Effect of Environmental Stress Factors and Recycling on the Lipid Composition of Brewer’s Yeast Mitochondria

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Abstract
The aim of this study was to investigate alterations in the content and composition of mitochondrial lipids of brewer’s yeast, which occur during brewing and repetitive recycling. The bottom-fermenting brewer’s yeast of Saccharomyces cerevisiae species obtained from industrial beer production was used throughout the study. The first three generations of recycled yeast were analysed. Differences between the generations were more pronounced in the phospholipid and neutral lipid composition than in fatty acid composition. Squalene was present in all generations in high concentrations. The results give insight into the response of yeast cells to stress factors and recycling.

Keywords
Brewer’s yeast, recycling, mitochondria, plasma membrane, lipids, squalene

1 Introduction
Water, malt, yeast, and hops are the basic raw materials for the production of beer. Yeast is responsible for the production of alcohol and carbon dioxide, but also for the production of fermentation by-products (higher alcohols, acids and esters), which are responsible for the bouquet of beer.

Most organisms are exposed to constant changes in their environment, and therefore their survival depends on the capacity to adapt to new conditions. This also applies to industrial yeasts, but in a much more pronounced way, since their cells are exposed simultaneously to different stressors during production processes. Among them, exposure to ethanol and oxygen-limited availability are of special importance for the bottom-fermenting brewer’s yeast. Modern brewing technology is based on wort fermentation with selected, aerobically propagated, pure yeast cultures. During aerobic growth, some yeasts (like those from Saccharomyces genus) in the second growth phase (called oxidative), spend ethanol produced from sugars in the first phase (called fermentative). Yeast culture used in one fermentation cycle, is usually reused in successive fermentation cycles. The number of cycles is theoretically unlimited, but in practice, it includes 8–10 fermentation cycles (often reduced to 3–5). The performance of brewer’s yeast declines with repeated use and the yeast is said to become “weak” and physiologically inactive. Recycled yeast is pitched in wort with limited oxygen concentration. In the case of the bottom-fermenting brewer’s yeast, characterized by high density, the aerobically propagated pure yeast culture undergoes anaerobic conditions during fermentations because the yeast biomass precipitates on the bottom of fermenter. Failure to provide sufficient molecular oxygen to the fermentation medium restricts yeast growth and viability, since the cells cannot produce unsaturated fatty acids (FAs) and sterols for membrane biosynthesis. Yet, brewing wort contains small portions of unsaturated FAs, which the yeast cells can use for membrane reconstruction.

Various physiological adaptations related to stress protection occur in yeasts, ranging from the alteration in membrane fluidity to the synthesis of detoxification enzymes. Membrane lipids play an essential role in the biophysical and biochemical characteristics of cell membranes; they determine permeability and fluidity, enable proper functioning of membrane-attached proteins, and participate in the cell signalling. The ability of yeast cells to alter sterol and phospholipid (PL) ratio, PL composition and location, as well as the degree of unsaturation of membrane lipids is an important factor in adaptation to environmental conditions.

The mitochondria are engaged in many cellular processes, one of the most important being oxidative degradation of nutrients for the production of energy. Since the cell growth and normal functioning depend on the generation of metabolic energy, well-developed mitochondria are of special importance for the cells. They are dynamic organelles whose functional integrity requires a coordinated supply of proteins and lipids adjusted to meet physiological and functional demands. Their lipid composition is of great importance since they contain a double membrane system, with a pronounced curvature of the inner membrane shaped by lipids.

Phosphatidycholine (PtdCho), phosphatidylethanolamine (PtdEtn) and phosphatidylinositol (PtdIns) are the main phospholipids in S. cerevisiae cells, including the mitochondria. Cardiolipin (CL), a typical mitochondrial phospholipid, is the key determinant in the maintenance of...
It is deposited at the Collection of Microorganisms of the University of Zadar. The strain used throughout the study was obtained from a local brewery.

2.2 Isolation of mitochondria

Crude mitochondria were isolated by differential centrifugation after enzymatic disruption of the cell wall using zymolase.24

2.3 Plasma membrane preparation

The plasma membranes were obtained by the method for the isolation of highly purified plasma membranes.25 For technical reasons, the first part of the method was modified, so that instead of mechanical disruption of the cell wall with glass beads, the enzymatic method with zymolase was applied, described in detail in our previous paper.23

2.4 Protein determination

Protein content of the plasma membrane was determined by the method of Lowry26 using bovine serum albumin as the standard.

2.5. Lipid extraction

Total lipids from the mitochondria and plasma membrane were extracted by Folch method.27

Chromatographic methods used for the lipid analysis are the standard ones for the determination of basic lipid composition and accepted by Saccharomyces cerevisiae – EUROFAN (European Function Analysis Network) 2 group.28

2.6 Phospholipid analysis

Total phospholipids (TPLs) in the lipid extract were determined spectrophotometrically as inorganic phosphorus. Samples of total lipid extracts were digested with concentrated nitric and sulfuric acid by gentle heating in a microwave oven for 20 min. After cooling, the solution was neutralized with solution of sodium hydroxide. The intensity of blue colour, developed by ascorbic acid, solution of ammonium heptamolybdate tetrahydrate and antimony potassium tartrate hemihydrate in sulfuric acid, was measured at 880 nm. The concentration of TPLs was calculated by multiplying mass concentration of phosphorus by 25 and expressed per mass of proteins. PL classes were determined by two-dimensional thin-layer chromatography on silica gel 60 plates, 20 × 20 cm, 0.2 mm. Chloroform/methanol/ammonia (in a volume ratio 15 : 7 : 1) was used as the first, and chloroform/acetone/methanol/acetic acid/
water (in a volume ratio 10 : 4 : 2 : 1) as the second solvent system. PLs were visualised by iodine staining, scraped off the plate, and quantified the same way as TPLs, using factor 27.1 for PtdIns, and 24.2 for other PLs, respectively.

2.7 Analysis of neutral lipids
Neutral lipids were separated by two-step TLC on silica gel 60 plates (200 × 100 × 0.2 mm). Lipid extracts and standards were applied by a sample applicator (Linomat V, Camag, Switzerland). The plates were developed by using light petroleum/ diethyl ether/ CH₂COOH (volume ratio 20 : 20 : 0.8) up to 1/3 of a plate as the first, and light petroleum/ diethyl ether (volume ratio 39.2 : 0.8) up to 2/3 of plate as the second solvent system. For the determination of components, the bands were visualised by post chromatographic derivatization. Plates were dipped for 4 s into developing reagent (0.63 g MnCl₂ · 4H₂O, 60 ml H₂O, 60 ml CH₃OH and 4 ml H₃SO₄, conc.), dried, and heated for 30 min at 100 °C. Quantification was carried out by densitometric scanning (GS-710 Calibrated Imaging Densitometer, Bio-Rad).

2.8 Fatty acid analysis
Fatty acid (FA) composition was determined by GC analysis of the corresponding methyl esters obtained by acid methanolation of lipid extracts with BF₃/methanol. GC analyses of FA methyl esters were carried out using an Auto System XL from Perkin-Elmer with flame-ionization detector (FID). An SP-2330 capillary column (Supelco, USA), 30 m × 0.32 mm × 0.2 µm and helium as carrier gas with split injection (100 : 1) were used. Hydrogen was obtained with a Claind hydrogen generator. The analyses were carried out in programmed temperature mode from 140 to 220 °C at 5 °C min⁻¹ and then isothermally for 25 min. The injector temperature was 300 °C and the detector 350 °C. For data acquisition, Chromatography Software from Perkin-Elmer Nelson (Turbochrom 4) was used. The results were expressed as mass fractions of individual FA in total FAs. The degree of unsaturation is expressed as unsaturation index (UI) which is calculated as follows:

\[
UI = w(\text{monoenic fatty acids}) + 2 w(\text{dienoic fatty acids}) + 3 w(\text{trienoic fatty acids})
\]

3. Results and discussion
Saccharomyces cerevisiae is used in beer production in the medium, ergosterol and squalene were the main

3.1 Phospholipids
Mass ratio of total PLs to proteins accounted for 0.020 mg mg⁻¹ in the mitochondria of the starter culture and 0.027, 0.031 and 0.066 mg mg⁻¹, respectively, in the first three recycled generations.

Phospholipid composition of the mitochondria was characterised by predominance of PtdCho, which was the main PL in the zero and first two recycled generations (ranging from 23–29 %) and the second one in the third generation, accounting for 20 %. In spite of anaerobic conditions, CL was present in relatively high concentrations (14–16 %), which is a characteristic of the well-developed mitochondria. PtdSer was present in the lowest concentrations, ranging from 5–7 %, which is in accordance with the literature since it is enriched in the PM. When the results of analysed yeast generations were plotted for individual PLs, regularity in the alteration became obvious: in the first generation, obtained after the first beer fermentation, the alteration was the greatest, followed by the alteration in the opposite direction in the second and third generations (Fig. 1).
Components. The results of the analysis are presented in Table 1.

Table 1 – Content of ergosterol and squalene in the mitochondria and plasma membrane of the zero and first three recycled generations of brewer’s yeast

<table>
<thead>
<tr>
<th>Yeast generation</th>
<th>Zero Nulla</th>
<th>1st</th>
<th>2nd</th>
<th>3rd</th>
</tr>
</thead>
<tbody>
<tr>
<td>mitochondria</td>
<td>ergosterol</td>
<td>16.2</td>
<td>23.0</td>
<td>24.8</td>
</tr>
<tr>
<td></td>
<td>squalene</td>
<td>4.7</td>
<td>7.4</td>
<td>7.6</td>
</tr>
<tr>
<td>plasma membrane</td>
<td>ergosterol</td>
<td>18.4</td>
<td>11.0</td>
<td>10.6</td>
</tr>
<tr>
<td></td>
<td>squalene</td>
<td>10.4</td>
<td>19.4</td>
<td>12.8</td>
</tr>
</tbody>
</table>

The results represent the mean value of double analysis of at least three independent experiments; ‘Čanadi Jurešić et al.\(^29\).

Rezultati predstavljaju srednju vrijednost dviju analiza najmanje tri nezavisne pokusa; ‘Čanadi Jurešić et al.\(^29\).

Ergosterol was present in a higher concentration (mass ratio of lipids to proteins 11–25 \(\mu g \text{mg}^{-1}\)) compared to squalene (2–8 \(\mu g \text{mg}^{-1}\)), but regarding the normal distribution of the lipids in the cells, squalene as a hydrocarbon should not be present in the membranous systems at all. We have previously determined high concentrations of squalene in the mitochondria and plasma membranes in a recycled generation of another industrial strain of brewer’s yeast,\(^20\) which we have explained as a way the yeast cells cope with anaerobic conditions. Probably, squalene is incorporated in the mid-plane of the lipid bilayer, thus decreasing membrane permeability, as was hypothesised for bacteria.\(^29\) Since the key step in sterol synthesis, the formation of squalene epoxide from squalene, occurs in aerobic conditions, the finding of squalene in the mitochondria of the zero generation (aerobically propagated) was quite unusual.\(^29\) Squalene was found in the plasma membranes of all analysed generations, as well, and its concentration was higher than in the mitochondria accounting for 10–19 \(\mu g \text{lipids/mg proteins}\) (Table 1). In the plasma membrane of all recycled generations, its concentration was markedly higher than the concentration of ergosterol. These results confirmed our hypothesis that squalene replaced ergosterol in the membranes of the bottom-fermenting brewer’s yeast in anaerobic and aerobic conditions, as well, as adaptation to the governing conditions during beer fermentation, which was achieved by careful strain selection during thousands of years.

3.3 Fatty acids

Several studies have reported on the fatty acid composition of brewer’s yeast along with its alterations under different conditions, mainly in relation to the ethanol concentration.\(^18\) The general conclusion for \(S. \text{ cerevisiae}\) is that the composition of unsaturated FAs is a significant determinant of ethanol tolerance, and that oleic acid is the most efficacious unsaturated FA in overcoming the toxic effect of ethanol.\(^31\) Published data concerning the mitochondria, are scarce. In our previous study of recycled yeast (the third generation) of another industrial strain of the bottom-fermenting brewer’s yeast, we have determined palmitic acid as by far the most represented, accounting for 43 % (and 41 % in the plasma membrane, respectively). The overall content of saturated acids was very high, accounting for 79 % and the C16/C18 ratio was 5.5.\(^20\) In the yeast analysed in this study, two acids were highly represented, palmitic and palmitoleic, accounting for 33–37 %, and 28–39 %, respectively (Table 2). Compared to the zero generation, the first recycled generation was characterised by a marked increase of palmitoleic acid mainly on the account of palmitic and oleic acid, which resulted in a marked increase in the C16/C18 ratio from 2.4 to 3.3. Similar to the plasma membranes, unsaturated fatty acids slightly predominated in all analysed generations, ranging from 53 % to 56 %, which was quite unusual for anaerobically grown brewer’s yeast.

Table 2 – Main fatty acids in the total mitochondrial lipids of the zero and first three recycled generations of the bottom-fermenting brewer’s yeast, and the main features of the composition (expressed as mass fraction of total identified fatty acids)

<table>
<thead>
<tr>
<th>Yeast generation</th>
<th>Zero Nulla</th>
<th>1st</th>
<th>2nd</th>
<th>3rd</th>
</tr>
</thead>
<tbody>
<tr>
<td>mass fraction /% maseni udjel /%</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>C16:0</td>
<td>37.2</td>
<td>33.7</td>
<td>35.9</td>
<td>33.2</td>
</tr>
<tr>
<td>C16:1</td>
<td>31.6</td>
<td>38.6</td>
<td>27.7</td>
<td>36.8</td>
</tr>
<tr>
<td>C18.0</td>
<td>6.9</td>
<td>7.8</td>
<td>9.1</td>
<td>8.5</td>
</tr>
<tr>
<td>C18:1</td>
<td>18.9</td>
<td>10.2</td>
<td>12.1</td>
<td>10.5</td>
</tr>
<tr>
<td>C18:2</td>
<td>3.2</td>
<td>3.5</td>
<td>10.1</td>
<td>7.4</td>
</tr>
<tr>
<td>saturated, SFAs</td>
<td>45.5</td>
<td>45.8</td>
<td>47.3</td>
<td>43.7</td>
</tr>
<tr>
<td>unsaturated, UFAs</td>
<td>54.5</td>
<td>54.2</td>
<td>52.7</td>
<td>56.3</td>
</tr>
<tr>
<td>U/FAs/SFAs</td>
<td>1.2</td>
<td>1.2</td>
<td>1.1</td>
<td>1.3</td>
</tr>
<tr>
<td>unsaturation index indeks nezasićenosti</td>
<td>0.59</td>
<td>0.58</td>
<td>0.62</td>
<td>0.65</td>
</tr>
<tr>
<td>C16</td>
<td>68.7</td>
<td>72.3</td>
<td>63.7</td>
<td>70.0</td>
</tr>
<tr>
<td>C18</td>
<td>29.0</td>
<td>21.7</td>
<td>32.1</td>
<td>27.0</td>
</tr>
<tr>
<td>C16/C18</td>
<td>2.4</td>
<td>3.3</td>
<td>2.0</td>
<td>2.6</td>
</tr>
<tr>
<td>C16:0/C16:1</td>
<td>1.2</td>
<td>0.9</td>
<td>1.3</td>
<td>0.9</td>
</tr>
<tr>
<td>C18:0/C18:1</td>
<td>0.4</td>
<td>0.7</td>
<td>0.8</td>
<td>0.8</td>
</tr>
</tbody>
</table>

The results represent the mean value of double analysis of at least three independent experiments; ‘Čanadi Jurešić et al.\(^29\).

Rezultati predstavljaju srednju vrijednost dviju analiza najmanje tri nezavisne pokusa; ‘Čanadi Jurešić et al.\(^29\).
recycled generations. In contrast to some other parameters of the lipid composition (e.g. phospholipid composition), the alterations of UFAs/SFAs were small and showed no trend, but the unsaturation index showed a marked increase in the second and third generations (Table 2). The FA analysis of individual PL classes showed that UFAs prevailed only in the main PLs, PtdCho and PtdEtn, of the first generation, while SFAs prevailed in the second generation and in other, less represented PLs (data not shown). These results suggest that neutral lipids were the pool of UFAs in recycled generations, which is in accordance with the findings of Sajbidor and Grego.

4 Conclusion
Alterations in the lipid composition of the mitochondria under unfavourable conditions – which come to the fore mostly during transition from aerobic propagation to anaerobic beer fermentation – are marked in all lipid classes. On the other hand, recycling has a strong effect on phospholipids and neutral lipids, but a very slight effect on the FA composition. Therefore, since unsaturation is generally considered as the most important factor in the yeast adaptation to ethanol, we may say that the recycled yeast had adapted well to the governing conditions during the production process. The changes that follow transition from propagation to fermentation may be considered as stress injury, and alterations in the first recycled generations as stress response and indicator of good adaptation of the yeast to new, unfavourable conditions.

ACKNOWLEDGEMENTS
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List of abbreviations
Popis kratica

CL – cardiolipin
FA – fatty acid
FAME – fatty acid methyl ester
PL – phospholipid
PtdOH – phosphatidic acid
PtdCho – phosphatidylcholine
PtdEtn – phosphatidylethanolamine
PtdIns – phosphatidylinositol
PM – plasma membrane
SE – steryl ester
TAG – triacylglycerol
TL – total lipids
TPL – total phospholipids
UI – unsaturation index, %

References
Literatura

SAŽETAK
Utjecaj okolišnih čimbenika stresa i recikliranja na lipidni sastav mitohondrija pivskog kvasca

Gordana Čanadi Jurešić, Suzana Popović, Josip Škara, Marin Glad i Branka Blagović

Cilj istraživanja bio je odrediti promjene u sadržaju i sastavu lipida mitohondrija pivskog kvasca do kojih dolazi za vrijeme vrenja i recikliranja biomase. U radu je upotrijebljen pivski kvasac donjeg vrenja vrste Saccharomyces cerevisiae dobiven u industrijskom procesu. Analizirane su prve tri generacije recikliranog kvasca. Razlike između pojedinih generacija bile su izražene u sastavu fosfolipida i neutralnih lipida nego u sastavu masnih kiselina. Skvalen je u svim generacijama bio prisutan u relativno visokoj koncentraciji. Rezultati istraživanja pružaju uvid u odgovor stanica na okolišne čimbenike stresa i recikliranja.

Ključne riječi
Pivski kvasac, recikliranje, mitohondriji, citoplazmatska membrana, lipidi, skvalen

Izvorni znanstveni rad