I1VWH9	505	Yes
2	<i>Anoxybacillus</i> sp. DT3-1 ¹⁸	AFI49456.1	I1VW10	505	Yes
3	<i>Anoxybacillus</i> sp. GXS-BL	AEQ38578.1	G4Y5W9	505	Yes
4	<i>A. flavithermus</i> AK1	EMT46721.1	M8D7E8	505	No
5	<i>A. kamchatkensis</i>	WP_019417912.1	UPI0002F2C1C0	504	No
6	<i>A. flavithermus</i> TNO-09.006	ELK21122.1	M5JAR3	504	No
7	<i>A. flavithermus</i> NBRC 109594	GAC90171.1	R4F9M5	505	No
8	<i>A. flavithermus</i> DSM 21510/WK1	ACJ34547.1	B7GLU6	505	Yes
9	<i>Bacillus</i> sp. m3-13	WP_010192073.1	UPI0001E8973B	517	No
10	<i>Paenispodosarcina</i> sp. TG-14	WP_017381610.1	UPI0003096FF2	514	No
11	<i>Geobacillus</i> sp. GHH01	AGE21200.1	L7ZU32	511	Yes
12	<i>G. caldoxylosilyticus</i>	WP_017436518.1	UPI0002F1817B	510	No
13	<i>Geobacillus</i> sp. WCH70	ACS23590.1	C5D6S3	510	Yes
14	<i>Geobacillus</i> sp. POT5	ABL77406.1	A8QL62	514	Yes
15	<i>Ornithinibacillus scapharcae</i>	WP_010094264.1	UPI000225AA66	506	No
16	<i>B. oceanisediminis</i>	WP_019381460.1	UPI0002D7E70C	512	No
17	<i>B. infantis</i> NRRL B-14911	AGX03407.1	U5L9V5	513	Yes
18	<i>Bacillus</i> sp. 2_A_57_CT2	EFV78894.1	E5WEA3	512	No
19	<i>G. kaustophilus</i> GBlys	GAD14452.1	U2X6P3	513	No
20	<i>Geobacillus</i> sp. G11MC16	EDY07654.1	B4BIQ2	511	No
21	<i>Geobacillus</i> sp. GXS1	ACK58047.1	B7UDC2	513	Yes
22	<i>Geobacillus</i> sp. Y412MC61	ACX78152.1	C9RXK9	511	Yes
23	<i>B. firmus</i> DS1	EWG12544.1	UPI0003EB955E	512	No
24	<i>Geobacillus</i> sp. MAS1	ESU72338.1	V6VD05	513	No
25	<i>G. thermoleovorans</i> CCB_US3_UF5 ²¹	AEV18110.1	G8N704	511	Yes
26	<i>Clostridiaceae bacterium</i> L21-TH-D2	EOD00789.1	R1CVX1	496	No
27	<i>Geobacillus</i> sp. C56-T3	ADI27796.1	D7D2F9	511	Yes
28	<i>Geobacillus</i> sp. JF8	AGT31044.1	S5YWG2	511	Yes
29	<i>G. thermoleovorans</i> NP54 ²⁰	AFK08971.1	I3QI14	513	Yes
30	<i>Geobacillus</i> sp. WSUCF1	EPR28542.1	S7SUP4	513	No
31	<i>G. kaustophilus</i> HTA426	BAD74992.1	Q5L238	513	Yes
32	<i>Bacillus coahuilensis</i>	WP_010171518.1	UPI0001850CB5	445	No
33	<i>G. thermodenitrificans</i> NG80-2	ABO65996.1	A4IKZ2	511	Yes
34	<i>B. boroniphilus</i> JCM 21738	GAE45850.1	W4RN42	510	No
35	<i>B. aquimaris</i> MKSC 6.2 ¹¹	AER68125.1	G8JA7	512	Yes
36	<i>Spirochaeta</i> sp. L21-RPul-D2	AHC13589.1	V5WDG2	490	Yes

esis. These residues including Phe-113 in CSR-I, Tyr-187 and Leu-189 in CSR-II, and Ala-161 in CSR-V (ASKA numbering) were substituted with valine, phenylalanine, isoleucine, and aspartic acid, respectively, which are more conserved in other GH13 subfamilies containing the α -amylase specificity.

Results

Analysis of the conserved sequence regions of α -amylases. ASKA, ADTA, Pizzo, GtamyII, GTA and BaqA α -amylases phylogenetically

cluster together, creating a new GH13 subfamily. Other homologs of this subfamily are summarised in Table 1, based on the BLAST results (see the Methods section). Most of these α -amylases are not well characterised. Using the thirty-six α -amylases shown in Table 1, seven conserved sequence regions (CSRs) typical for the members of this new α -amylase family GH13²³ were identified in these sequences and the weblogo covering the CSRs was created (Figure 1). Compared with the representatives of this new GH13 subfamily, ASKA and all its closely homologous subfamily counterparts have

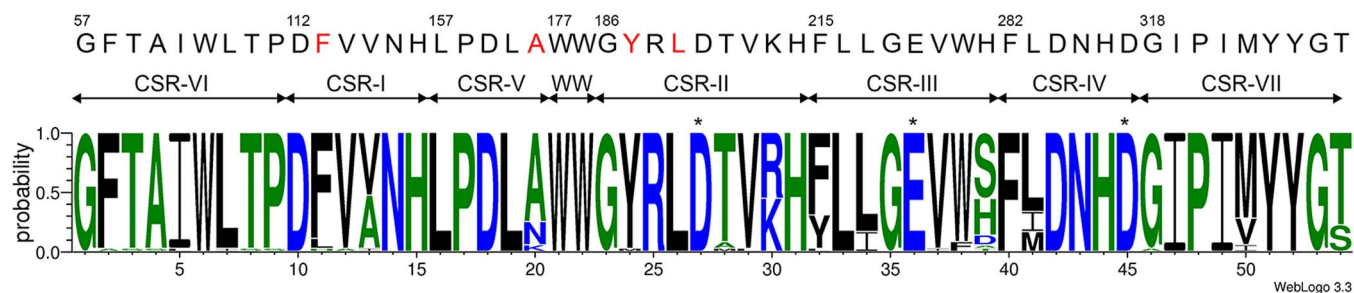


Figure 1 | Sequence logo for the seven CSRs of the GH13 subfamily represented by ASKA. CSR-I, residues 10–15; CSR-II, residues 23–31; CSR-III, residues 32–39; CSR-IV, residues 40–45; CSR-V, residues 16–20; CSR-VI, residues 1–9; and CSR-VII, residues 46–54. The two characteristic, consecutive tryptophans are also shown. The catalytic triad of the GH13 family, i.e., the catalytic nucleophile (No. 27, aspartic acid), the proton donor (No. 36, glutamic acid) and the transition-state stabiliser (No. 45, aspartic acid) are indicated with asterisks. The logo is based on the sequences of 36 proteins including both the characterised α -amylases and very similar hypothetical proteins (Table 1). The corresponding ASKA CSR sequences are stated above the logo and the mutated residues described in this study are shown in red.

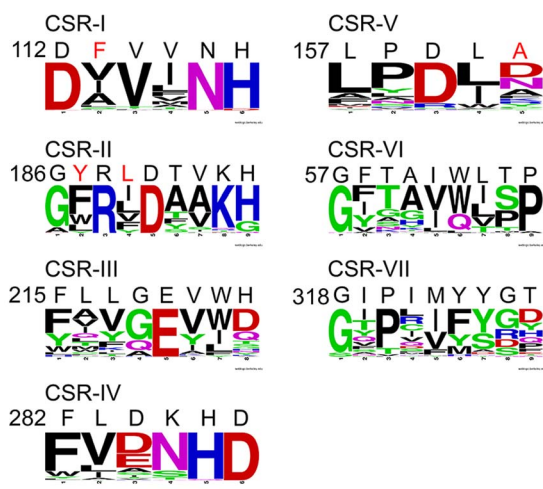


Figure 2 | Sequence logos of the seven CSRs based on the mixture of various GH13 subfamily α -amylases (Table S1). The size of the single letter amino acid code in each sequence logo represents the occurrence of a particular amino acid at a particular position. The corresponding amino acid sequences of ASKA were stated above each sequence logo and the mutated residues performed in this study were shown in red.

similar CSRs (Figure 1). A separate weblogo with α -amylase sequences from earlier known GH13 subfamilies was built and compare with ASKA's CSRs (Figure 2). ASKA clearly contains different residues in several positions of the CSRs (Figure 2). To investigate their functional role, Phe-113, Ala-161, Tyr-187, and Leu-189 of ASKA were thus selected and replaced with valine, aspartic acid, phenylalanine, and isoleucine, respectively. A total of four single-point mutant α -amylases were designed and denoted as F113V, A161D, Y187F, and L189I (Table 2).

Figure 3a illustrates the position of CSR-I to CSR-VII *in silico*. Phe-113 is the second residue of CSR-I (Figure 1; position 11 in the logo), which is located on strand β 3 of the catalytic (β/α)₈-barrel domain A²³. This position was chosen for the mutagenesis study because a valine or other aliphatic hydrophobic residue has frequently been observed at the corresponding position, e.g., in the α -amylases from subfamilies GH13_1 (fungi), GH13_5 (liquefying enzymes from bacteria), GH13_6 (plants) and GH13_7 (archaeons)^{23,24}. The next two mutant positions, Tyr-187 and Leu-189 (logo positions 24 and 26, respectively; Figure 1), are residues from CSR-II, which covers strand β 4 of the catalytic barrel²³. While the former position is well-conserved as an aromatic one throughout the entire α -amylase family GH13, it is mostly occupied by phenylalanine or tryptophan^{23,24} and the presence of the tyrosine (Figure 2) here in ASKA seems to be unique for the GH13 subfamily¹¹. Because tryptophan is highly characteristic of the GH13 subfamilies 6 and 7, which include plants, archaeons and flavobacteria^{23–25}, phenylalanine, which is found in the GH13 subfamilies 15 and 24 (animals) as well as 27, 28, 32 and 36 (various groups of bacteria), was chosen as the substitute for Tyr-187 in the mutant. The latter mutant position in CSR-II, Leu-189, is a conserved position, but if it is replaced then isoleucine (Figure 2) is

often found^{23,24}. The fourth residue selected for mutation, Ala-161, is positioned in CSR-V near the C terminal of domain B, i.e., in the loop connecting strand β 3 to helix α 3 of catalytic (β/α)₈-barrel domain A²³. Although various residues can be seen at that position, aspartic acid or asparagine may represent a consensus^{23,24}. It should therefore be of interest to investigate the role the alanine may play in ASKA (Figure 1; position 20 in the logo) because an alanine is typically present in the corresponding position only in the animal α -amylases (from vertebrates) from the subfamily GH13_24²³.

Characterisation of the mutant α -amylases. The purified enzymes were subjected to sodium dodecyl sulphate polyacrylamide gel electrophoresis (SDS-PAGE) to determine the purity of the proteins (Figure 4). The mutant enzymes were pure with single bands at the molecular mass of 50 kDa in agreement with that of native α -amylase¹⁸. The zymogram analysis was performed using native PAGE, and the gel was stained with iodine solution. For all the enzymes, a single, clear band was observed upon staining. The zymogram of the purified proteins shows that the mutant enzymes retained amylolytic activity (Figure 4a). After the purification procedures, the enzyme activity and the protein concentration for all α -amylase variants were determined, and the specific activity of each α -amylase variant was calculated (Table 2).

The effect of temperature on the activity of the ASKA and mutant variants was tested at temperatures ranging from 30 to 90°C at pH 8.0 (Supplemental file). The optimum reaction temperature for all the α -amylase variants was 60°C. None of the amino acid substitutions affected the optimum temperature activity. At 70°C, the enzymes maintained 54–87% of the initial activity. As the incubation temperature rises, the activity of the enzymes drops to 13–24% at 80°C and 10–18% at 90°C. Compared to ASKA, mutants Y187F and L189I had a slight decline in activity of approximately 32% and 23%, respectively at a temperature of 70°C. The thermostability of the enzymes was studied at the temperatures of 60°C and 65°C for 72 h. The activity half-life of all mutants at their optimum temperature of 60°C is unaffected by the mutation (Supplemental file). The mutant A161D retained 94.6% of its total activity after 72 h of incubation at 60°C, and this value is slightly higher compared to ASKA, which retained 91% of its activity. The activity half-life at 65°C is displayed in Table 2, the ASKA had a half-life of 48 h, while Y187F and L189I had a slight reduction in thermostability with half-lives of 43 and 36 h, respectively. The activity half-lives at 65°C for the mutants F113V and A161D were 51 and 53 h, respectively.

The effect of pH on α -amylase activity and stability were tested at a pH range of 4.0 to 11.0 using sodium acetate buffer (pH 4–5), sodium phosphate buffer (pH 6–7), Tris-HCl buffer (pH 8–9) and glycine-NaOH buffer (pH 10–11) at a temperature of 60°C. ASKA and all mutant α -amylases exhibited maximum activity at pH 8.0. The optimum pH is thus unaffected by the amino acid substitutions. This result is expected because very frequently the optimum pH of an enzyme is not easily affected by a single residue replacement. The ASKA and mutant α -amylases were stable within a wide pH range of 7.0 to 10.0 for 30 min, and the enzyme activities were at least 90% of the initial enzyme activity at these pH values. The pH stability of the mutants was similar to the ASKA. The pH stability of ASKA was unaffected by the mutations.

Table 2 | Specific activities and kinetic parameters of ASKA and mutant α -amylases

Protein	CSR	Domain, secondary structure	Half-life, $t_{1/2}$ at 65°C (h)	Specific activity (U/mg)	k_{cat} (s^{-1})	k_{cat}/K_m ($ml\ mg^{-1}\ s^{-1}$)
ASKA			48	34	17,984	14,790
F113V	I	A, β strand 3	51	33	24,211	7,271
A161D	V	B, loop connecting β stand 3 to α helix 3	53	76	39,108	57,463
Y187F	II	A, β strand 4	43	30	5,258	4,093
L189I	II	A, β strand 4	36	27	3,429	3,717

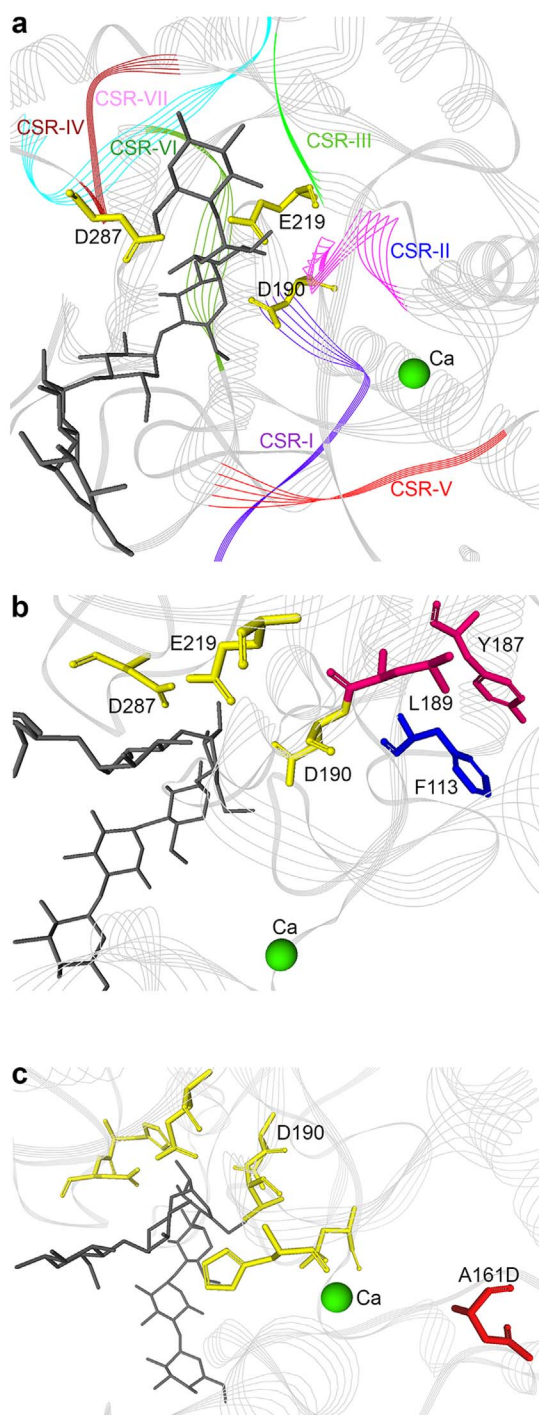


Figure 3 | (a). Structural prediction of native ASKA. Green sphere, Calcium; grey ligand, substrate; yellow, three catalytic residues D190, E219 and D287; blue violet, CSR-I; violet, CSR-II; apple green, CSR-III; maroon, CSR-IV; red, CSR-V; forest green, CSR-VII; blue, CSR-VII. (b). The mutation sites F113, Y187 and L189 are located adjacent to the catalytic residues. (c). Structure prediction for mutant A161D. The putative residues for subsites -1 and $+1$ are shown in yellow, and the side chain of mutant A161D is shown in red. The average distance from the A161D side chain to catalytic residue D190 is 18 Å.

The specific activities, turnover rate (k_{cat}) and catalytic efficiency (k_{cat}/K_m) of ASKA were compared to its mutant variants (Table 2). ASKA had a specific activity of 34 U/mg-protein. Of the mutant α -amylases, the mutant A161D had the highest specific activity at 76 U/mg with an increase of approximately 1.23 times. The V_{max}

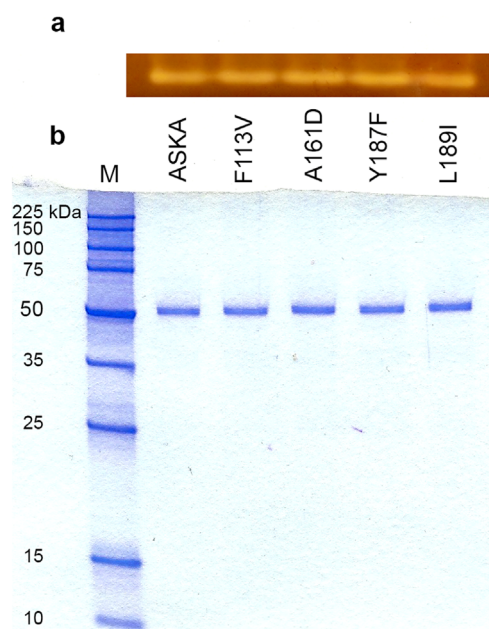


Figure 4 | (a). Zymogram analysis of the purified native and mutant α -amylases. (b). SDS-PAGE. Lane M: Promega™ Broad Range protein marker, Lane 1: Native ASKA, Lane 2: Mutant F113V, Lane 3: Mutant A161D, Lane 4: Mutant Y187F, Lane 5: Mutant L189I.

as well as the k_{cat}/K_m were drastically changed by the A161 mutation. The mutant A161D exhibited the highest k_{cat} of 39,108 s^{-1} , which is 1.17-times higher than the k_{cat} value of ASKA. Moreover, the k_{cat}/K_m of A161D was also higher than that of ASKA by 2.88-times, at 57,463 $\text{ml mg}^{-1} \text{s}^{-1}$. Of the four mutants constructed using site-directed mutagenesis, only mutant A161D exhibited enhanced k_{cat} and k_{cat}/K_m values (Table 2). The L189I substitution had an undesirable effect as the k_{cat} and k_{cat}/K_m values were reduced by 81% and 74%, respectively. Such deterioration was the worst observed of the mutants. The adjacent mutant Y187F also exhibited a reduced efficiency in the catalytic reaction. Both of these residues were located near the catalytic Asp-190 (Figure 3b), and their mutations have caused a detrimental effect to the catalytic property of the enzyme. These results are in agreement with the findings of earlier reports^{26,27}, which stated that any mutations near the catalytic residue will most likely influence the catalytic property of enzyme and likely cause an overall negative effect on enzyme catalysis.

All mutant enzymes were subjected to an enzymatic reaction to study the effect of the mutation on product specificity. Hydrolysis of 1% (w/v) and 4% (w/v) gelatinised soluble-, tapioca-, potato-, corn-, and sago-starch was carried out at 60°C for 24 h. The total sugar production (mg/ml) is shown in Table 3. At the lower starch concentration of 1%, the yield of total sugars generated by mutants Y187F and L189I was significantly lower than that of the ASKA; however, this effect was less significant at the higher concentration of starch (4%). With the 1% starch solution, the total sugars yielded by mutant A161D were quite similar to the ASKA most likely because the amount of substrate was not in excess. Interestingly, at a higher substrate concentration, for instance 4% corn-starch, the total sugar formed by mutant A161D was 27.8 mg/ml, which is higher than that of the ASKA (16.3 mg/ml) (Table 3). The k_{cat} and k_{cat}/K_m values for mutant A161D are better than for ASKA, and the HPLC analysis confirms the outperformance of A161D.

The percentages of individual sugars that were formed after 24-h reactions in 1% (Figure 5a) and 4% corn-starch (Figure 5b) were studied. Previously, we reported that ASKA is a maltose-forming α -amylase¹⁸, and a negligible percentage of glucose (G1), maltohex-



Table 3 | Total sugar production (mg/ml) from the hydrolysis of different starches by ASKA and mutant α -amylases using 1% (w/v) and 4% (w/v) of starches

Type of starches	1% (w/v) of starches					4% (w/v) of starches				
	ASKA	F113V	A161D	Y187F	L189I	ASKA	F113V	A161D	Y187F	L189I
Corn	12.5	12.3	13.3	10.9	11.1	16.3	17.0	27.8	15.0	15.7
Soluble	12.1	10.4	12.1	9.5	10.5	16.3	15.8	24.9	14.9	15.1
Tapioca	11.0	11.4	12.8	9.5	10.5	16.2	16.9	25.8	14.2	14.7
Potato	11.2	11.7	12.0	10.0	10.4	16.1	16.7	24.5	14.2	14.9
Sago	11.8	12.0	12.9	11.4	10.7	16.1	16.4	27.4	15.5	15.5

ase (G6) and maltoheptaose (G7) were usually detected in 1% gelatinised corn-starch (Figure 5a). However, when the ASKA was reacted with 4% corn-starch for 24 h, a small percentage of G6 and G7 were detected. The product specificity spectrum for mutant A161D changed significantly for the reaction that was carried out in 4% starch. The percentage of maltose forming dropped, while a substantial increase in the percentage of glucose was observed (Figure 5b).

Discussion

α -Amylases, in addition to all members of the α -amylase GH13 family, contain four to seven conserved sequence regions (CSRs), roughly covering strands β 2, β 3, β 4, β 5, β 7 and β 8 of the catalytic TIM-barrel and a short stretch near the C-terminus of domain B protruding out of the barrel in the β 3- α 3 loop. The individual α -amylases often possess unique sequence features in their respective CSRs that distinguish them from each other, and thus these regions may be connected to the specific biochemical properties of each protein, such as substrate preferences, product specificities, and thermostability^{9–11}.

Anoxybacillus sp. SK3-4 was isolated from a Malaysian hot spring²⁸ and its genome was sequenced using Illumina MiSeq²⁹. Most of the glycoside hydrolases of the cell are expressed intracellularly; however, ASKA was described as an extracellular hydrolase¹⁸. Compared with well-recognised α -amylases, the ASKA protein sequence has a low similarity of 49% to the *G. stearothermophilus*³⁰ (BSTA, subfamily GH13_5, UniProt Accession number P06279), 47% to the *Aspergillus oryzae*³¹ (TAKA, GH13_1, P0C1B3) and 37% similarity to the *B. licheniformis*³² (BLA, GH13_5, P06278) α -amylases. Despite the fact that ASKA is a high maltose producing α -amylase, it is a genuine α -amylase. In fact, the two types of family GH13 amylases forming mainly maltose (EC 3.2.1.133) should be distinguished from each other, i.e., maltogenic amylases and maltogenic α -amylases²³. Whereas the former enzymes are members of the so-called neopolulanase subfamily (subfamily GH13_20), the latter ones are closely related to cyclodextrin glucanotransferases (CGTase) (GH13_2). A typical GH13_20 maltogenic amylase possesses in its CSR-V the signature MPKLN⁹, contains at its N-terminus a carbohydrate-binding module from the family CBM34³³ and is able to hydrolyse cyclodextrins³⁴. Because the ASKA has a different CSR-V (157_LPDLA; Figure 1), lacks the CBM34 N-terminal domain and is not active on cyclodextrins¹⁸, it is likely not related to this type of maltogenic amylase. For the maltogenic α -amylases from the subfamily GH13_2, the CSR-V signature (LADLS) from *B. stearothermophilus*²³ is more similar to that seen in ASKA, but it typically shares its five-domain organisation with CGTase³⁵; i.e., it has two C-terminal domains in addition to domains A, B and C that are present in the ASKA. Thus, ASKA and most likely the entire newly established α -amylase family GH13 group¹¹ (i.e., ADTA, BaqA and homologues from *G. thermoleovorans*) may represent a novel type of high maltose producing amylase.

ASKA shares 69% similarity with the α -amylase from *G. thermoleovorans* subsp. *stromboliensis* Pizzo²², 69% to *G. thermoleovorans* GtamyII²⁰ and 60% to a marine *B. aquimaris* MKSC 6.2 α -amylase (BaqA)^{11,36}. Recently, the crystal structure of the α -amylase *G. thermoleovorans* (GTA) was reported²¹. The sequence similarity between ASKA and GTA is as high as 69%. ASKA, Pizzo, GtamyII, GTA, and BaqA as well as another *Anoxybacillus* α -amylase ADTA from our laboratory¹⁸ can be classified as a new sub-group of the α -amylase family GH13. Taken together, the reported experimental data of these α -amylases, this new subgroup has at least three unique characteristics. Firstly, they are high maltose producing enzymes as seen in ASKA, ADTA and Pizzo and potential in GTA since Pizzo shares an identical protein sequence with GTA²¹. Secondly, these enzymes are able to degrade raw starches as reported for the Pizzo, GtamyII, and BaqA amylases^{11,20,22}. Although the two consecutive tryptophans were proposed to be the putative raw starch binding residues¹¹, the binding action has yet to be experimentally proven. Thirdly, the C-terminal of the ASKA protein sequence was predicted to be a putative transmembrane region that could possibly anchor to the wild type *Anoxybacillus* sp. SK3-4¹⁹. Recent work on the deletion of this transmembrane region of GtamyII elucidated its role in raw starch adsorption²⁰.

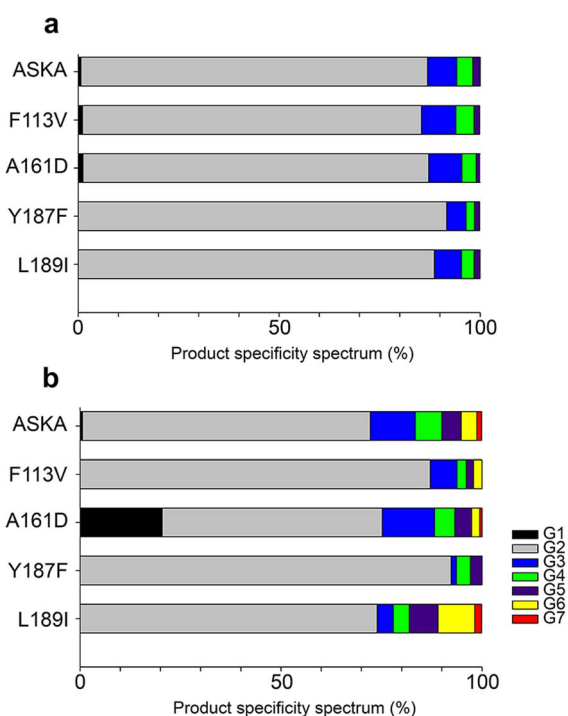


Figure 5 | End product distribution of ASKA and mutant α -amylases from the hydrolysis of a) 1% (w/v) and b) 4% (w/v) corn-starch. The data are shown as the percentage (%) hydrolysed. Black: glucose (G1), grey: maltose (G2), blue: maltotriose (G3), green: maltotetraose, (G4), purple: maltopentaose (G5), yellow: maltohexaose (G6) and red: maltoheptaose (G7).



The main goal of this current work is to address the reason that the enzymes from this new GH13 subgroup, for instance ASKA, are a high maltose forming. The CSRs of ASKA were compared to other reported α -amylases, and four single point mutations were carried out. Interestingly, the product specificity of mutant A161D was altered causing the overall ratio of maltose to drop, while the percentage of glucose increased 28-times in 4% corn-starch. Based on these findings, Ala-161 is proposed to be responsible for the high maltose-producing characteristic of ASKA. In the maltose producing ADTA, Pizzo and GTA α -amylases, Ala is well conserved at an identical position. The other homologs, i.e., GtamyII and BaqA also contained Ala, but their product specificity has yet to be determined.

Such drastic change was unexpected because the location of 161D in the predicted 3D structure is 18 Å from the nearest catalytic residue, Asp-190 (Figure 3c). Moreover, the superimposition of the mutant A161D model with the *G. thermoleovorans* α -amylase (4EZO) elucidated that 161D has no direct interaction with the pseudo substrate acarbose. At this stage, it is not clear the true reason why product specificity is associated with Ala-161. In CGTase, the product specificity is often related to the type of amino acids at the subsite interacting residues^{37–39}, yet A161 is not likely to be a part of any subsites. The *in silico* position of residue 161D is located close to the putative calcium (Ca^{2+})-binding site (Figure 3c). A161D may stabilise the secondary structure or the folding of this Ca^{2+} -binding pocket, therefore the half-life of this mutant was higher than that of ASKA (Table 2). Nevertheless, at this point, we are unable to determine why A161D can alter the product specificity in a drastic manner. We are now conducting further examination of this site by creating other mutations to examine the effects of various amino acid side chains on product specificity. Besides X-ray crystallisation will be performed in the near future. We hope that these experiments will lead to more insight to the importance of A161 in product specificity.

Several mutations have previously been made to the CSRs of other α -amylases. For instance, Chen *et al.* (2010) mutated Asp-236 of the *Bacillus* sp. strain TS-23 α -amylase⁴⁰. This mutated residue corresponds to Ala-161 in ASKA numbering, in CSR-V. The replacement of Asp-236 with asparagine resulted in a 33% reduction in the specific activity of the enzyme. Interestingly, this finding contradicts the A161D mutation results obtained in this study. In addition, Chen *et al.* (2010) mutated Asp-234, which is also located in CSR-V. This substitution resulted in an 86% reduction in the specific activity of the enzyme. It seemed from that study that two residues in CSR-V (Asp-234 and Asp-236) played a significant role in the catalytic properties of the enzyme⁴⁰. Unfortunately, both trials disrupted the specific activity, and the authors suggested that Asp-234 and Asp-236 were involved in Ca^{2+} -binding, and any changes to these Ca^{2+} -binding residues will affect the enzyme's structural integrity, and stability as well as its α -amylase activity. In another study, the substitution of Asp-233 of the *B. amyloliquefaciens* α -amylase (corresponds to ASKA's Ala-161) significantly affected the specific activity and thermostability of the enzyme. There was an 84.4% reduction in specific activity compared to the ASKA. From this result, the authors deduced that Asp-233 is important for the catalytic activity of the enzyme⁴¹.

Despite mutant F113V exhibiting a similar specific activity to ASKA, the k_{cat} value for the mutant was 35% higher. ASKA and its homologues ADTA, Pizzo, GtamyII, BaqA, and GTA have a Phe at this particular position of CSR-I, while most other α -amylases have another type of amino acid (Figure 2). Several mutations of CSR-I residues were performed previously, for instance, H137L (His-117 in ASKA) of the *Bacillus* sp. TS-23 α -amylase resulted in a complete loss of amylolytic activity⁴². Chang *et al.* (2003) explained that His-137 could be the essential catalytic residue in the amylolytic reaction. The site directed mutagenesis of another CSR-I residue, Asn-104 of BLA (corresponds to ASKA's Asn-116) was replaced with Asp. This mutant had an 11% decrease in specific activity at 30°C. But, inter-

estingly, the mutant had a 30% higher specific activity than the ASKA at the higher temperature of 70°C⁴³.

This study found that the mutation of CSR-II, i.e., mutant Y187F and L189I caused a more than 70% reduction in the k_{cat} and k_{cat}/K_m (Table 2). For mutant Y187F, the chemical structures of tyrosine and phenylalanine are structurally close except that phenylalanine lacks a hydroxyl -OH group at end of the aromatic side chain. The removal of the hydroxyl group caused a severe disturbance during the starch hydrolysis process. The hydroxyl group is likely involved in stabilising of the intermediate state during the catalytic mechanism. These results elucidate that the amino acid residues Tyr-187 and Leu-189 are associated with ASKA activity, most likely because they are located adjacent to catalytic residue 190 (Figure 3b). Similarly, the mutation of the *Bacillus* sp. TS-23 α -amylase, in CSR-II at His-269 (TS-23 numbering), which is adjacent to catalytic Asp-265, led to the complete loss of amylolytic activity. The report proposed that this conserved histidine may play a substrate binding role³⁷, or possibly, this stretch of CSR-II is sensitive to alteration.

Methods

Bioinformatics analysis. The 36 amino acid sequences (Table 1) that are currently thought to form the GH13 subfamily represented by ASKA were retrieved from GenBank database⁴⁴. The sequences were collected from a BLAST search⁴⁵ using as the query the stretch of 89 residues from the ASKA sequence¹⁸ spanning the polypeptide chain covering CSR-I (strand β 3) and CSR-II (strand β 4), including the entire domain B (GenBank Acc. No.: AFI49455.1: Asp-129-His-217). A sequence was considered to belong to this GH13 subfamily if it contains the features ascribed to the closely related α -amylase from *B. aquimaris*¹¹, especially the two consecutive tryptophan residues in helix α 3 of the catalytic TIM-barrel and the specific sequence in CSR-V positioned in domain B.

The sequences were aligned using the program Clustal-Omega⁴⁶ available on the European Bioinformatics Institute's server (<http://www.ebi.ac.uk/>). For the whole set of 36 sequences, a sequence logo for their seven CSRs typical of the members of the α -amylase family GH13²³ was created using the WebLogo 3.0 server⁴⁷ (<http://weblogo.berkeley.edu/>).

Preparation of starting structure and molecular dynamics simulation. The protein structure of ASKA was constructed using the SWISS-MODEL⁴⁸ program (<http://swissmodel.expasy.org/>), and the X-ray crystal structure of *G. thermoleovorans* α -amylase (PDB code: 4EZO)²¹ was selected as the template. Refinements of the model were performed using energy minimisation and dynamic simulation (MD). Prior to the MD simulation using the GROMACS software package⁴⁹, calcium and glucose were removed from the structure. The system was solvated with SPC/E water in a dodecahedral box with 10 Å between the protein structure and the box. The system was subsequently neutralised by the addition of sodium ions. The MD simulation was performed using an OPLS-AA force field. The system was energy minimised and subsequently simulated in NVT and NPT ensembles. The temperature was maintained at 333 K. Upon completing the system equilibration, a production run of 10 ns was performed.

Mutagenesis. The α -amylase mutant gene was generated using the Overlap Extension PCR (OE-PCR) method. The external forward primer contained an *EcoRI* restriction site, and the external reverse primer contained an *XhoI* restriction site. The change in nucleotide sequence was introduced by incorporating nucleotide changes into the overlapping oligo primers, as shown in Supplemental file. During the cloning process, the native signal peptide DNA sequence was excluded, as the pET-22b(+) contains a pelB leading sequence. Phusion[®] PCR Master Mix (NEB, Ipswich, Massachusetts, USA) was used to perform the PCR. PCR reaction mixtures containing 25 ng ASKA DNA template, 25 μ l 2 \times Phusion[™] Flash PCR Master Mix, and 0.5 μ M each primer were added to a final volume of 50 μ l. A SpinPrep[™] PCR Clean-Up kit (Novagen, Darmstadt, Germany) was used to purify the PCR products. Purified PCR products and pET-22b(+) (Novagen, Darmstadt, Germany) were digested with *EcoRI* and *XhoI* (NEB, Ipswich, Massachusetts, USA) and subsequently purified. The ligation was performed using T4 DNA ligase (NEB, Ipswich, Massachusetts, USA) and then transformed into *E. coli* DH5a. The recombinant plasmid was confirmed by sequencing. The recombinant plasmid was then transformed into the expression host *E. coli* BL21 (DE3) for later analysis.

Protein expression and purification. The recombinant *E. coli* BL21 (DE3) was cultured overnight in LB medium supplemented with 100 μ g/ml ampicillin at 37°C with agitation at 200 rpm. Induction was performed by adding 0.01 mM IPTG, and the cultivation was continued at 25°C with agitation at 200 rpm for 48 h⁵⁰. The cell free enzyme was harvested after centrifugation (10000 \times g, 20 min, and 4°C). The crude enzyme solution was concentrated using the Vivaflow 50 crossflow system from Sartorius Stedim Biotech (Göttingen, Germany) using the polyethersulfone (PES) membrane with a 10 kDa molecular weight cut-off (MWCO). The retentate was loaded onto affinity columns, and chromatography was conducted using the



ÅKTAprime plus system (GE Healthcare, Uppsala, Sweden) according to the method reported previously¹⁸.

Characterisation of ASKA and mutant enzymes. The protocols used to characterise the mutant α -amylases were also performed using ASKA unless otherwise stated¹⁸. One unit of α -amylase was defined as the amount of enzyme that releases 1 μ mol maltose per min per ml under the assay conditions specified. The kinetic parameters of the ASKA and mutant α -amylases were determined by measuring the amount of maltose production using the standard assay conditions. The samples were reacted with soluble starch (Kanto Chemical, Tokyo, Japan) at concentrations from 0.1 to 2.0 mg/ml in 0.1 M Tris-HCl buffer (pH 8.0). SigmaPlot v11.0 was used to build the Hanes plot.

The products of starch hydrolysis by α -amylase were analysed using High-performance liquid chromatography (HPLC). First, 0.1 ml purified enzyme was incubated with 1.0 ml 1% gelatinised starch in 0.1 M Tris-HCl buffer (pH 8.0) at 60°C for 24 h. The efficiency of degradation on various starches (soluble-, tapioca-, potato-, sago- and corn-starch) was compared. After 24 h of incubation, the reaction mixture was boiled for 15 min to stop the reaction and cooled to room temperature. The mixture was filtered through a 0.45 μ m nylon syringe filter (Whatman, Little Chalfont, United Kingdom) to remove insoluble particles. The hydrolysis products were analysed using a Waters HPLC system with a Zorbax Carbohydrate analysis column, 4.6 \times 250 mm (Agilent, Santa Clara, California, USA). An acetonitrile (HPLC grade, Merck, White House Station, New Jersey, USA) to water mixture (70 : 30, v/v) was used as the mobile phase, and the flow rate was maintained at 1.0 ml/min. The column was kept at 30°C, and a Waters 2414 refractive index detector was used to detect the reaction products. The data were recorded by the integrated computer system attached to the HPLC. Glucose, maltose, maltotriose, maltotetraose, maltopentaose, maltohexaose and maltoheptaose (Sigma-Aldrich, St. Louis, Missouri, USA) were used as standards. Additionally, the efficiency of starch concentration on the hydrolysis products was investigated by incubating the ASKA and mutant enzymes in a 4% gelatinised starch in 0.1 M Tris-HCl buffer (pH 8.0) at 60°C for 24 h. The end products were analysed using HPLC. The assays and data collection were performed at least in triplicate, unless otherwise specified.

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Author contributions

S.J., S.S., R.N.Z.R.A.R., K.G.C. and K.M.G. conceived and designed the experiments. V.R., K.P.C. and K.G.C. performed the experiments. V.R., S.J., S.S. and K.M.G. analyzed the data. S.J., R.N.Z.R.A.R., K.G.C. and K.M.G. contributed reagents/materials/analysis tools. V.R., S.J., K.P.C. and K.M.G. wrote the paper. V.R., S.J., K.P.C. and K.M.G. prepared figures 1–5. All authors reviewed the manuscript.

Additional information

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