

Construction of efficient multifunctional enzymes for Bioethanol production: Various approaches and Futuristic Developments

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Abstract:

Increase consumption of non-renewable sources i.e. the fossil fuels has caused worldwide global warming and pollution. Bioethanol is one kind of biofuel produced from biological raw materials and lignocellulosic biomass. Bioethanol production can decrease the burden on non-renewable resources for energy. The chief constituents of the plant based lignocellulosic biomass are cellulose, hemicellulose and lignin. Cellulase and hemicellulases are responsible for hydrolysis of cellulose and hemicellulose components into monomeric sugars. The monomeric i.e. glucose and xylose can be fermented by using yeast for production of bioethanol. Moreover, the cellulases and hemicellulases are also used extensively for other industrial purposes such as biobleaching, food industry etc. However in spite of their larger industrial application, these present enzymes have low catalytic efficiency and high production cost of multiple enzymes, Due to these noteworthy efforts have been initiated for improving the performance and production rate of the present enzymes. Therefore, present chapter provides an overview of current scenario and

development in the field of plant cell wall degrading enzymes and their application. The chapter discusses different approaches of protein engineering tools to study the underlying mechanism of action of these enzymes and the current techniques utilized for refining their activity. The chapter also discusses recent development in production of efficient multifunctional chimeras for degradation of complex polysaccharide in a single reaction. Furthermore, the chapter also highlights the synthetic biology approaches that have been used for designing artificial designer cellulosomes containing distinct cellulases and xylanases domains which showed enhanced synergistic action against plant based biomass. Finally, the chapter covers various challenges and futuristic applications of these engineered enzymes in the field of renewable energy production.

Keywords (4-5): Lignocellulose biomass, bioethanol, protein engineering, designer cellulosome

1.1.0 Introduction

The extensive use of fossil fuels is the leading cause of global warming and pollution worldwide. The use of renewable energy sources is one of the solutions to decrease the increasing harmful effects.

Nowadays, the main focus of mankind is to make activities and life on earth more at ease by addition of essential products generated through biological processes. In the current scenario, each year millions of tons of plant based raw materials or lignocellulosic biomass are unused by burning them without providing benefits to living organisms. Therefore, it is highly important that these raw materials and lignocellulosic biomass should be efficiently utilized for generating renewable energy source in the form of biofuel by developing advanced scientific strategies. The lignocellulosic biomass are comprised of complex polysaccharides mostly cellulose, hemicelluloses and lignin. These complex polysaccharides are broken down by different sets of

enzymes called cellulase, hemicellulase and laccase in simpler monomers. The monomers mostly glucose and xylose are utilized by yeast in fermentation process to produce alcohol known as bioethanol. The bioethanol is the alternative form of energy that can be used as energy source instead fossil fuels. The enzymatic process involved in depolymerization of complex polysaccharides is one of the critical step in the production of bioethanol. Moreover, the low catalytic efficiency and production multiple enzymes involved in complete degradation of carbohydrate polymer is [one of the chief concern among scientific community](#).

Nowadays, major advancement in enzyme technology has resulted in development of efficient enzymes having high catalytic efficiency and multifunctionality. These efficient engineered enzymes are utilized for complete degradation of carbohydrate polymer into monomers.

1.1.1 Lignocellulosic biomass and its components

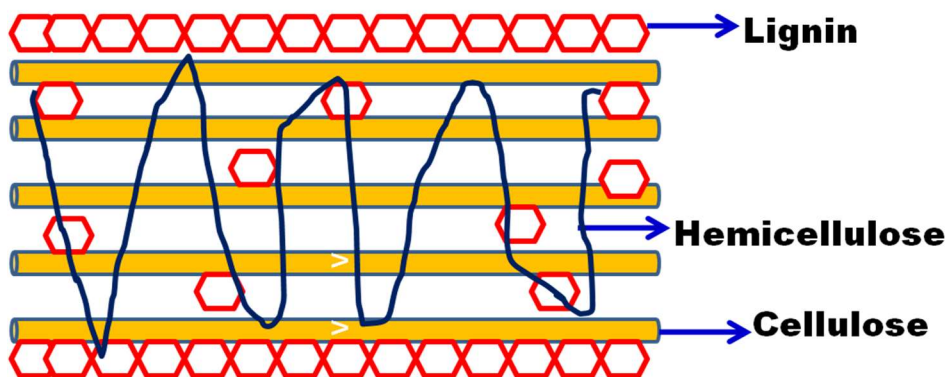


Figure: 1 Diagrammatic representations of components of the plant based lignocellulosic biomass

Plant based lignocellulosic biomass are cheap and readily available. From the biofuel production view lignocellulosic biomass is of particular interest. The major [polymeric constituents of lignocellulosic](#) biomass are cellulose, hemicelluloses and lignin as shown in Figure. 1 [1]. Generally, cellulose fibrils are encrusted with hemicelluloses

forming an open system, the remaining open spaces are filled up with lignin (Figure. 1). The different components of lignocelluloses biomass are discussed below.

1.1.1.1 Cellulose

Cellulose is the chief component plant based biomass. The cellulose content in most plant based biomass is on an average 33% [2]. The cellulose is extracted from biomass materials for biofuel production [2]. Cellulose is an organic polymer consisting of linear chain of glucose moieties connected together by β (1 \rightarrow 4) linkage as shown in Figure 2 [3].

The microfibrils of the cellulose are insoluble and are arranged into a cable-like structures (Figure 1) which comprised of around 24 hydrogen-bonded chains comprising β -(1,4)-linked glucose moieties as shown in Figure 2 (Guerriero *et al.*, 2010, Fernandes *et al.*, 2011, Nso *et al.*, 2017). The glucose polysaccharides chain is made up of glucose molecules forms parallel and successive chain of glucose moieties. The cellobiose is a disaccharide of glucose and is the repeating component of cellulose in which there is a 180° rotation of each glucose molecule with respect to its neighbouring glucose moiety as shown in Figure 3 [3]. The deconstruction of cellobiose is an estimation of β -glucosidase activity. The above arrangement described is responsible for a flat, inflexible and ribbon-like crystalline structure of glucan chains and is merged together by Van der Waals forces and hydrogen bonds to form microfibrils. Hydroxyl groups present in cellulose macromolecules are connected via enormous number of intra- and intermolecular hydrogen bonds, which caused in several ordered crystalline arrangements of cellulose chain into flat ribbon like conformations [4].

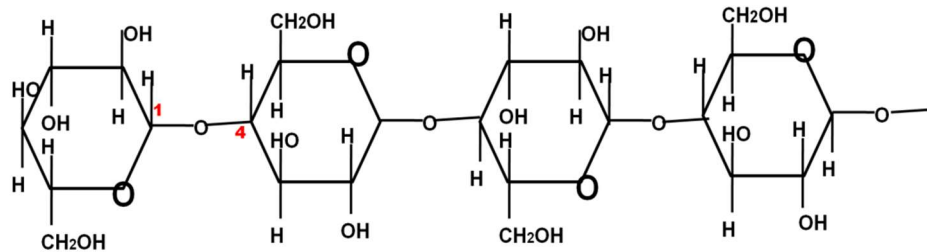


Figure 2: Demonstration of Cellulose chain in plant based lignocellulosic biomass

1.1.1.2 Hemicellulose

Hemicelluloses is the second most abundant polysaccharide representing 25-30% plant based biomass [5]. The hemicelluloses are made up of pentoses (xylose, arabinose), hexoses (mannose, glucose, galactose), and sugar acids [5]. The xylan is the most abundant hemicelluloses known. The hemicelluloses are not homogenous like cellulose, mostly they are heterogenous for example the hardwood hemicelluloses contains xylan however the softwood contains glucomannans [6]. The heteropolysaccharides xylan backbone is made up of homopolymeric chain of xylopyranose units associated via β -1,4 linkages as shown in Figure 3. Based on the xylan source, the backbone chain of xylan constitutes branches of arabinose, glucuronic acid or its 4-O-methyl ether, and ferulic acetic and p-coumaric acids (Figure 3).

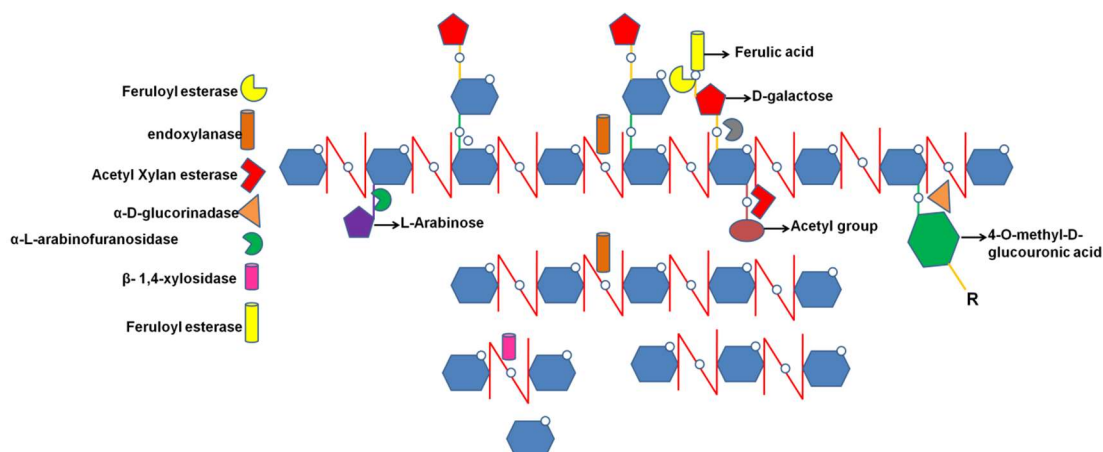


Figure 3. Xylan structure with different substituents and sites of attack by xylanases [6]

1.1.1.3 Lignin

Lignin is the main non-carbohydrate polymer present in plant based biomass. It is constitute (15-40%) of lignocellulosic biomass [7]. Lignin is accountable for protecting the plant cell wall from degradation by action of microbial community. It provides mechanical strength and hydrophobicity to the plant cell wall. The lignin is made up of from polymerization of different aromatic groups Guaiacyl (G), Syringyl (S), and hydroxyphenyl (H) to form monolignols [7]. The lignin is responsible for recalcitrance in plant based biomass. The cellulose accessibility is disrupted by higher content of lignin in lignocellulosic biomass thus reducing the availability of sugar residues for enzymatic saccharification reaction [7]. The lignin is connected to the carbohydrate polymer i.e. cellulose and hemicelluloses to form lignin carbohydrate complex or LCC through different linkages known as lignin carbohydrate complex linkages or LCC linkages [7] for example as shown in Figure. 4 lignin and carbohydrate is connected by an ester linkage.

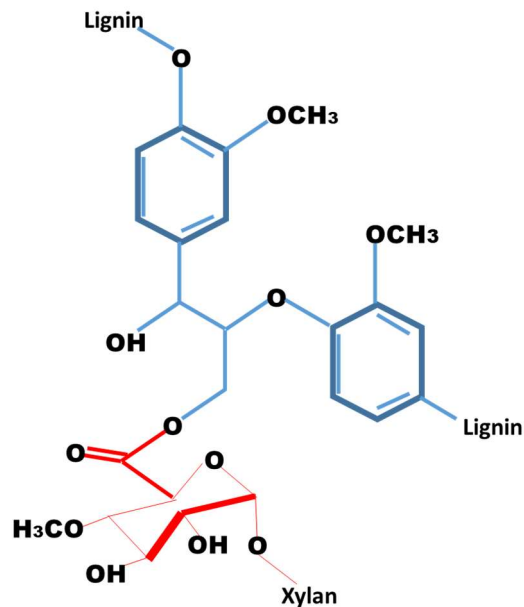


Figure 4 Ester Lignin-Carbohydrate Complex [25]

1.1.2 Enzymes for Lignocellulosic Hydrolysis

The plant based lignocellulosic biomass is recalcitrant in nature. Several factors are involved for biomass recalcitrant for example, high content of lignin, protection of cellulose by lignin, cellulose sheathing by hemicelluloses and high crystallinity and degree of polymerization [8]. The accessibility of plant carbohydrates for enzymatic saccharification is prevented by biomass recalcitrance. However, various pretreatment strategies showed improved accessibility of plant carbohydrates thus increasing the enzymatic hydrolysis step for releasing total reducing sugars for further bioethanol production [8]. Moreover, the complete degradation of the plant carbohydrates mostly cellulose and hemicelluloses are achieved by utilizing multiple number of enzymes discussed in the subsequent sections [9].

1.1.2.1 Cellulases

Cellulose upon depolymerization by the combined action of endoglucanase, cellobiohydrolase and β -glucosidase gives the product glucose [9]. Cellulase enzymes belong to different glycoside hydrolase families (GH) well described in CAZy database (www.cazy.org) [9]. Cellulase enzymes mostly constitute of two different modules namely, the catalytic module, which contains the active site and the non-catalytic carbohydrate binding module (CBM) as shown Figure 5. The two modules are connected by flexible linkers (Figure 5), comprising mostly of serine and threonine residues [11].

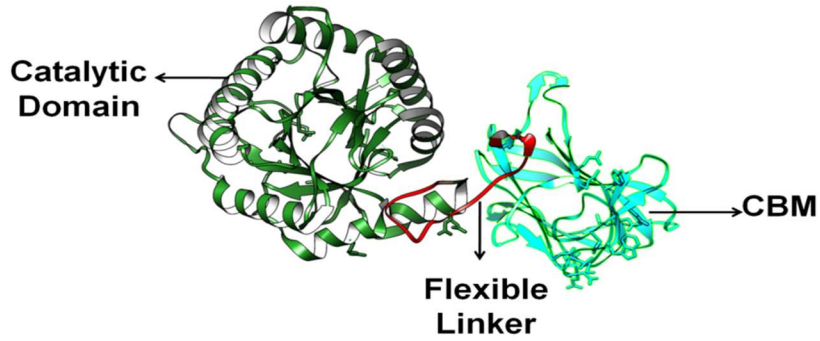


Figure: 6 Structure overview of a cellulase enzyme [11]

Moreover, cellulolytic enzymes act synergistically, in which endo- β -1,4-glucanase randomly acts on the cellulose chain and generates cellodextrins which are the larger cellooligosaccharides as a hydrolyzed products [12] (Figure 6). Cellobiohydrolase acts at the terminal of the cellodextrin chain and discharges cellobiose as the main product [12] (Figure 6). Moreover, ultimately the β -glucosidase acts on the cellobiose to release two molecules of glucose [12] (Figure 6). The following sections demonstrate different cellulases and their mode of actions.

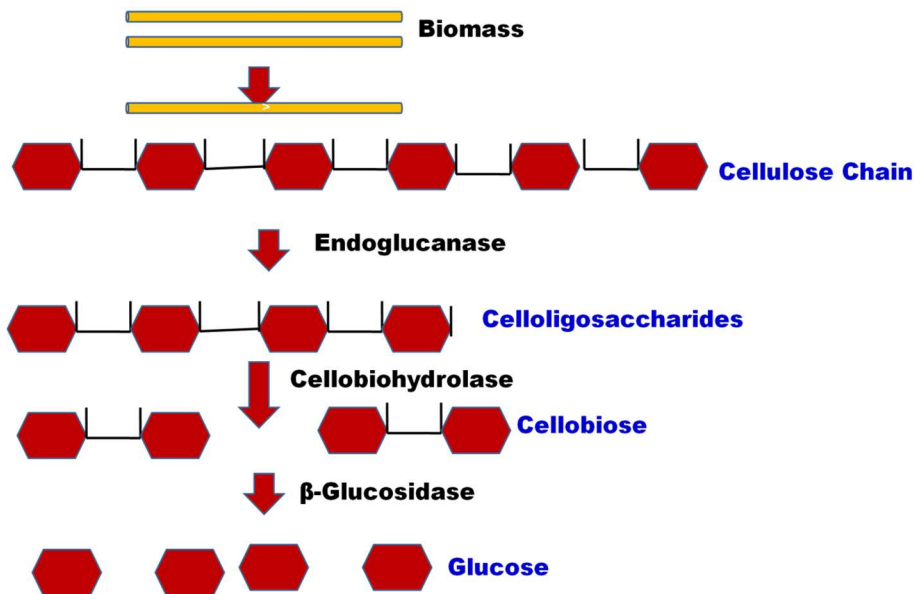


Figure 7: Demonstration of the action different cellulases in depolymerization of cellulose chain

[12]

1.1.2.2 Hemicellulases

Hemicellulases play an important role in the degradation of plant-based biomass. Hemicellulases depolymerize xylans, xyloglucans, arabinoxylans, and glucomannans of plant-based biomass [5]. Hemicellulases are modular structures with catalytic and functional domains as shown in Figure 7. In a previous study on extracellular β -L-arabinofuranosidase (HypBA2) which belongs to glycoside hydrolase (GH) family 121 from *Bifidobacterium longum* [13] was found to be a multidomain protein consisting of mostly N-terminal domain (amino acid residues 38-123, Brown), a N-domain from residue 124-303, Cyan followed by linker region from residue 304-347, yellow, a catalytic region comprising of $(\alpha/\alpha)_6$ barrel domain (348-771, green) followed by a C-terminal domain (772-870, blue).

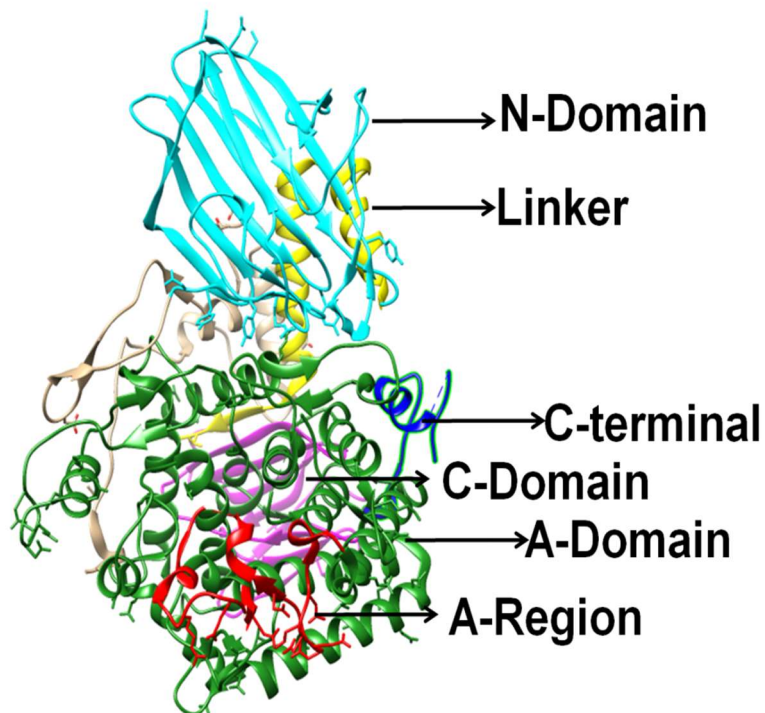


Figure 8: Multimodular structure of extracellular β -L-arabinofuranosidase (HypBA2) from glycoside hydrolase (GH) family 121 from *Bifidobacterium longum* [13]

The complete breakdown of hemicellulosic matter in plant biomass requires synergistic action of multiple enzyme due to its branched and organized structure as shown in Figure. 3

1.1.2.3 Lytic polysaccharide monoxygenases (LPMOs)

Lytic polysaccharide monoxygenases (LPMOs) are the mono-copper based enzymes which are broadly dispersed in nature that are responsible for catalysis of the glycosidic bonds hydroxylation in polysaccharide through oxidative reaction [14]. The LPMOs are auxillary enzymes and are distributed in CaZy database belonging to families AA9, AA10, AA11, AA13, AA14 and AA15 as demonstrated in Figure 9. In family AA10 the bacterial LPMOs are found which are responsible for cleaving cellulose and Chitin [15].

Protein Name	EC#	Reference	Organism	GenBank	Uniprot	PDB/3D
lytic chitin monoxygenase (AsLPMO10A; VSAL_I10134)	1.14.99.53	pubmed	<i>Allivibrio salmonicida</i>	CA080888.1	B5EQ86	
lytic chitin monoxygenase B (AsLPMO10B; VSAL_I10217)	1.14.99.53	pubmed	<i>Allivibrio salmonicida</i>	CA080921.1	B5EQ26	Z0KR[A]
lytic chitin monoxygenase (ChbB; BaAA10A; BaCBM33; Rbam17540; BAMF_1859)	1.14.99.53		<i>Bacillus amyloliquefaciens</i>	CB142885.1	E1UJX3	ZYQW[A,B] ZIDJ[A,B] ZYDT[A,B] SILJ[A,B]
lytic chitin monoxygenase (C1-specific) / chitin binding protein (ChbB; Cbp; B6CBP; BAA10A; BL00521; BL00145)	1.14.99.53	pubmed	<i>Bacillus licheniformis</i>	AAU39477.1 DSM 13 = ATCC 14580	DOEW57 O6Z0N7 O65N87	5TWE[A]
lytic chitin monoxygenase (Cbp; BRLPMO10A-FL; BRCBP)	1.14.99.53	pubmed	<i>Bacillus thuringiensis</i>	serovar <i>kurstaki</i> ACW83015.1	DOEW65	
chitin-active lytic monoxygenase (Lpmo10A; BRLPMO10A)	1.14.99.53		<i>Bacillus thuringiensis</i>	serovar <i>kurstaki</i> ACCC 10066	AJPM62637.1	ADAPCSK362 ADAP66FRT6
lytic cellulose monoxygenase (C1/C4-hydroxylating) (C1LPMO10; Cell_0270)	1.14.99.54 1.14.99.56	pubmed pubmed	<i>Cellulomonas fimi</i>	ATCC AEE44415.1	F4H6A3	484

Figure 9: Demonstration of LPMOs are auxillary enzymes and are distributed in CaZy database (<http://www.cazy.org/>)

The LPMOs produces holes on the crystalline surface of cellulose through oxidation reaction as shown in Figure 9. Thus LPMOs produce varied degree of cello-oligosaccharides [15]. Hence, the released cello-oligosaccharides are acted more efficiently by cellulases. The LPMOs acts synnergistically with cellulases for efficient breakdown of cellulose polymer in enzymatic hydrolysis reaction. Thus, nowadays the LPMOs act as a potent candidate in plant based biomass degrading enzyme cocktail [16].

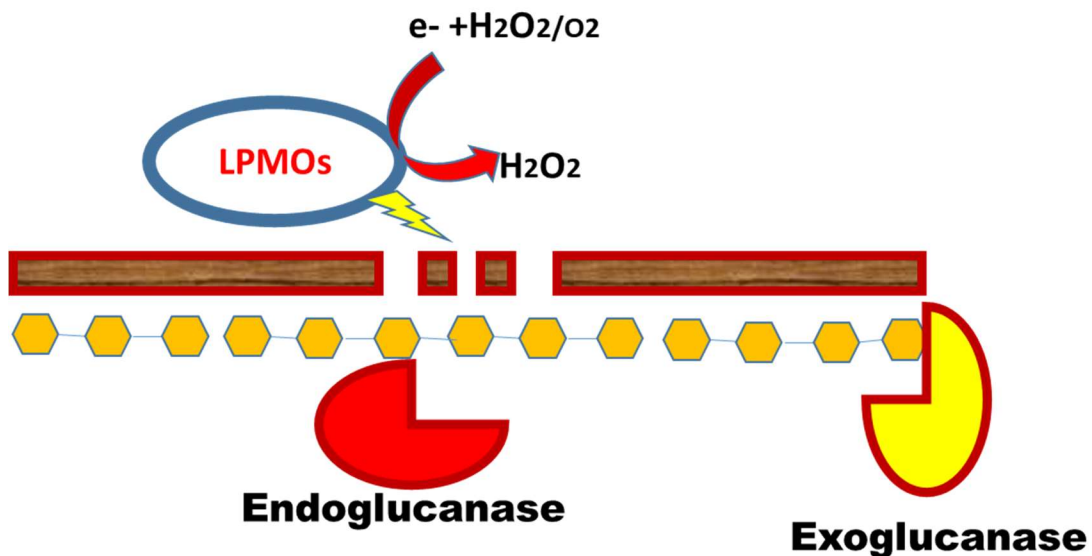


Figure 9: Diagrammatic representations of LPMOs and glycosidase in synergistic degradation of carbohydrate polymer [16]

1.2.0 Protein engineering existing plant cell wall degrading enzymes

1.2.0.1 Strategies for protein engineering

Cell wall degrading enzymes are widely used in enormous number of industrial purposes and therefore significant amount of efforts have been put for improving their performances and rate of production. Nowadays, protein engineering has been utilized to

study the underlying mechanism of action of these enzymes, as well as efforts have been made for improving their activity. Protein engineering comprises the mutagenesis of potential binding site residues and their kinetic analysis. Moreover, inactive mutants are mostly utilized for studying the protein-ligand complexes at their three-dimensional level. The protein engineering can be achieved by three strategies mainly, the directed evolution, the rational designing and by construction of chimeras having multiple functionality. In the directed evolution strategy of protein engineering random mutations of the target genes were executed and the variant with the enhanced activity were selected [2] as shown in **Figure.10**. In rational designing, the three-dimensional complexes of enzymes have been used to design new strategies for modification and exploitation of the glycoside hydrolases for engineering the enzymes and modifying their functions [17] as shown in **Figure.10**. The chimeras with multifunctional activity were constructed by utilizing molecular biology techniques. In multifunctional chimeras two or more modules were fused in a single polypeptide chains [18] as shown in **Figure.10**. The multifunctional chimera constructed can result in reduction the cost of production of several which are required for complete degradation of plant based lignocellulosic biomass [19]

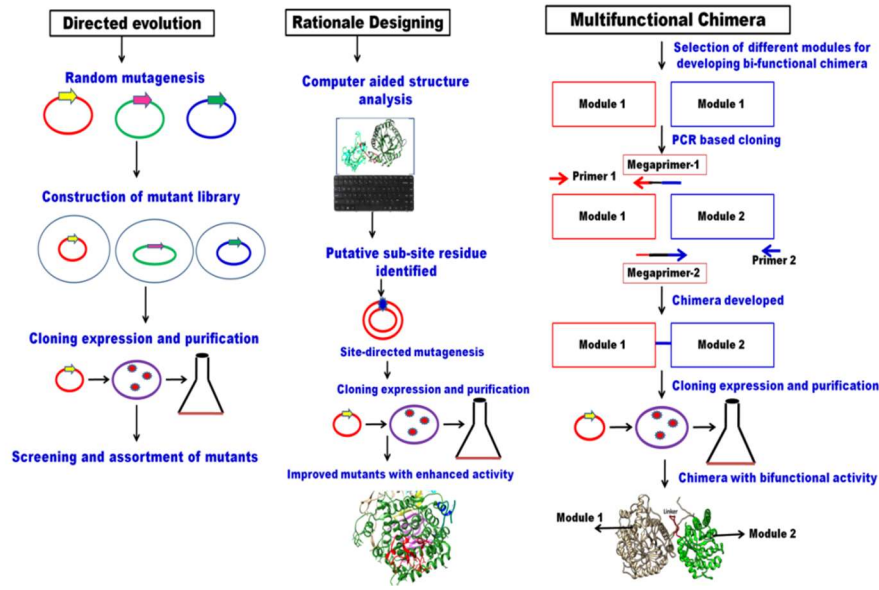


Figure 10: Diagrammatic representation of protein engineering strategies

1.2.0.1 Engineered multi enzyme complexes

Multi enzyme complex or cellulosome are large complex entity and are mostly found in anaerobic bacteria [20]. The multi enzyme complex or cellulosome is made up different carbohydrate degrading enzymes such as cellulase and xylanase [20]. The different domains in the cellulosome are attached by dockerin domain to a cohesion domain of main scaffoldin protein as shown in **Figure 11**. This complex structure is connected to the surface of microorganism. The cellulosome facilitates the microorganism to degrade the insoluble form of cellulose to soluble form so that the later could be absorbed by bacterial cell wall.

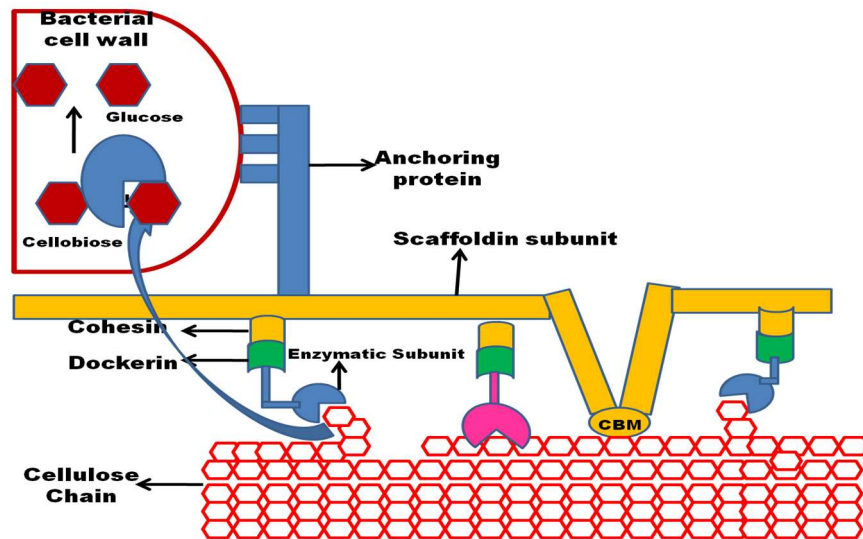


Figure 11: Structure and organization of cellulosome [20]

1.2.0.1.1 Construction of designer cellulosome

Designer cellulosome or engineered multi enzyme complexes are developed by attaching carbohydrate active enzyme to a scaffoldin discussed in the previous section with the help of strong cohesion and dockerin interaction [21]. Designer cellulosomes have efficient hydrolytic activity due to improved enzyme substrate closeness [21]. The designer cellulosome contains various domains which includes cellulases, hemicellulases, LPMOs and laccases [22]. The designer cellulosome are constructed by meticulous incorporation of desired catalytic activities based on the target substrate. The designer cellulosome can be constructed using various steps discussed below in the flow chart (Figure 12) [26]. The first step includes selection of domains and designing of designer cellulosome enzymes and scaffoldin protein. The second step consist of cloning of the cellulosome components by using their coding sequences. In the next steps the different components are assembled in the final multi enzyme complex. Lastly, in the final step the catalytic activity was determined for new assembled designer cellulosome.

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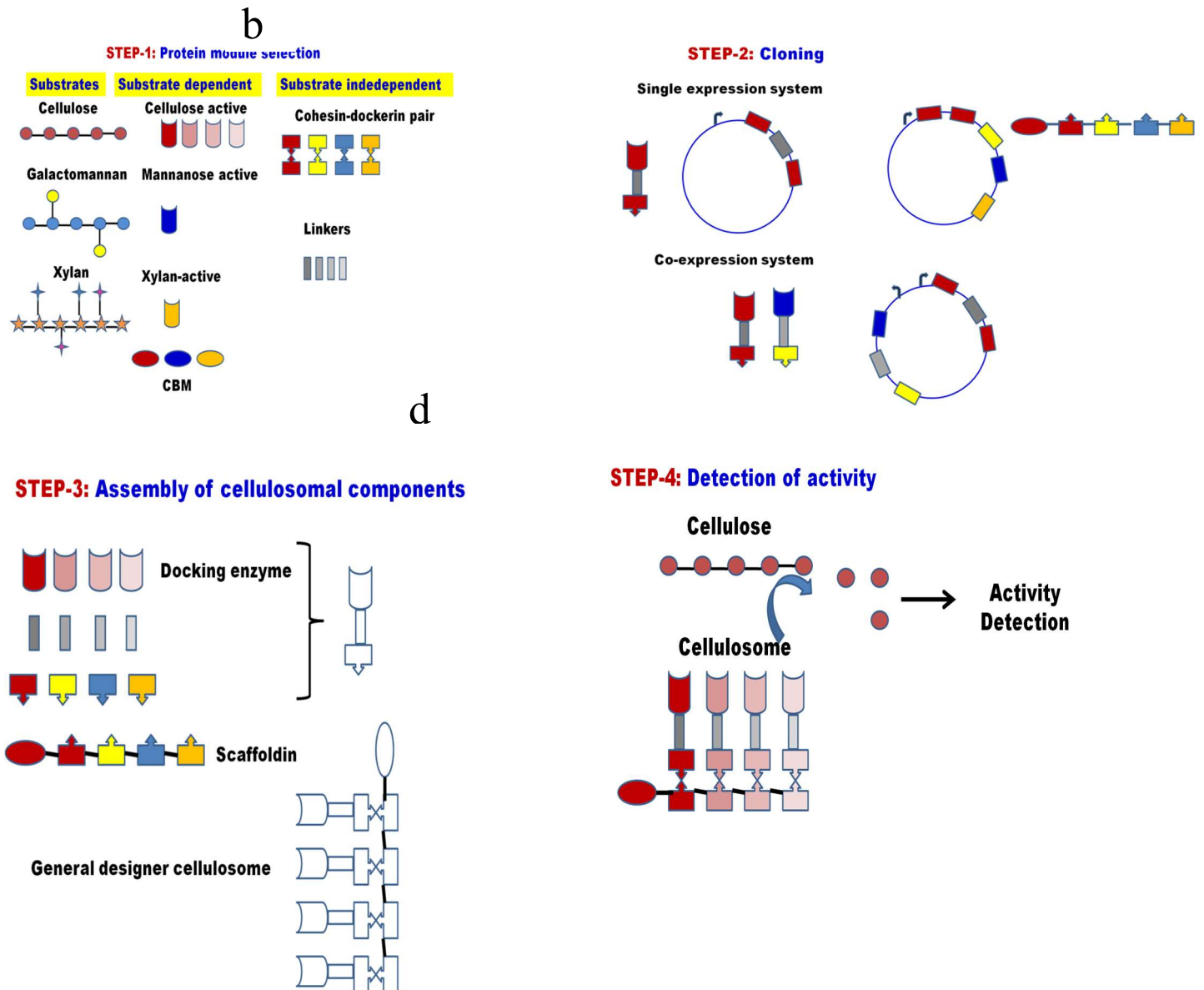


Figure: 12 Flow chart elaborating the steps for design, construction and evaluation of designer cellulosome [26].

1.2.0.1.2 Applications of designer cellulosome

1.2.0.1.2.1 Lytic polysaccharide monooxygenases in designer cellulosome

The synthetic biology approaches are used for designing efficient cellulosomal complex. The LPMOs are responsible for boosting cellulose degradation. Moreover, in anaerobic bacteria the cellulase enzymes are assembled into large

complex called cellulosome. Additionally, LPMOs are mostly known to be found in aerobic bacteria thus, it cannot be advantageous to the cellulosomal complex during cellulose degradation. In one of the study the LPMOs are incorporated into the cellulosomal complex through synthetic biology approach. In the study chimeric enzymes were developed by fusing LPMOs from the bacterium *Thermobifidafusca* using different dockerin in the cellulosomal system [23, 25]. The study showed 1.7 fold increase in release of soluble sugar as compared to the wild-type enzymes [23].

1.2.0.1.2.1 Xylan binding domain in designer cellulosome

The xylanase were employed to deconstruct the hemicelluloses component plant based lignocellulosic component [5]. Xylanases engineered in designer cellulosome system in deconstruct the cellulosic biomass more efficiently [5]. Xylanases are important because it is used in different industry for example textile, biorefinary, food and pharmaceutical [6]. In one of the study, an entire xylanolytic complex of the bacterium *Thermobifida fusca* has been incorporated into an artificial cellulosome using designer cellulosome approach [24, 26].

1.3.0 Conclusion

The efficient enzyme with multi-functionality can be used for efficient degradation of cellulosic substrates. The multi-functionality can be introduced through protein engineering and designing artificial designer cellulosome. The engineered enzymes are utilized as cost effective process for enzymatic hydrolysis in deconstructing plant based lignocellulosic biomass and the technology can be transferred to biorefinery industry.

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