The engineering of xylo-oligosaccharide-consuming *Saccharomyces cerevisiae* strains is a promising approach for more effective utilization of lignocellulosic biomass and the development of economic industrial fermentation processes. Extending the sugar consumption range without catabolite repression by including the metabolism of oligomers instead of only monomers would significantly improve second-generation ethanol production. This review focuses on different aspects of the action mechanisms of xylan-degrading enzymes from bacteria and fungi, and their insertion in *S. cerevisiae* strains to obtain microbial cell factories able of consume these complex sugars and convert them to ethanol. Emphasis is given to different strategies for ethanol production from both extracellular and intracellular xylo-oligosaccharide utilization by *S. cerevisiae* strains. The suitability of *S. cerevisiae* for ethanol production combined with its genetic tractability indicates that it can play an important role in xylan bioconversion through the heterologous expression of xylanases from other microorganisms.

**Keywords:** *Saccharomyces cerevisiae*, xylo-oligosaccharides, lignocellulosic ethanol, xylanases, xylose

**BACKGROUND**

High fermentative capacity and robustness make *Saccharomyces cerevisiae* the microorganism of choice for ethanol production. *S. cerevisiae* strains are the most widely used in the ethanol industry with a well-characterized genome sequence, besides being an efficient expression system for recombinant enzyme production (Hou et al., 2012; Fang et al., 2017; Oh and Jin, 2020). Industrial *S. cerevisiae* strains are highly tolerant against various stresses in industrial environments, such as low pH, high osmotic pressure, high alcohol concentration, and phage contamination. In addition, this yeast presents high ethanol productivity, resulting from the naturally selective process that occurs during the successive fermentations involving pitching between fermentation tanks (Della-Bianca et al., 2014; Nielsen, 2019).

Economic production of second-generation biofuels should include the efficient and simultaneous co-fermentation of all hydrolysable sugars derived from cellulose and hemicellulose (Li et al., 2015). Lignocellulose-derived ethanol provides environmental and
economic benefits, as significant reductions in the disposal of solid wastes, and less air pollution, besides representing a promising industry in the expected transition from fossil to renewable fuels and chemicals. This biofuel is an environmentally friendly liquid fuel because the exhaust carbon dioxide is being taken up by growing plant biomass, contributing to the reduction of CO₂ content in the air, which in turn, contributes to the equilibrium of the earth’s atmosphere (Palmqvist and Hahn-Hägerdal, 2000a; Jacobsen and Wyman, 2000; Galbe and Zacchi, 2002).

Hemicellulose and cellulose are the major components of the secondary layers of the cell wall in wood fiber, shaping the well-known natural composition of, lignocellulosic biomass, along with lignin and minor components such as extractives and minerals (Gírio et al., 2010). Lignocellulose represents the most abundant source of renewable material on earth. This material can be found in agricultural residues, forestry waste, municipal solid waste, woods, and grasses, making them widely available at low cost, which is advantageous to the industrial context (Palmqvist and Hahn-Hägerdal, 2000b; Dahlman et al., 2003). Hemicellulose is a heterogeneous group of polysaccharides that comprises 15–35% of plant biomass (Gírio et al., 2010). Achieving 31.4% in switchgrass (Sun and Cheng, 2002), 29.3% in willow (Jorgensen et al., 2007), 28.6% in sugarcane bagasse (Fernandes Pereira et al., 2011), 22.1% in corn stover (Jorgensen et al., 2007), 19.7% in birchwood (Jorgensen et al., 2007) and 18% in spruce (Tengborg et al., 1998). Besides different amounts, the distribution of hemicellulose varies significantly between different plants. Depending on the source of the biomass (softwoods or hardwoods), its structure and composition can also vary. Softwood hemicellulose (pine and spruce, for example) presents a higher proportion of mannose and glucose units than hardwood hemicellulose (such as Eucalyptus, willow, and oak), which in turn, has a higher ratio of xylose units typically acetylated (Palmqvist and Hahn-Hägerdal, 2000b; Dahlman et al., 2003). The dominant hemicellulose polymer in hardwood biomass, xylan, is composed of repeating β (1–4)-linked xylose residue backbone, with acetyl and (methyl)glucuronic acid side groups. However, variations exist in its structures between different species (Rennie and Scheller, 2014; Wierzbicki et al., 2019).

Until recently, xylan represented the main component of plant biomass that cannot be efficiently utilized for biofuels production by fermentation using modified S. cerevisiae yeast strain. However, in 2004, Katahira and coauthors first demonstrated that a xylose-consuming S. cerevisiae strain expressing xylanolytic enzymes was able to produce ethanol from hemicellulose fraction although in lower levels (Katahira et al., 2004). Xylo-oligosaccharides (XOS)-consuming S. cerevisiae strains can represent an essential step to reach a more cost-effective second-generation ethanol production, conferring three significant advantages: 1) less intensive pre-treatment conditions would be required – harsh lignocellulosic pretreatment has been applied to release monomers (fermentable sugars), however during this process several yeast growth inhibitors are formed, such as furans, organic acids, phenols, and inorganic salts. Different aspects can interfere with the severity of the pretreatment process, which include holding time, pH, and temperature (Pedersen and Meyer, 2010). The lower severity process can result in high amounts of oligosaccharides, lower monosaccharides, and lower inhibitors compounds, as presented by Brenelli et al. (2020). In their study, the authors evaluated the effect of a mild decayetation treatment accomplished by hydrothermal pretreatment of raw sugarcane straw and achieve 81.5% of soluble hemicellulose with XOS yields up to 9.8% (w/w of an initial straw). These investigators found that an increase in the pretreatment temperature from 180 to 210°C, achieving a severity factor greater than 4, was accompanied by an increase in xylose production and lower oligosaccharides production. Under a lower severity factor condition (3.95) the crude hydrolysate yielded approximately 13.5 g L⁻¹ soluble XOS as well low amounts of arabinose, xylose, formic acid, acetic acid, and furfural were obtained. Increased temperature is related to an increase in the severity of the treatment, resulting also in the formation of inhibitors for both the enzymatic and fermentation processes (Pedersen and Meyer, 2010). It is worth mentioning that the depolymerization of cellulose and solubilization of hemicellulose and lignin vary according to the proposed pretreatment process and the severity factor applied in the respective process (Lynd et al., 2002). Moreover, the preparation of hemicellulose hydrolysate includes acid addition, high pressure and temperature which cause environmental pollution and equipment corrosion; therefore, successful ethanol production through XOS fermentation would make the process more environmentally friendly (Woodward and Wiseman, 1982; Gueguen et al., 1997; Nevoigt, 2008; Li et al., 2013); 2) lower demand for xylanolytic enzymes would be required, achieving lower production costs—the biomass enzymatic hydrolysis is a crucial step in the overall process due to its relatively large contribution to the total cost of lignocellulosic-derived ethanol (Nieves et al., 1997; Galbe and Zacchi, 2002); to maximize xylose yield and minimize the production of inhibitors, higher amounts of xylanolytic enzymes are required for total degradation of xylan and XOS which is prohibitively expensive on an industrial scale (Galbe and Zacchi, 2002), milder pretreatment methods have been described and 3) industrial competitive advantages (mainly for recombinant microorganism which are able to uptake and consume XOS internally)—it is expected that XOS-consuming S. cerevisiae strains would have a competitive advantage concerning other microorganisms, such as contaminating bacteria and wild Saccharomyces and non-Saccharomyces species that naturally use xylan as carbon source (Cabirini and Gallo, 1999; Amorim et al., 2011). It is important to point out that, in order to obtain a second-generation ethanol cost-competitive with first-generation ethanol, it is crucial to obtain microorganisms with unique genotype features to hydrolyze hemicellulose internally through recombinant DNA technology, which represents the best option to overcome the barriers to the commercial exploitation of lignocellulosic bioethanol.

Heterologous expression of xylose and XOS-producing enzymes in S. cerevisiae has been extensively reported. However, only one study has reported an S. cerevisiae strain able to break xylan down in an intracellular environment (Li et al.,
2015). For this reason, although there are engineering efforts to improve direct xylan utilization by this microorganism, some limitations still remain, such as the affinity between XOS and cell membrane transporters, and the understanding of metabolic pathways regulation. This overview examines all strategies reported to date adopted for the re-construction of XOS assimilation in S. cerevisiae yeast strains, focusing on those that bioethanol could be bio-converted from hemicellulose fraction.

LATEST TRENDS IN XYLOSE-UTILIZING S. CEREVISIAE

Although S. cerevisiae strains present all genes required for the xylose fermentation, i.e., xylose reductase (XR), xylitol dehydrogenase (XDH), and xylulokinase (XKS1), only XKS1 has been functionally expressed. XKS1 phosphorylates xylulose into xylulose-5-phosphate which is introduced into the central metabolism through the pentose-phosphate pathway. Previous studies have reported that the wild type of S. cerevisiae is capable of naturally assimilating xylulose as a sole carbon source, although at a low rate, under aerobic conditions. However, xylulose is a rare pentose not widely available in nature and probably due to this, the challenge of directing xylulose fermentation by S. cerevisiae has received little attention (Jeffries, 1983; Eliasson et al., 2000; Mittelman and Barkai, 2017; Patiño et al., 2019). Furthermore, xylose fermentation by S. cerevisiae requires additional interventions in endogenous genes expression and/or kinetic properties (Patiño et al., 2019).

With a focus on second-generation bioproducts, it is not surprising that many studies have attempted to develop laboratory and industrial engineered S. cerevisiae strains capable of simultaneous glucose and xylose fermentation by the expression of heterologous xylose consumption genes (Eliasson et al., 2000; Kuyper et al., 2005; Kwak and Jin, 2017; Li et al., 2019). Many studies have shown that different mutations can improve xylose fermentation by yeast. In Table 1 we benchmark five xylose-utilizing strains with superior ethanol yields on xylose metabolism.

In nature, pentose assimilation is widespread across many prokaryotes and eukaryotes, such as Pseudomonas fragi (Weinberg, 1961), Kluyveromyces lactis (Margaritis and Bajpai, 1982), Scheffersomyces stipitis (Toivola et al., 1984), Candida shehatae (Toivola et al., 1984), Pachysolen tannophilus (Smiley and Bolen, 1982; Toivola et al., 1984), Trichoderma sp. (Kulkarni et al., 1999), Aspergillus sp. (Kulkarni et al., 1999), Cryptococcus adeliiae (Petrescu et al., 2000), Pseudooleromas halolanklitis (Van Petegem et al., 2002), Hansenula polymorpha (Ryabova et al., 2003), Bacillus halodurans (Honda and Kitaoka, 2004), Bacillus subtilis (Collins et al., 2006), Caulobacter crescentus (Stephens et al., 2007), Plectosphaerella cucumerina (Zhang et al., 2007), Haloferax volcanii (Johnsen et al., 2009), Aureobasidium pullulans (Yegin, 2017). To date, three different pathways for xylose assimilation have been identified in these microorganisms and they are differentiated by the involvement of a phosphorylation step (Figure 1). In the first possibility, xylose is isomerized to xylulose and then phosphorylated to form xylulose-5-P. Two metabolic pathways have been identified which involve this strategy: the redox pathway, involving the combined activity of pyridine-nucleotide-dependent xylose reductase (XR) and xylitol dehydrogenase (XDH), and the isomerization pathway involving the redox-cofactor-independent xylose isomerase (XI). The main

| Strain | Parental strain | Relevant genotype/features | Culture conditions | Xylose specific consumption rate | Ethanol production rate | Ethanol productivity | Ethanol yield (g gxylose −1) | Reference |
|--------|----------------|----------------------------|-------------------|-------------------------------|------------------------|----------------------|--------------------------|----------|
| LYY34.4 | PE-2 (MATa) | XIX-Orpinomyces sp., Xyla, XKS1, TAL1, RKO1, TKL1, RPE1, Delta3, evolved | Microaerobic batch, YPX, 3% xylose, ICW 0.25 g DCW L−1 | 1.320 (g g−1 h−1) | 0.620 (g g−1 h−1) | ND | 0.460 | Dos Santos et al. (2016) |
| XUSE | BY4741 (MATa his3 leu2 met15 ura3) | XIX-Promyces sp. Xyla, XKS1, TAL1, Delta3, apho13, evolved | Microaerobic batch, YSC, 2% xylose, OD600 10 | ND | ND | ND | 0.400 | Tran Nguyen et al. (2018) |
| IMU078 | CEN.PK113-5D (MATa ura3) | XIX-Promyces sp. Xyla, RPE1, RKO1, TAL1, TKL1, NQM1, TKL2, XKS1, Delta3 | Anaerobic batch, Synthetic medium with L-aspargin instead of ammonium sulfate, 2% xylose, ICW 0.02 g DCW L−1 | ND | ND | ND | 0.406 | Bracher et al. (2019) |
| SR8N | D425-2 (MATa his3 leu2 ura3) | XR/XDH-S. stipitis Xyl1, Xyl2 and YL31, Lactococcus lactis NoxE, Delta13, Aald6 | Microaerobic batch, YNB, 4% xylose, OD600 10 | ND | ND | 1.220 (g L−1 h−1) | 0.391 | Kim et al. (2013b), Zhang et al. (2017b) |
| YRH1490 | PE-2 (MATa) | XR/XDH-S. stipitis Xyl1 and Xyl2, XKS1 | Microaerobic batch, YPX, 8% xylose, OD600 1 | ND | ND | 0.310 (g L−1 h−1) | 0.330 | Dias Lopes et al. (2017) |

ICW, initial cell weight. OD600 = Initial OD600. ND, no data available.

TABLE 1 | Literature data on engineered, xylo-oxidoreductase and xylose-isomerase -based S. cerevisiae strains.
difference between these pathways is the dependence on cofactors (oxido-reduction pathway) or not (isomerization pathway). Both metabolic pathways have been used extensively as targets in engineered *S. cerevisiae* and have been reviewed in detail (Kim et al., 2012, Kim et al., 2013; Harner et al., 2015; Kwak and Jin, 2017; Bracher et al., 2019). The generation of xylulose-5-phosphate via the oxidoreductase pathway allows a link to glycolysis, the central carbon flux, through the non-oxidative part of the pentose phosphate pathway (Stincone et al., 2015). Optimized *S. cerevisiae* recombinant strains overexpressing XI or XR/XDH have been reported (Table 1). The success of these strategies enables new perspectives on the carbon-source range assimilated by *S. cerevisiae* to be considered.

Since xylose assimilation by engineered *S. cerevisiae* strains has become well-established, new approaches have been adopted to enable *S. cerevisiae* to consume XOS instead of xylose and glucose (La Grange et al., 2000, 2001; Fujita et al., 2002; Qian et al., 2003; Katahira et al., 2004; Lee et al., 2009; Sun et al., 2012; Li et al., 2015; dos Reis et al., 2016; Sekar et al., 2016; Zhang et al., 2017a). Scientific interest in this field is increasing steadily, but still much must be done to obtain an efficient XOS-consuming *S. cerevisiae* strain.

An additional xylose assimilation possibility is the so called Weimberg pathway, which is characterized as an oxidative but non-phosphorylating metabolic pathway without xylose to xylulose isomerization. This pathway received much less attention when compared with oxido-reductase and isomerase pathways but recently has gained major attention from research groups (Shen et al., 2020). This pathway provides an alternative entry point for xylose into yeast central metabolism with possibilities to produce new compounds that are intermediates or derivatives from the TCA cycle and provides an interesting route for the production of xylose-derived α-ketoglutarate (Figure 1) (Weimberg, 1961). Recently, the Weimberg pathway enzymes derived from *Caulobacter crescentus* and *Corynebacterium glutamicum* were functionally expressed in *S. cerevisiae*; however, pathway intermediates were detected, indicating that this pathway needs further optimization (Borgström et al., 2019). Some of the drawn hypotheses could explain this incompatibility between prokaryotic and eukaryotic proteins, such as deficiency of enzymatic cofactors, posttranslational modifications of the protein, differences in the internal pH of the parental and the host cell (Sarthy et al., 1987), and the improper folding of the protein (Gárdonyi and Hahn-Hägerdal, 2003; Xia et al., 2016).

### XYLANOLYTIC ENZYME SYSTEMS

The most selective method for the conversion of poly- to monosaccharides is by using enzymes. Most studies on hemicellulases have focused on xylanolytic enzymes, which are responsible for xylan hydrolysis (Gírio et al., 2010). The study of microorganisms able to hydrolyze xylan started more than 130 years ago, probably in 1889 by Hoope-Seyler (Whistler and Masak Jr., 1955). Since then, many organisms with the
ability to colonize and grow on plant biomass have been identified. Xylanolytic enzyme producers are widespread, such as fungi, bacteria, yeast, marine algae, protozoans, snails, crustaceans, and insects, of which bacterial and fungal xylanases have the most important role concerning heterologous expression in *S. cerevisiae* (Jeffries, 1983; Biely, 1985; Kulkarni et al., 1999; Katahira et al., 2004; Schmoll and Schuster, 2010). *Trichoderma reesei*, *Trichoderma atroviride*, *Trichoderma virens* (Beg et al., 2001), *Aspergillus niger* (La Grange et al., 2001), *Neurospora crassa* (Li et al., 2015), *Aspergillus foetidus* (Whistler and Masak Jr., 1955), *Bacillus pumilus* (Pan et al., 1991) are some examples of potent xylanolytic enzymes producers. These enzymes have potential for the application of xylanases in several industries, such as in the pulp and paper, food additives, animal feed, textiles, drinks industries, ethanol, and xylitol production (Polizeli et al., 2005). The search for newer microbial xylanases producers is ongoing, together with molecular biology studies on the regulation of xylanases expression and their heterologous expression in non-xylanolytic microorganisms.

Xylans represent a family of complex non-cellulosic branched polysaccharides that consists structurally of linear homopolymeric β-(1,4)-xylopyranosyl units with a diversity of substituted groups, which vary quantitatively and qualitatively according to the plant or the method of isolation. They can be comprised of 4-O-methyl-D-glucuronoxylan, α-L-arabinofuranosyl, acetyl, feruloyl and/or p-coumaroyl groups (Figure 2) (Wong et al., 1988; Collins et al., 2005; Biely et al., 2016). Wood xylan exists as O-acetyl-4-O-methylglucuronoxylan in hardwoods and as arabin-4-O-methylglucuronoxylan in softwoods, which represent the two major forms of xylan in wood, whereas xylans in grasses and annual plants are typically arabininoxylans (Kulkarni et al., 1999). On the other hand, in esparto grass, tobacco stalks, and guar seed husk another type of xylan has been identified, the homoxylans which are composed exclusively of xylosyl residues (Sunna and Antranikian, 1997).

Due to the heterogeneity and complex chemical nature of xylan, its complete breakdown requires the action of a consortium of enzymes with diverse specificities and modes of action (Figure 2). Thus, it is not surprising that xylan-degrading organisms produce a multienzyme system of xylanases that present diverse structures, different mechanisms of action, substrate specificities, hydrolytic activities, and physicochemical characteristics (Beg et al., 2001; Collins et al., 2005; Moreira and Filho, 2016). It is interesting to note that microorganisms can produce multiple forms of the same xylanase family, showing that some factors such as differential processing of mRNA, post-translational modification, proteolytic digestion, and differential expression by distinct alleles of one gene, or even by completely separate genes affect this multiplicity of xylanases secreted (Polizeli et al., 2005; de Vries et al., 2017).

The carbohydrate-active enzymes (CAZy—www.cazy.org) database collectively complies and assigns xylanases are glycoside hydrolases (GH) that catalyze the hydrolysis of 1,4-β-D-xylosidic linkages in xylan. Sequence-based classification has grouped xylanases in two major families GH10 and GH11, but xylanases are also found in other GH families, 3, 5, 8, 9, 10, 11, 12, 16, 26, 30, 43, 44, 45, 51, 62, 98, and 141 (Collins et al., 2005; Chakdar et al., 2016). GH10 members are highly active on short XOS, thereby indicating small substrate-binding sites. The major enzymes of this family are endo-β-1,4-xylanases (Collins et al., 2005). In contrast, the enzymes that belong to the GH11 family are most active on long-chain XOS. Furthermore, this family is monospecific, which means they are exclusively active on D-xylene-containing substrates (Collins et al., 2005; Chakdar et al., 2016). Some genera of fungi and bacteria produce more than one subfamily of xylanases. However, the majority of the bacterial xylanases belong to the GH10 family, whereas fungal xylanases majorly belong to the GH11 family (Liu et al., 2011).

There is a great diversity of xylanases produced among bacterial genera with *Bacillus* presenting a potential source of these enzymes (Subramaniam and Prema, 2002; Gupta and Verma, 2015; Chakdar et al., 2016). Xylanases obtained from bacterial sources are known to be active and stable in a wide range of pH and temperature, such as temperature from 30 to 60°C, pH 5.2-6.0. **Figure 2** | Xylan degradation scheme. The arrows represent each enzyme activity for a determined substrate. Adapted from Bhardwaj et al. (2019).
Table 2: Characteristics of xylanase from different microorganisms functionally expressed in S. cerevisiae.

| Organism        | Enzyme                  | Cloning process                  | Host                      | Optimum Temperature | pH  | Activity in S. cerevisiae | Reference                     |
|-----------------|-------------------------|----------------------------------|---------------------------|---------------------|-----|--------------------------|-------------------------------|
| B. pumilus PLS  | β-xylosidase            | pDLG12—2µ yeast plasmid          | S. cerevisiae Y294        | 45–50°C             | 6.6 | 0.09 nkat ml⁻¹           | La Grange et al. (1996, 1997) |
|                 |                         |                                  |                           |                     |     |                          |                               |
| B. pumilus IPO  | Xylanase (xynA)         | pNAX2—2µ yeast plasmid           | S. cerevisiae NA87-11A    | 40°C                | 6.5 | 0.36 U (mg protein)⁻¹    | Panbanged et al. (1983), Pan et al. (1991) |
|                 |                         |                                  |                           |                     |     |                          |                               |
| B. pumilus IPO  | β-xylosidase (xynB)     | pYXB—2µ yeast plasmid            | S. cerevisiae NA87-11A    | ND                  | ND  | 0.28 U (mg protein)⁻¹    | Pan et al. (1991)              |
|                 |                         |                                  |                           |                     |     |                          |                               |
| Caldoccillium   | Xylanase (xynA)         | pFGxyn—2µ yeast plasmid          | S. cerevisiae STX929-3A   | ND                  | ND  | 90 U (mg protein)⁻¹      | Donald et al. (1994)           |
| saccharolyticum |                         |                                  |                           |                     |     |                          | Kim et al. (2003)              |
| Bacillus sp.    | Xylanase (xynD)         | pBX45—2µ yeast plasmid           | S. cerevisiae SEY2102     | 58°C                | 6.2 | 8.5 nkta mL⁻¹            | Nuyens et al. (2001)           |
|                 |                         |                                  |                           |                     |     |                          |                               |
| Penicillium     | Xylanase (xynA)         | pYEplac181—integrating plasmid   | S. cerevisiae Y3M35::RY171| ND                  | ND  | 4.52 U mL⁻¹              | Ma and Ptashe (1987), Chávez et al. (2002) |
| purpurogenum    |                         |                                  |                           |                     |     |                          |                               |
| ATCC No. MYA-38 |                         |                                  |                           |                     |     |                          |                               |
| Bacillus spp.   | Xylanase (xynG)         | pAGX3—2µ yeast plasmid           | S. cerevisiae SEY2102     | ND                  | ND  | 7.56 U mL⁻¹              | Lee et al. (2007)              |
|                 |                         |                                  |                           |                     |     |                          |                               |
| Bacillus spp.   | Xylanase (xynH)         | pAX4-1—2µ yeast plasmid          | S. cerevisiae SEY2102     | ND                  | ND  | 9.8 U mL⁻¹               | Lee et al. (2009)              |
|                 |                         |                                  |                           |                     |     |                          |                               |
| Bacillus sp.    | Xylanase (XyP)          | pGMF-xyp—2µ yeast plasmid        | S. cerevisiae SEY2102     | ND                  | ND  | 70.1 U mL⁻¹              | Kim et al. (2013a)             |
| HY-20           |                         |                                  |                           |                     |     |                          |                               |
| Bacillus sp.    | Xylanase (XyP)          | pGMF-xyp—2µ yeast plasmid        | S. cerevisiae FY833       | ND                  | ND  | 42.4 U mL⁻¹              | Kim et al. (2013a)             |
| HY-20           |                         |                                  |                           |                     |     |                          |                               |
| Bacillus sp.    | Xylanase (XynHβ)        | pHBM367H—rDNA-mediated integration | S. cerevisiae INV         | ND                  | ND  | 255 U (g DWcell)⁻¹       | Fang et al. (2017)             |
| HBp8            |                         |                                  |                           |                     |     |                          |                               |
| Fungal species  | Xylanase (XLN)          | pVT100—2µ yeast plasmid          | S. cerevisiae T109-3C     | ND                  | ND  | 1.3 U (mg protein)⁻¹     | Moreau et al. (1992)           |
| Cryptococcus    |                         |                                  |                           |                     |     |                          |                               |
| albidus         |                         |                                  |                           |                     |     |                          |                               |
| Aspergillus     | β-xylosidase (xynC)     | pJC1—2µ yeast plasmid            | S. cerevisiae Y294        | 60°C                | 3   | 300 nkat ml⁻¹            | Crous et al. (1995)            |
| kawachi         |                         |                                  |                           |                     |     |                          |                               |
| IFO4308         |                         |                                  |                           |                     |     |                          |                               |
| Aureobasidium   | Xylanase II (XynA)      | 2µ yeast plasmid                 | S. cerevisiae INVSc1      | ND                  | ND  | 32.9 U ml⁻¹              | Li and Ljungdahl, (1996)       |
| pullulans Y-2311–1 |                       |                                  |                           |                     |     |                          |                               |
| A. nidulans G191 | Xylanase (xlnA)         | pYLA1—2µ yeast plasmid           | S. cerevisiae OL1         | ND                  | ND  | 65 U ml⁻¹                | Pérez-González et al. (1996)   |
|                 |                         |                                  |                           |                     |     |                          |                               |
|                 | Xylanase (xlnB)         | pYLB1—2µ yeast plasmid           | S. cerevisiae OL1         | ND                  | ND  | 25 U ml⁻¹                | Pérez-González et al. (1996)   |

(Continued on following page)
### TABLE 2 (Continued) Characteristics of xylanase from different microorganisms functionally expressed in S. cerevisiae.

| Organism           | Enzyme           | Cloning process               | Host                                | Name                  | Optimum Activity in S. cerevisiae | Activity in S. cerevisiae | Remarks                          | Reference             |
|--------------------|------------------|-------------------------------|-------------------------------------|-----------------------|----------------------------------|---------------------------|---------------------------------|------------------------|
| T. reesei RutC-30 | α-Arabinofuranosidase (AbfB) | p17SA—2µ yeast plasmid       | S. cerevisiae                      | DBY746               | MATa his3 Δ1 leu2-3 112 ura3-52 trp1-289 cyHR | ND ND                       | 171.1 nkat ml−1         | Margolies-Clark et al. (1996) |
| A. niger ATCC 90196 | β-xylosidase   | pMLU1—2µ yeast plasmid       | S. cerevisiae                      | Y294                  | MATa leu2-3 112 ura3-52 his3 trip1-289 Δpfr1 | ND ND                       | 91 nkat ml−1           | Luttig et al. (1997)         |
| T. reesei          | Xylanase II (XYNII) | pCAS1—2µ yeast plasmid       | S. cerevisiae                      | MT8-1                 | MATa ade leu2 ura3 his3 trip1 | ND ND                       | 1.78 µmol min−1 (g DWcell)−1 | Fujita et al. (2002)       |
| Trichoderma spp.  | Endoglucanase   | pAGX1—2µ yeast plasmid       | S. cerevisiae                      | SEY2120               | MATa leu2-112 ura3-52 his4-519 suc2-Δ9 leu2 | ND ND                       | 0.6 U ml−1            | Lee et al. (2007)            |
| C. flavus I-11    | Xylanase (CfXYN) | Yep351PGK—2µ yeast plasmid   | S. cerevisiae                      | MFL                   |                                         | 50°C 3                      | 2.5 U ml−1           | Parachin et al. (2009)        |
| A. niger IME-216   | Xylanase        | pUJPXR—integrating plasmid   | S. cerevisiae                      | YS2_2                 | Industrial ethanol producing strain | ND ND                       | 74.8 U ml−1           | Tian et al. (2013)           |

**Ethanol production has been reported**

**Fungal species**

| Organism           | Enzyme           | Cloning process               | Host                                | Name                  | Optimum Activity in S. cerevisiae | Activity in S. cerevisiae | Remarks                          | Reference             |
|--------------------|------------------|-------------------------------|-------------------------------------|-----------------------|----------------------------------|---------------------------|---------------------------------|------------------------|
| A. oryzae NiaD300 | β-xylosidase (Xyla) | pUCSIIIXA—cell-surface expressing plasmid | S. cerevisiae                      | MT8-1                 | MATa ade leu2 his3 ura3 can1 SoXY1 SoXYL2 SoXXS1 | ND ND                       | 234 U (g DWcell)−1 | Katahira et al. (2004)       |
| T. reesei QM9414   | Xylanase II (XYNII) | pAUR-XSD—2µ yeast plasmid   | S. cerevisiae                      | MA-D4                 | MATa leu2 his3 ura3 can1 SoXY1 SoXYL2 SoXXS1 Δaur1 | ND ND                       | 16 U (g DWcell)−1 | Fujii et al. (2011)          |
| T. reesei QM9414   | β-xylosidase     | pUCSIIYXII—integrating plasmid | S. cerevisiae                      | OC-2                  | MATa a SoXYL1 SoXYL2 SoXXS1 | 60°C ND                       | 6 nmoi min−1 (mg protein)−1 | Sakamoto et al. (2012) |
| T. reesei          | β-xylosidase (XylI) | p8W- GPAGXyI—integrating plasmid | S. cerevisiae                      | MT8-1                 | Mata ade leu2 trip1 ura3 SoXY1 SoXYL2 SoXXS1 | ND ND                       | 41.2 U (g DWcell)−1 | Sakamoto et al. (2012) |
| A. oryzae          | β-xylosidase (Xyla) | pHGXYLA—integrating plasmid  | S. cerevisiae                      | EBY100 ( INVitrogen, Carlsbad, CA) | SoXYL1 SoXYL2 SoXYL3 | ND ND                       | 16.8 U (g DWcell)−1 | Sun et al. (2012)           |
| T. reesei QM9414   | Xylanase II (XYNII) | pYD1—2µ yeast plasmid       | S. cerevisiae                      | EBY100 ( INVitrogen, Carlsbad, CA) | SoXYL1 SoXYL2 SoXYL3 | ND ND                       | ND ND               | ND                   |
| A. niger DSM821 A. niger DSM821 | β-xylosidase (XlnD) | α-arabinofuranosidase (AbfB) | S. cerevisiae                      | DBY746               | MATa his3 Δ1 leu2-3 112 ura3-52 trp1-289 cyHR | ND ND                       | ND ND               | ND                   |
| A. terreus          | Xylanase         | pSK2—2µ yeast plasmid       | S. cerevisiae                      | INVSc1                | MATa leu2 trip1-289 ura3-52 C2XR | ND ND                       | ND ND               | Li et al. (2013)          |
| N. crassa FGSC 2489 | β-xylosidase (GH43-2) | pXDO8.7—2µ yeast plasmid   | S. cerevisiae                      | SR8U                  | MATa ura3 SoXYL1 SoXYL2 SoXYL3 Δpfr13 Δald6 | ND 7                     | ND ND               | Kim et al. (2013), Li et al. (2015) |

(Continued on following page)
TABLE 2 | (Continued) Characteristics of xylanase from different microorganisms functionally expressed in *S. cerevisiae*.

| Organism          | Enzyme                  | Cloning process          | Host          | Remarks                     | Optimum Temperature | Activity in *S. cerevisiae* | Reference         |
|-------------------|-------------------------|--------------------------|---------------|-----------------------------|---------------------|-----------------------------|--------------------|
| U. beomyces       | Xylanase 1 (XNA1)       | P423—2µ yeast plasmid    | *S. cerevisiae* | Mata trp1 can1              | 30°C                | 1.94 U mg⁻¹                 | Lee et al. (2015)  |
|                   | Xylosidase 2 (X2D2)     | P424—2µ yeast plasmid    | *S. cerevisiae* | cy1 gal+ leu2::             | 30°C                | ND                          |                    |
|                   | Arabinofuranosidase (ABF)| P424—2µ yeast plasmid    | *S. cerevisiae* | LEU2-TDH3P-PxYX1-TDH3ST     | 30°C                | ND                          |                    |
|                   |                         |                          |               | ura3::URA3-TDH3P-PxYX2-TDH3ST|                     | ND                          |                    |
|                   |                         |                          |               | Tgy3::G418-PxYX3            |                     | ND                          |                    |
|                   |                         |                          |               | YOR202w::hnpNT1             |                     | ND                          |                    |
| T. reesei QM6a    | Endoxylanase (Xyn2)     | pVSDis-TrXyn2—cell-      | *S. cerevisiae* | Mata AGA1::                  | 30°C                | 1.197 U mg⁻¹                | Tabanag et al. (2018)|
|                   |                         | surface expressing       |               | GAL1-AGA1::                  |                     |                             |                    |
|                   |                         | plasmid                  |               | URA3 ura3-S2                 |                     |                             |                    |
|                   |                         |                          |               | trp1 leu2-A200              |                     |                             |                    |
|                   |                         |                          |               | his3-A200                    |                     |                             |                    |
|                   |                         |                          |               | pep4::HI3                    |                     |                             |                    |
|                   |                         |                          |               | prb11.6R can1                |                     |                             |                    |
|                   |                         |                          |               | GAL1 PrX1 PrXK1              |                     |                             |                    |
|                   | β-xylosidase (Bxl1)     | pVSDis-TrBxl1—cell-      | *S. cerevisiae* | EBY100                      | 30°C                | ND                          |                    |
|                   |                         | surface expressing       |               |                                |                     |                             |                    |
|                   |                         | plasmid                  |               |                                |                     |                             |                    |
|                   | Acetyl esterase (Axe1)  | pVSDis-TrAxe1—cell-      | *S. cerevisiae* |                                | 30°C                | ND                          |                    |
|                   |                         | surface expressing       |               |                                |                     |                             |                    |
|                   |                         | plasmid                  |               |                                |                     |                             |                    |
|                   | α-glucoronidase (Glr1)  | pVSDis-TrGlr1—cell-      | *S. cerevisiae* |                                | 30°C                | ND                          |                    |
|                   |                         | surface expressing       |               |                                |                     |                             |                    |
|                   |                         | plasmid                  |               |                                |                     |                             |                    |
|                   | α-arabinofuranosidase   | pVSDis-TrAbf1—cell-      | *S. cerevisiae* |                                | 30°C                | ND                          |                    |
|                   | (Abf1)                  | surface expressing       |               |                                |                     |                             |                    |
|                   |                         | plasmid                  |               |                                |                     |                             |                    |

*The final goal of this work was xyitol production from xylan. Li et al. (2015) achieved a xyitol yield of 0.71 g xyitol (g xylan)⁻¹, and S. cerevisiae recombinant strain, Sc-K2, produced 1.94 g L⁻¹ xyitol when cultivated in YPD supplemented with 3 g L⁻¹ xylan.

Ss, S. stipitis; Sc, S. cerevisiae; Ct, Candida tropicalis; Pr, Prevotella ruminicola.

ND, no data available.

from 5.0 to 9.0 (See some examples in Table 2). These enzymes are produced alone mostly, thereby reducing the downstream process time (Chakdar et al., 2016).

Despite the great diversity of bacterial xylanase producers, filamentous fungi are the major commercial source due to their higher levels of xylanase secretion (Kulkarni et al., 1999; Polizeli et al., 2005). Some works have demonstrated that many fungal species produce xylanase when cultured on cellulose (Mishra et al., 1984; Biely, 1985; Wong et al., 1988), perhaps because the cellulose substrates contain traces of hemicellulose (Polizeli et al., 2005). Nevertheless, in an opposite scenario, several fungal species produce specific xylanases with little or no cellulose background in the presence of xylan, which indicates the xylanolytic and cellulolytic systems are likely to be under separate regulatory control (Biely, 1985; Wong et al., 1988). And interestingly XOS reduced the efficiency of cellulase hydrolysis by cellulase (Zhang et al., 2012; Wang et al., 2018) which cannot be effectively relieved by increasing the loading of the cellulose substrate or cellulase (Wang et al., 2018). However, some fungi species require low pH for the growth and production of xylanases which necessitates additional steps in the subsequent stages which make fungal xylanases less attractive (Chakdar et al., 2016).

Among xylanases, endo-β-xylanases (xylanase or endo-β,1,4-xylanases) and β-d-xylosidases (β-xylosidases or xylosidase) have been most extensively studied. Endo-β-xylanases (EC 3.2.1.8) randomly cleave the β-1,4 linkages between the xyloprenosyl units from the xylan backbone, producing mixtures of XOS (Biely, 1985; Kulkarni et al., 1999). β-d-xylosidases (EC 3.2.1.37) are known to be the major component of xylanase systems. They are produced by biodegradative microorganisms to hydrolyze XOS releasing d-xylose; however, usually, they do not hydrolyze xylan, with their best substrate being xylobiose and their affinity for XOS being inversely proportional to its degree of polymerization. They act on the non-reducing ends of their substrate, XOS and/or xylobiose (Wong et al., 1988; Collins et al., 2005; Polizeli et al., 2005). A high concentration of xylose in the fermentation broth can inhibit the activity of β-d-xylosidases (Fujii et al., 2011) which leads to the inefficient hydrolysis of hemicellulose and the accumulation of XOS and xylobiose by using microorganisms that do not consume xylose rapidly. Some β-d-xylosidases have been reported to possess α-L-arabinofuranosidase activity, e.g., the enzymes from *A. niger*, *T. reesei*, *T. ethanolicus*, and *Penicillium wortmannin* (Sunna and Antranikian, 1997). Other enzymes, such as α-L-arabinofuranosidase (EC 3.2.1.55), α-d-glucuronidases (EC 3.2.1.139), acetylxyylan esterases (EC 3.1.1.72), ferulic acid esterases (EC 3.1.1.73), and p-coumaric acid esterases (EC 3.1.1.-) catalyze the removal of xylan side groups (Collins
et al., 2005). All these enzymes act cooperatively to convert xylans into xylose, XOS, O-acetyl, L-arabinose, acetic and glucuronic acids (Beg et al., 2001; Polizeli et al., 2005).

Understanding the enzymes xylanolytic microorganisms produce for hemicellulose breakdown may become an important tool for re-construction of their XOS degradation pathway in non-xylanolytic microorganisms. As mentioned before, complete degradation of xylan is achieved by a variety of modular enzymes. Although many hemicellulolytic enzymes have been studied extensively, little is known about how microorganism cells sense the presence of xylan and uptake hemicellulose-derived products (Polizeli et al., 2005; Delmas et al., 2012; Najjarzadeh et al., 2020). The induction signal for the synthesis of xylanolytic enzymes is supposed to involve transporters of xylose and short XOS released by the action of little amounts of the enzymes produced constitutively, along with lactose, glucose, and even cellulose, that are able to cross the membrane and induce the regulatory machinery (Biely and Petráková, 1984; Royer and Nakas, 1989; Chandra Raj and Chandra, 1995; Christakopoulos et al., 1996a, 1996b; Kulkarni et al., 1999), which suggests a complex induction mechanism of xylanases. In their study, Delmas et al. (Delmas et al., 2012) studied the strategy of the filamentous fungus A. niger employ to degrade complex polysaccharides. They showed that wheat straw itself is not initially detected by the A. niger. According to their findings, the overall strategy appears to be an induction of a specific, small scale, sensory response by the onset of carbon starvation, mediated at least partially by alleviation of CreA-dependent catabolite repression, that triggers the release of a small subset of degradative enzymes which initiate degradation on a small scale, in turn releasing sugars that cause the fungus to express its full degradative arsenal (Delmas et al., 2012). More recently, Najjarzadeh and collaborators reported that xylotetraose is more effective than other substrates inducing endoxylanase, while xylohexaose and endoglucanases (two from T. aurantiacus) is considered a key enzyme (Banerjee et al., 2010), since XOS accumulation can reduce the efficiency of cellulases, such as cellobiohydrolase I (CBHI, from Thermosascus aurantiacus), cellobiohydrolase II (CBHII, from Trichoderma reesei) and endoglucanase II (from T. aurantiacus) (Zhang et al., 2012), which would affect conversion yields in simultaneous saccharification and fermentation (SSF) or consolidated bioprocessing (CBP). Therefore, reducing the concentration of XOS using β-xylanolytic enzymes represents the best strategy to prevent enzyme inhibition.

The heterogeneous nature of hemicellulose represents a challenge for hemicellulase enzymes. Considering the first-generation ethanol industry, a huge amount of lignocellulose-based materials is formed during ethanol production, especially corn stover, sugarcane straw, and sugarcane bagasse, which are particularly attractive as second-generation ethanol feedstock. As mentioned earlier, their hemicellulose content can achieve up to 28% in sugarcane bagasse (Fernandes Pereira et al., 2011), 27–31% in sugarcane straw (Almeida and Colombo, 2021), and 22% in corn stover (Jorgensen et al., 2007). According to the chemical structure of hemicellulose, they present a very similar composition, which includes a high content of acetyl as side groups. Therefore, it is important to highlight that other enzymes would allow more efficient degradation of these hemicellulosic derived materials, such as acetylxylan esterases (EC 3.1.1.72) in combination with xylanase and xylosidase for hydrolyzing pretreated hardwood hemicellulose. The presence of acetyl group degrading enzymes may increase the accessibility of the xylose chain to xylanases.

**HETEROLOGOUS EXPRESSION OF XYLANOLYTIC ENZYME IN THE YEAST S. CEREVISIAE**

Several reports have described the expression of heterologous endo-β-xylanases and β-xylanolases from both prokaryotes and eukaryotes in *S. cerevisiae* to enable the conversion of xylan or XOS into xylose by this species (Table 2). Usually, the complete hydrolysis of xylan requires at least these two enzymes (Biely, 1985): an endo-β-xylanase that cleaves xylan into XOS with diverse degrees of polymerization, followed by the breakdown of XOS to xylose by β-xylanolases (Wang et al., 2018). Although endo-β-xylanases are important for the hydrolysis process, β-xylanolases is considered a key enzyme (Banerjee et al., 2010), since XOS accumulation can reduce the efficiency of cellulases, such as cellobiohydrolase I (CBHI, from Thermosascus aurantiacus), cellobiohydrolase II (CBHII, from Trichoderma reesei) and endoglucanase II (from T. aurantiicus) (Zhang et al., 2012), which would affect conversion yields in simultaneous saccharification and fermentation (SSF) or consolidated bioprocessing (CBP). Therefore, reducing the concentration of XOS using β-xylanolases represents the best strategy to prevent enzyme inhibition.

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**S. CEREVISIAE AS A PLATFORM TO PRODUCE XYLANOLYTIC ENZYMES**

To the best of our knowledge, the first report of xylan-degrading genes expression in *S. cerevisiae* was in 1991, when Pan et al. (1991) described the expression of two enzymes, xylanase (a xynA gene product) and β-xylanolase (a xynB gene product) from Bacillus pumilus in yeast cells. Thereafter, physiological data related to the expression of the intracellular β-xylanolase from *B. pumilus* in *S. cerevisiae* has been published (Crous et al., 1995; Crous et al., 1996; La Grange et al., 1996; La Grange et al., 2000). A successful expression of a bacterial β-xylanolase from *B. pumilus* (xynB)
in yeast cells was achieved by its fusion to a native secretion signal sequence named mating pheromone α-factor (MFα1s) (La Grange et al., 1997). The native open-reading frame of these enzymes starts with the codon TTG which is not recognized by S. cerevisiae for the initiation of translation. Even after replacing the TTG codon with an ATG starting codon, no β-xilosidase activity could be detected by the recombinant S. cerevisiae Y294 (La Grange et al., 1997).

Although there has been some success in functional expression of bacterial enzymes in S. cerevisiae (Table 2), this can be problematic, possibly because of incompatibility with eukaryotic chaperones (Sarthy et al., 1987; Gárdonyi and Hahn-Hägerdal, 2003; Xia et al., 2016). The heterologous expression of eukaryotic xylanases in S. cerevisiae naturally shows more compatibility since fungal species share many features, particularly related to transcription, translation, and protein stability (Frommer and Ninnenmann, 1995). The first recombinant S. cerevisiae strain (namely IAF130) expressing heterologous eukaryotic xylanase was described by Moreau et al. (1992). However, because the cells could not catabolize xylose, the majority of the early reports of recombinant S. cerevisiae expressing β-xilosidases only demonstrated the conversion of xylan and XOS into xylose, xylobiose, and xylotriose, but not to ethanol.

Heterologous expression of the T. reesei xylanase II (XYNII) anchored on the cellular surface was described in the S. cerevisiae strain MT8 by using a cell surface engineering system based on α-agglutinin which consist of the fusion of the protein with the C-terminal half region of an agglutinin. The recombinant strain MT8-1/pCAS1-XYNII was able to hydrolyze birchwood xylan into xylose and xylotriose. The proposed work did not aim to present a hydrolytic process during the growth of MT8-1/pCAS1-XYNII in a medium containing complex sugar. Instead, xylanase activity was measured in both supernatant and pellet fractions from pre-cultured strain. XYNII activity was detected in the cell pellet with no leakage into the supernatant medium (Fujita et al., 2002).

**ETHANOL PRODUCTION FROM XOS BY EXTRACELLULAR EXPRESSION OF Xylanolytic Enzymes in S. Cerevisiae Strains**

The first example of the concept of CBP applied to ethanol production from xylan using recombinant S. cerevisiae strain without the addition of exogenous xylan-degrading enzymes was described in 2004 (Katahira et al., 2004). In their work, xylanase II (XYNII) from T. reesei QM9414 and β-xilosidase (Xyla) from Aspergillus oryzae NiaD300, were co-displayed on the cell surface of xylose-consuming S. cerevisiae harboring genes encoding the oxidoreductase pathway from S. stipitis and native xylulokinase (XKS) from S. cerevisiae. To obtain this strain, the C-terminal region of α-agglutinin was fused to both xylanolytic enzymes. The constitutive expression of XYNII and Xyla enabled xylan consumption and ethanol production without a lag-phase. The recombinant strain MT8-1/pUCSXIIXA/pWX1X2XK produced 7.1 g L\(^{-1}\) of ethanol after 62 h of fermentation in semi defined medium supplemented with birchwood xylan corresponding to 100 g of total sugar per liter as the sole carbon source. Despite the significant ethanol production, a large amount of xylan remained in the growth medium, suggesting that this strain needs further optimization.

Microbial surface display technology allows the expression of peptides and proteins on the surface of living cells in which proteins are expressed extracellularly, however, the enzymes remain fused at the cell with no leakage into the culture medium (Fujita et al., 2002; Katahira et al., 2004; Tafakori et al., 2012). Other examples of coexpression of xylanolytic enzymes anchored on the S. cerevisiae cellular surface considered the expression of bifunctional minihemicellulolysomes, with several assembled modules included (Sun et al., 2012). Sun and coauthors constructed a recombinant yeast that directly produced ethanol from birchwood xylan through the expression of bifunctional minihemicellulolysomes. This recombinant strain co-displays two complementary xylanases, XYNII and an A. niger xylosidase (XDNL) as well as a mini scaffolding (CipA3), which served as the basis to establish interaction between the enzymes and cell surface. According to their findings, the HZ3345 strain was able to ferment xylan into ethanol. The recombinant S. cerevisiae strain also contained an integrated xylose-utilizing pathway (XR, XDH, and XK from S. stipitis) to ensure the xylose assimilation. Interestingly, xylose production was immediately observed from XOS, without any lag phase, as previously observed (Katahira et al., 2004). The recombinant HZ3345 strain, produced 0.95 g L\(^{-1}\) of ethanol from approximately 3.0 g L\(^{-1}\) birchwood xylan after 80 h of cultivation under anaerobic conditions in YPBX (YP supplemented with birchwood xylan) supplemented with Tween and ergosterol.

In a more recent study, a blended bioprospecting approach was applied (Lee et al., 2015) along with rational and evolutionary engineering to improve xylan assimilation in an engineered xylan-catabolizing S. cerevisiae strain. The extracellular expression of xylan active enzymes (xylanase 1—XNA1, xylosidase 2—XD2, and arabinofuranosidase—AFB) from Ustilago bevmoyces were cloned into 2-μm plasmids, p423 and p424 under the control of the GPD promoter. These plasmids were transformed into the S. cerevisiae YXS3 Δhis3 strain, which has the xylose consumption pathway genes from S. stipitis integrated. Before applying the evolutionary approach, the recombinant strains grew slowly on xylan as a sole carbon source, producing 0.26 ± 0.008 g L\(^{-1}\) ethanol from YPXN (YP supplemented with 20 g L\(^{-1}\) xylan) after 5 days of cultivation. To improve its ability to assimilate xylan, serial-subcultures in the xylan medium were used over 3 weeks. After selecting clones with improved traits, the evolved strain was able to produce 23% more ethanol in complex media (YPXN, 2% xylan), 0.32 ± 0.028 g L\(^{-1}\). These results demonstrate the capacity to use whole-cell adaptive evolution to improve xylan metabolism by the cell.
The often-emphasized advantage of the xylose isomerase pathway in comparison with the oxidoreductase pathway was considered by Mert et al. (2016). In the earlier studies (Katahira et al., 2004; Fujii et al., 2011; Sun et al., 2012), engineered *S. cerevisiae* strains with XR/XDH were modified by the introduction of xylanolytic enzymes. Although the authors observed ethanol production from XOS, large amounts of xylose remained in the fermentation broth, probably resulting from a redox imbalance and/or inefficient xylose uptake; this, in turn, can inhibit β-xylanolase activity (Fujii et al., 2011; Peng et al., 2017; Niu et al., 2019). However, early attempts to express xylanases in engineered *S. cerevisiae* strains harboring XI had failed to produce ethanol from beechwood xylan (5%) as the sole carbohydrate source under aerobic growth over 28 days (Mert et al., 2016). The recombinant strain, Y294 [YMXI], which carries *T. reesei* endoxylanase (XYNII), *A. niger* β-xylanolase (*xlnD*), *S. stipitis* xylulokinase (*xyl3*), and the codon-optimized xylose isomerase encoding gene (*xylA*) from *Bacteroides thetaiotaomicron* was able to break down xylan into trisaccharide, disaccharides, and monosaccharides. However, the growth rate was low probably due to the low consumption of xylose. The small amounts of xylose consumed supported cell biomass synthesis only; ethanol, xylitol, glycerol, and acetic acid production were negligible. It is worth noting that when this recombinant strain Y294 [YMXI] was cultivated under similar conditions but using xylose (2%) as the sole carbon source, higher biomass was obtained, and larger amounts of xylose were consumed in a lower cultivation time. Moreover, xylitol production was also observed (Mert et al., 2016). Unfortunately, whether the expressed xylanolytic enzymes were secreted or expressed intracellularly in the Y294 [YMXI] strain is not clear. It is important to mention that in previous studies published by this research group, using a *S. cerevisiae* expressing an endoxylanase encoding gene (*xyn2*) and a xylanolase encoding gene (*xlnD*), enzyme activities were detected in the culture supernatant (La Grange et al., 2001), suggesting that these enzymes were secreted by the strain.

Recombinant gene expression can promote a nonspecific metabolic burden which reduces the maximum specific growth rate and production yield of the host, as previously observed (Görgens et al., 2001). In this research, *T. reesei* xylanase II (XYN2) was expressed in two recombinant *S. cerevisiae* strains, Y294 [PGK1−XYN] and Y294 [ADH2−XYN], using two 2-μm yeast plasmids under the control of either the yeast glycolytic phosphoglycerate kinase (*PGK1*) or alcohol dehydrogenase II (*ADH2*) promoters, respectively. *ADH2* is a strong promoter inducible in the absence or at low concentrations of glucose, while *PGK1* is a constitutive promoter. However, no significant difference was observed for XYN2 expression by Y294 [ADH2−XYN] and Y294 [PGK1−XYN] strains. After 80 h of cultivation in a defined medium (Verduyn et al., 1992) containing 20 g L⁻¹ glucose, specific xylanase production levels were 3.2 and 2.6 mg (g biomass)⁻¹, respectively. The fermentation parameters of Y294 [PGK1−XYN] and Y294 [ADH2−XYN] were compared with those of the reference strains. In all Y294 [PGK1−XYN] and Y294 [ADH2−XYN] cultivations, a reduction in yeast biomass, ethanol, and glycerol yields were observed as well as specific consumption and production rates of glucose and ethanol, compared with the reference strains. Therefore, the expression of XYN2 from either *PGK1* or *ADH2* promoters resulted in a significant metabolic burden on the host metabolism.

These findings might explain the results obtained by Mert et al. (2016), who found lower biomass production in the engineered strain Y294 [YMXI] during cultivation on xylan than in xylose as sole carbohydrate source. It is likely that the metabolic burden associated with the expression of xylanolytic enzymes impacted xylose isomerase activity. Unfortunately, xylanase activity assays during cultivation on xylan were not reported (Mert et al., 2016). The influence of *ADH2* and *PGK1* on xylanase expression was also examined by (Nuyens et al., 2001). They also used two 2-μm yeast plasmids named pFN3 and pFN4 to insert endoxylanase (xynA) of *B. pumilus* PLS into *S. cerevisiae* Y294 strain under the control of these two different promoters. The two engineered yeast strains did not exhibit any xylanase activities until the gene encoding uracil phosphoribosyl transferase (*FUR1*) was disrupted. This step ensured auto-selection of the *URA3*-bearing expression plasmid in a rich growth medium since mutants by *FUR1* disruption allow the growth of the recombinant yeasts in a complex medium without the risk of losing the plasmid (La Grange et al., 1996). However, unlike the work of (Görgens et al., 2001), Y291 [pFN3 fur1:LEU2], in which the xylanase was under the control of the *ADH2* promoter, exhibited better xylanase activity (and presumably, expression) in the culture supernatant than Y291 [pFN4 fur1:LEU2], specifically 8.5 nkat ml⁻¹ and 4.5 nkat ml⁻¹, respectively.

Recently Niu et al. (2019), reported ethanol production from an efficient xylose-utilizing strain, BSPX042, expressing a xylose isomerase gene derived from a bovine rumen metagenomic study (Ru-xyla1), cloned in an episomal plasmid (pJXIH-PC, *URA3* as a select marker) carrying the β-xylanolase from *Penicillium oxalicum* (xyl3A) and the signal peptide fragment *INU* from *Kluyveromyces* sp. The recombinant strain, BSGIBX, cultivated in a selective synthetic complete medium supplemented with 20 g L⁻¹ XOS, immediately converted XOS into xylobiose and xylotriose after inoculation. The highest ethanol concentration, approximately 4.37 g L⁻¹, was reached at 36 h. When the XOS were pretreated with xylanase, the ethanol concentration reached approximately 9 g L⁻¹. Another important study involving the use of the XI pathway and xylanases is the work of (Tabañag et al., 2018). They expressed five different hemicellulases: endoxylanase (*XYNII*), β-xylanolase (*Bsl1*), acetyxyylan esterase (*Axel*), α-D-glucuronidase (*Glrl*) and α-L-arabinofuranosidase (*Ablf*), all from *T. reesei*, bound to the cell surface of a XI-expressing *S. cerevisiae* strain. Since hemicellulose is a complex structure that requires a consortium of enzymes to break it down completely, the authors explored accessory enzymes to make the main–chain more accessible to main–chain cleaving hemicellulases. The recombinant strain grew on xylan substrates as their sole carbon source and achieved an ethanol titer of 0.96 g L⁻¹ after 160 h of cultivation.

In the context of a lignocellulosic biorefinery, in order to make full use of cellulose and hemicellulose to produce ethanol, Lee et al. (2007) investigated constitutive co-expression of
endoxyylanase (xynA) from Bacillus spp. and endoglucanase (egl6) from Trichoderma spp. in S. cerevisiae SEY2102 strain. The expression levels of endoxylanase and endoglucanase were investigated during aerobic cultivation on YPD medium. Although fermentative parameters were not investigated, 5.6 U mL\(^{-1}\) of endoxylanase was secreted into the extracellular medium, and 1.96 U mL\(^{-1}\) was intracellular after 48 h cultivation. However, these findings are still far from achieving the goal of ethanol bioconversion from cellulosic biomass. The most promising strategy for converting cellulosic biomass to ethanol in yeast is certainly the concerted heterologous expression of all main types of hemicellulases and cellulases enzymes to maximize their synergies and improve ethanol production (Nevoigt, 2008).

Similarly to the previous report, Saitoh et al. (2011) have reported the expression of β-xilosidase and β-glucosidase from T. reesei on the yeast cell surface based on α-agglutinin engineering system, obtaining the engineered industrial S. cerevisiae strain OC2-AXYL2-ABGL2-Xyl2 which also contains the oxidoreductase pathway for xylose consumption. The highest ethanol concentration, 12.5 g L\(^{-1}\), was observed after 48 h in YPKX medium (40 g L\(^{-1}\) KC-flock and 40 g L\(^{-1}\) xylan from Birchwood) containing 30 G L\(^{-1}\) cellulose. The ethanol yield was 0.52 g (g sugar consumed)\(^{-1}\). Another example of the coexpression in S. cerevisiae of cellulase and hemicellulase enzymes, including an endoxylanase, xylsidase, and glucosidase was reported by (Sakamoto et al., 2012). These authors expressed endoxylanase from T. reesei, β-xilosidase from Aspergillus oryzae, and β-glucosidase from A. aculeatus anchored on the surface cell of the laboratory xylose-assimilating S. cerevisiae strain MN8140/XBX. Therefore, the recombinant strain, MN81/XBXX, expressed XR and XDH from S. stipitis, xylulokinase from S. cerevisiae, in addition to xylanases, and cellulase enzymes. The strain was reported to ferment cellulose and hemicellulose giving a high ethanol yield, 0.32 g g\(^{-1}\), and concentration of 8.2 g L\(^{-1}\) after 72 h, from rice straw under oxygen-limited conditions and initial cell concentration of 100 g of wet cells L\(^{-1}\). When 0.2 g L\(^{-1}\) of a commercial hemicellulase was added at the medium, the recombinant strain reached 10.3 g L\(^{-1}\) ethanol after 72 h, and the ethanol yield was 0.41 g g\(^{-1}\). The addition of the commercial hemicellulase allows complete hydrolysis of xylolbiose after 72 h of fermentation, which in turn increased xylolose content in the medium. Although the depletion of all xylolbiose, after 72 h of fermentation 2.2 g L\(^{-1}\) xylose remained in the fermentation medium (Sakamoto et al., 2012).

Lastly, Xiao et al. (2019) also reported the co-expression of both cellulase and xylanase enzymes in S. cerevisiae (unfortunately, the authors did not specify what cellulase or xylanase was used in their work). As reported herein, there is a synergistic action between cellulase and xylanase during lignocellulosic hydrolysate (Zhang et al., 2012; Wang et al., 2018; Xiao et al., 2019). In the presence of such synergies, pretreated lignocellulosic substrate degradation is more efficient because, since XOS can inhibit cellulase activity, the co-expression strategy would reduce the cellulase activity inhibitors (Zhang et al., 2012; Wang et al., 2018). Inside the concept of synergistic effect, it is worth noting that β-glucosidase did not affect xylanase activity as demonstrated by Chen et al. (2019b). The recombinant strains of Xiao et al. (2019), INVSc1-CBH-CA and INVSc1-CBH-TS, were cultivated using partly delignified corn stover (PDCS) producing 1.66 g L\(^{-1}\) and 1.90 g L\(^{-1}\) of ethanol after 120 h cultivation, respectively. This was approximately 4 times higher than the control (a strain that expressed a single cellulase or xylanase). Although the ethanol production did not exceed that of other published works (Katahira et al., 2004; Fujii et al., 2011; Niu et al., 2019) the effective synergistic effect of those enzymes could improve the saccharification of lignocellulose and increase the ethanol yield during fermentation by S. cerevisiae.

It is important to note that most of the investigations using engineered xylan-consuming S. cerevisiae cells have been carried out using laboratory strains, except by the work of (Saitoh et al., 2011). However, based on the industrial conditions for ethanol production, i.e., lignocellulosic inhibitors (Almeida et al., 2007), high osmolarity, and low pH, industrial host backgrounds would present more advantages as compared to laboratory strains (Della-Bianca and Gombert, 2013; Cola et al., 2020).

**ETHANOL PRODUCTION FROM INTRACELLULAR XOS UTILIZATION IN S. CEREVISIAE STRAINS**

Economic bioethanol production from lignocellulose requires complete and rapid conversion of both cellulose and hemicellulose on an industrial scale (Li et al., 2015; Mert et al., 2016). This generally includes the pretreatment of lignocellulosic biomass to increase enzyme accessibility, which improves the amount of fermentable sugars from the enzymatic digestion for biomass-to-bioethanol microbial conversion, (Palmqvist and Hahn-Hägerdal, 2000a; Katahira et al., 2004; Lynd et al., 2008). Ultimately, engineered S. cerevisiae expressing XOS-transporters and producing active xylanolytic enzymes for the intracellular depolymerization of XOS to xylose are important for reducing the xylanases inhibition by its end products and for tackling the issues of microbial contamination in industrial conditions, as well as for taking full advantage of all of the sugars in lignocellulosic hydrolysate (Fujii et al., 2011; Niu et al., 2019). There is a large amount of data on the expression of xylanolytic enzymes in S. cerevisiae strains, however, few of them report the expression of intracellular XOS hydrolysis system in this yeast (Li et al., 2015). Besides the reduction of xylanases inhibition by its end products, XOS internalization represents an additional advantage over extracellular hydrolysis. The engineered S. cerevisiae would grow faster than other contaminant microorganisms in the fermentation tank.

Neither XOS transporters in S. cerevisiae nor expression of heterologous XOS-transporters expression have been reported in the works mentioned above. Li et al. (2015) reported the first engineered S. cerevisiae strain able to consume XOS intracellularly following uptake by an oligosaccharide-transporter. The recombinant strain expressed two β-xilosidasises, GH43-2, and GH23-7, and one transporter, CDT-2,
from *Neurospora crassa* as well as XR/XDH from *S. stipitis* to ensure the internal breakdown of XOS into ethanol (SR8U) (Kim et al., 2013b; Li et al., 2015). The expression of both β-xylosidases was essential for converting XOS into xylose as the XR acted as an XOS reductase, producing xylosyl-xylitol as a potential dead-end product. Although GH43-7 had weak β-xylosidase activity, it rapidly hydrolyzed xylosyl-xylitol into xylose and xylitol. Anaerobic fermentation with this strain, expressing CDT-2, GH43-2, and GH43-7 in an optimized minimum medium (oMM) containing 4% xylose and 3% XOS, produced more than 30 g L\(^{-1}\) of ethanol in 72 h of cultivation, after supplying an additional 50 g L\(^{-1}\) xylose at hour 48 (Li et al., 2015). The authors also performed a co-fermentation of sucrose plus XOS with the strain SR8U carrying the plasmid pXD8.7. According to their report, the recombinant strain could increase 3 g L\(^{-1}\) of ethanol concentration comparing cultivations performed in oMM media containing approximately 60 g L\(^{-1}\) sucrose (control cultivation) and the media containing approximately 60 g L\(^{-1}\) sucrose and 10 g L\(^{-1}\) XOS of which 4.2 g L\(^{-1}\) represent xylobiose and 2.3 g L\(^{-1}\) xylotriose (Li et al., 2015).

Although Fujii et al. (2011) did not have express a specific XOS transporter, their recombinant strain could hydrolyze xylobiose and xylotriose to xylose intracellularly. They have reported an XOS-fermenting yeast strain, D-XSD/XKXDHXR, that was constructed by expression of intracellular *T. reesei* XKXDHXR, that was constructed by expression of their recombinant strain could hydrolyze the plasmid pXD8.7. According to their report, the recombinant strain could increase 3 g L\(^{-1}\) of ethanol concentration comparing cultivations performed in oMM media containing approximately 60 g L\(^{-1}\) sucrose (control cultivation) and the media containing approximately 60 g L\(^{-1}\) sucrose and 10 g L\(^{-1}\) XOS of which 4.2 g L\(^{-1}\) represent xylobiose and 2.3 g L\(^{-1}\) xylotriose (Li et al., 2015).

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As evident from Table 2, chromosomal integrative approaches have not been widely applied to construct the XOS-utilizing *S. cerevisiae* strain. Although laborious, this approach represents the only feasible strategy for engineering genetically stable yeast strains without a select marker, for industrial applications (Nevoigt, 2008; Fang et al., 2017). Fang et al. (2017) reported a promising approach to obtain yeast with a chromosomal multicopy expression of a *Bacillus* sp. xylanase (*xynHB*) in *S. cerevisiae* strain A13. rDNA-mediated integration was used in their work, providing stable expression over 1,011 generations of cultivation, and higher copy numbers of the target gene in the chromosome than from integrating plasmids, i.e., 13.64 copies of *xynHB* gene were found in the A13 genome. Yeast genome contains around 100 rDNA repetitive units which provide ideal homologous recombination sites for the target gene. It is worth noting that gene stability is only observed when the integrated plasmid is smaller than the size of the rDNA unit (9.1 kb) (Lopes et al., 1996). The A13 strain lacked the enzymes required to form xylose from xylose, therefore ethanol production was not the goal of Fang’s research.

### CONCLUDING REMARKS AND PROSPECTS

Although xylan-degrading enzyme systems have been studied extensively, there are much more missing points to connect than cellulose-degrading enzyme systems, probably because the structure of xylan is more complex and varies from plant to plant. However, the xylanolytic enzyme system deserves the same attention as the cellulolytic systems because their biotechnological potential is equally important. *S. cerevisiae* wild type strains are not suitable for producing bioethanol, even from a lignocellulosic hydrolysate with minimized production of inhibitors and high concentrations of hemicellulose/cellulose-derived oligosaccharides. A xylan-oligosaccharide-assimilating pathway has been demonstrated to be effective to generate *S. cerevisiae* strains able to convert polysaccharides into monomers. The effective intracellular hydrolysis of XOS has been demonstrated, however, the development of a strain capable of transporting large molecules of XOS is a crucial challenge. Screening for non-glucose specific transporters, such as xylose and xyl-oligomers specific transporters, and intracellular endoxylanases might advance strain improvement for efficient biomass conversion.

Significant improvements towards ethanol production from hemicellulose have been achieved in recent years, as the synergistic effect of overexpressing a combination of β-xylosidases, xylanases, and cellulases have been established (La Grange et al., 2000; Chen et al., 2019b; Xiao et al., 2019); however, there are potential limitations for efficient ethanol production from xylan by engineered *S. cerevisiae*, for example, β-Xylosidases with lower inhibition by products are needed for future *S. cerevisiae* engineering in order to achieve the complete conversion of xylobiose or xylotriose into xylose (Niu et al., 2019). In addition, it needs to be recognized that xylan in hemicellulose is typically branched and decorated, requiring accessory enzymes for their removal. Nevertheless, improvements in the technology to engineer and evolve *S. cerevisiae*, together with our current state of knowledge suggest that there is a high potential for the application of xylanolytic enzymes to obtain mono- and oligosaccharides from pretreated lignocellulose followed by fermentation into ethanol, since the cost of such sugars has historically been far too high to attract industrial interest.

Concerning the actual large-scale bioethanol production process from sugar- and starch-containing feedstock, hemicellulose-derived bioethanol is still in its infancy due to low ethanol yield achieved by the abovementioned engineered *S. cerevisiae* strains. However, it is worth mentioning that lignocellulosic hydrolysate contains a mix of carbon sources, such as cello-oligosaccharides, XOS, and monomers as glucose, and xylose. Naturally, *S. cerevisiae* can consume some mono- and disaccharides, such as D-glucose, D-galactose, D-fructose,
D-mannose, maltose, sucrose, and trehalose (Lagunas 1993), which do not represent the totality of the sugar derived from lignocellulosic hydrolysate. It is well known in the scientific literature that economically feasible bioethanol production might include the use of all sugars available in the lignocellulosic biomass. Thereby, despite the poor fermentation performance of XOS-utilizing S. cerevisiae strains as compared to glucose/sucrose fermentations, the simultaneous co-fermentation of lignocellulosic-derived sugars may result in higher ethanol titers and will maximize the use of the carbon sources available in lignocellulosic feedstock.

AUTHOR CONTRIBUTIONS

This review arose out of the PhD work of DPP, supervised by TOB, in which she spent time working in the laboratories of DJL.

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