Ashbya gossypii is a filamentous fungus that naturally overproduces riboflavin. Indeed, engineered strains are currently used for the industrial production of riboflavin, replacing the chemical synthesis processes formerly used. The utilization of A. gossypii for biotechnological applications affords significant advantages that involve low-cost media use and cheap downstream processing for some applications. Although A. gossypii cannot be considered a bona fide oleaginous microorganism, the accumulation of lipid droplets within hyphae has been described. In view of the genomic and molecular tools available for its manipulation, the metabolism of A. gossypii was engineered aiming to increase total lipid accumulation. Blocking the β-oxidation pathway through the knock-out of the AgPOX1 gene was sufficient to obtain strains with high lipid yields, comparable to those of the best oleaginous microorganisms. Thus, the poxΔ strain of A. gossypii constitutes a novel promising tool for the production of microbial oils in forthcoming modified A. gossypii strains.

Keywords: Ashbya gossypii, metabolic engineering, microbial oil, biolipids, single-cell oil, biodiesel

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Ashbya Gossypii as a Novel Organism for Microbial Oil Production

Our society is strongly dependent on crude oil for the production of fuels, fine chemicals, and many other commodities. However, fossil oil supplies are limited, and their use has negative environmental consequences. Thus, in recent years new technologies have been developed for the production of both fuels and chemicals from renewable sources, such as plant oils, animal fats or microbial oils.

Microbial oils (also called single-cell oils, SCOs) have several advantages over other oil sources: i) the fermentation process of microorganisms is independent of season and climate; ii) many industrial by-products can be used as fermentation substrates; and iii) microorganisms can be efficiently modified via metabolic engineering approaches, hence allowing the optimization of the amount and properties of the SCOs.

In light of the above, biotechnological research has focused on oleaginous microorganisms, which can accumulate at least 20% of their cell dry-weight (CDW) as lipid content. Additionally, some oleaginous microorganisms can reach higher levels of lipid, accumulating up to 80% of their CDW when cultured in oil-containing media. However, the biotechnological optimization of some oleaginous microorganisms for industrial applications is difficult owing to the lack of appropriate metabolic engineering tools. Accordingly, a potential alternative is the manipulation of industrial microorganisms that can be engineered to afford lipid yields comparable to oleaginous microorganisms. In this sense, some model organisms, such as Escherichia coli and Saccharomyces cerevisiae, have been engineered to overproduce lipids.

Ashbya gossypii is a filamentous hemiascomycete that is extensively used for the industrial production of riboflavin. Indeed, the use of A. gossypii in industry is considered a paradigm of sustainable white biotechnology for the industrial overproduction of riboflavin and other vitamins.

Importantly, a large number of genomic, bioinformatic and biotechnological tools are available for A. gossypii, thus allowing the development of systems metabolic engineering approaches to industrial applications of the fungus. In addition, the use of A. gossypii for
microbial fermentation has other biotechnological advantages such as the ability to grow both in some industrial by-products and low-cost oils, the partial autolysis of hyphae at late growth phases, and the harvesting of mycelia by simple filtration. Moreover, A. gossypii can be grown by pellet fermentation, which reduces broth viscosity and improves mixing and mass transfer, saving both costs and energy.

A. gossypii has not been considered an oleaginous fungus because it is not able to accumulate more than 20% of its cell dry weight in the form of fatty acids. However, as previously mentioned the fungus can be manipulated to overproduce different metabolites. Thus, in our recent paper entitled “Strain Design of Ashbya gossypii for Single-Cell Oil Production” we described the generation of A. gossypii strains that were engineered to accumulate up to 70% of their CDW as lipids in glucose+oil-containing media (1% glucose + 2% oleic acid). This represents a novel promising microbial platform for the production of single-cell oil.

**Lipid accumulation in A. Gossypii**

The concept of lipid accumulation refers to a metabolic imbalance between lipid biosynthesis and degradation. The biosynthesis of lipids mainly depends on a continuous supply of cytosolic acetyl-CoA, which is the substrate for the acetyl-CoA carboxylase (ACC) to form malonyl-CoA. Next, the fatty acid synthase (FAS) multienzymatic complex catalyzes the elongation of the acyl-CoA chain by condensing malonyl-CoA molecules and acetyl-CoA (Fig. 1). In this respect, oleaginous microorganisms are equipped with the enzyme ATP-citrate lyase, which converts citrate into cytosolic acetyl-CoA. Additionally, the biosynthesis of lipids also relies on a sufficient supply of NADPH as the essential reducing power for fatty acid biosynthesis. In contrast to lipidogenesis, β-oxidation is the main catabolic pathway of fatty acids and, consequently, its activity competes with the accumulation of lipids. Beta-oxidation comprises a 4-step oxidative cycle that removes 2 carbons (an acetyl-CoA molecule) from the acyl-CoA chain in each cycle.

The wild-type strain of A. gossypii is able to accumulate about 5% of its CDW as lipids in glucose-rich media (8% glucose), while it can accumulate almost 20% of its CDW as lipids in glucose+oil-containing media (1% glucose + 2% oleic acid). Lipid accumulation can be optimized both by increasing the availability of lipid precursor and by blocking the competing pathways of lipid accumulation.

*A. gossypii* lacks the enzyme ATP-citrate lyase, which is a hallmark of oleaginous microorganisms. In our work, *A. gossypii* was endowed with ATP-citrate lyase (ACL) activity through the heterologous expression of the *ACL1* and *ACL2* genes from *Yarrowia lipolytica*, which code for the 2 subunits of the ACL enzyme. Expression of the *Y. lipolytica* ACL genes in the wild-type strain of *A. gossypii* induced a 30% increase in lipid accumulation in glucose-rich medium (8% glucose) after 3 days of culture. Moreover, the β-oxidation pathway was abolished in *A. gossypii* through the deletion of the *POX1* and *FOX1* genes, which control the first and second steps of β-oxidation, respectively. The deletion of both the *POX1* and *FOX1* genes in *A. gossypii* did not exert any effect on biomass production either in glucose-rich medium or in glucose+oil-containing medium; however, growth was absent in soybean oil medium (2% soybean oil) (Fig. 2), indicating that the β-oxidation pathway supports growth from culture media with oil as the only carbon source. Accordingly, oil-containing medium for the quantification of lipid accumulation in either the *pox1Δ* or *fox1Δ* strains of *A. gossypii* must also include a non-lipid carbon source in their formulations (typically 1% glucose). The elimination of β-oxidation in *A. gossypii* resulted in a 2-fold increase in lipid accumulation both in glucose-rich and glucose+oil-containing media after 3 days of culture. However, the expression of the *Y. lipolytica* ACL enzyme in a *pox1Δ* strain did not show any increase in the lipid yield.

Regarding the levels of acetyl-CoA in the engineered strains, the expression of the *ACL* genes in the wild-type induced a significant increase in the intracellular acetyl-CoA pool (2-fold more acetyl-CoA). Interestingly, we found that the *pox1Δ* strain had levels of acetyl-CoA 4-times higher than those of the wild-type strain and, unexpectedly, the *pox1Δ* strain had higher levels of acetyl-CoA (approximately 2.5-fold higher) than the *ACL* overexpressing strain. In addition, the *pox1Δ* strain showed a lower increase in acetyl-CoA levels (50% more acetyl-CoA) after combined expression of the *ACL* genes from *Y. lipolytica*. Accordingly, lipid accumulation was not correlated with the

![Figure 1. Simplified diagram of the lipid metabolism in *A. gossypii*. The ACL (ATP:citrate lyase) activity (gray) is not present in *A. gossypii*. ACS, acetyl-CoA synthase; ACC, acetyl-CoA carboxylase; FAS, fatty acid synthase; Pox1, fatty-acyl coenzyme A oxidase; Fox2, 3-hydroxyacyl-CoA dehydrogenase and enoyl-CoA hydratase; Fox3, 3-ketoacyl-CoA thiolase. The elongation system comprises the elongases ELO624 and ELO586 which elongate C18:0 fatty acids up to C26:0 (C > 18). The desaturation system is composed by 2 desaturases (Δ9 and DES589). Saturated fatty acids (Cn:0) can be desaturated by Δ9 to form Cn:1 fatty acids (where n is the number of carbons of the acyl chain). The Δ12 DES589 can further desaturate the Cn:1 fatty acids to form Cn:2 (mainly C18:2, linoleic acid). Dashed arrows indicate multistep pathways.](image-url)
increase in acetyl-CoA levels in the different strains. Furthermore, the \textit{pox1}\textDelta strain expressing ATP-citrate lyase did not show any increase in the total lipid content, indicating that an additional metabolic bottleneck exists after the limitation in acetyl-CoA synthesis has been overcome. The block in the \(\beta\)-oxidation pathway in the \textit{pox1}\textDelta strain of \textit{A. gossypii} was sufficient to trigger a huge increase in fatty acid accumulation in glucose+oil-containing medium. After 7 d of culture in glucose+oil-containing medium, the \textit{pox1}\textDelta strain was able to accumulate up to 70\% of the cell dry weight as lipid content (Fig. 3A). Indeed, the \textit{pox1}\textDelta strain showed a characteristic phenotype of its hyphae when growing on glucose + oil-containing media due to the high lipid accumulation (Fig. 3B). In contrast, the \textit{pox1}\textDelta strain showed a lower total fatty acid yield (about 60\% of CDW) when it was engineered to overexpress both the \textit{ACLI} and \textit{ACL2} genes from \textit{Y. lipolytica}.\textsuperscript{14} This effect was inversely correlated with higher levels of acetyl-CoA in that strain, again supporting the idea that acetyl-CoA does not limit lipid accumulation under those conditions.

These results suggest that substrate limitation for lipid synthesis in \textit{A. gossypii} can be partially prevented by the expression of ATP-citrate lyase. However, once acetyl-CoA levels are no longer a limiting factor for \textit{de novo} lipid synthesis, other mechanisms seem to affect lipid production. The increased pool of intracellular acyl-CoA might be responsible for these limitations since it has been demonstrated that high levels of acyl-CoA can inhibit fatty acid synthesis in other microorganisms.\textsuperscript{17,18}

Accordingly, the enzymes involved in the esterification of the acyl-CoA to sterols or triacylglycerol might be good candidates to further manipulate the oleaginous nature of \textit{A. gossypii}. Also, NADPH levels might not be sufficient to provide enough reducing power for an increased biosynthesis of fatty acids. Thus, a combined strategy aimed at increasing both the levels of acetyl-CoA and NADPH has been used for the production of fatty acid ethyl esters and polyhydroxybutyrate.\textsuperscript{19-21}

Many culture medium formulations for microbial fermentations are based on waste/low-cost sugars and it is therefore important to improve lipid accumulation from sugar-based media.\textsuperscript{22} In this sense, the \textit{pox1}\textDelta strains generated in our work were able to produce up to 9\% of their CDW as lipids in rich glucose-rich medium (Fig. 3). Thus, although the total oil yield in \textit{A. gossypii} growing in glucose-rich media is still low, it is significantly higher than that seen for some oleaginous yeasts such as engineered \textit{Y. lipolytica}, which can accumulate up to 4.8 +/- 0.4\% of its CDW as fatty acids in a similar rich medium (YPD).\textsuperscript{23} Nonetheless, \textit{de novo} lipid biosynthesis must be increased by optimized medium formulations and/or by additional metabolic engineering manipulations. For example, a wild-type strain of \textit{Y. lipolytica} is able to attain 16.3\% of its cell dry weight CDW as lipids under optimized conditions in glucose-based media and up to 88\% of lipid content after extensive metabolic engineering.\textsuperscript{24} Similarly, the oleaginous yeast \textit{Rhodosporidium toruloides} accumulates up to 9.8\% and 63\% of its CDW as fatty acids in glucose-based media and in nitrogen-limited media respectively.\textsuperscript{25} These comparisons suggest that oil production in \textit{A. gossypii} from sugar-based media need...
to be further improved both by medium optimization (i.e. nitrogen-limited conditions) and by systems metabolic engineering.

Regarding the use of oil-containing media, the increased rate of lipid accumulation in such media suggests that the incorporation of extracellular lipids is a very active process in \textit{A. gossypii}. This is worth noting within the context of the industrial process called oil biotransformation, which is the fermentative conversion of a cheap oil into a high-value oil. This process relies on cells’ ability to modify the extracellular fatty acids that have been incorporated into the cytosol. For example, the activity of fatty acid elongases can tune the length of fatty acids, while desaturases can catalyze the formation of double bonds within the acyl chain. The activity of fatty acid desaturases can catalyze the formation of double bonds within the acyl chain. Furthermore, the activity of fatty acid elongases can incorporate into the cytosol. For example, the extracellular fatty acids that have been processed rely on cells’ ability to modify the extracellular fatty acids of industrial relevance.\textsuperscript{14,26} Additionally, metabolic engineering has also been applied to both the elongase and desaturase systems with a view to generating novel \textit{A. gossypii} strains that are able to accumulate polyunsaturated fatty acids, which are undesired features in biodiesel blends.\textsuperscript{26}

Taking the above observations together, we propose \textit{A. gossypii} as a novel and promising candidate for biolipid production that can offer an economically feasible alternative to some oleaginous yeasts. It has been demonstrated that modified strains of \textit{A. gossypii} are able to accumulate high amounts of lipids and, additionally, it is possible to adapt the lipid profiles of such strains for them to become enriched in fatty acids of industrial relevance.\textsuperscript{14,26} Nonetheless, more research is being carried out to increase lipid production from glucose-based media as well as to optimize the biotransformation of low-cost oils into high-value oils.

Disclosure of Potential Conflicts of Interest

No potential conflicts of interest were disclosed.

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