IMPACT OF BIOREACTOR SCALE ON LOVASTATIN BIOSYNTHESIS
BY *ASPERGILLUS TERREUS* ATCC 20542 IN A BATCH CULTURE

Marta Pawlak*, Marcin Bizukojć, Stanisław Ledakowicz

Technical University of Lodz, Faculty of Process and Environmental Engineering, Department of Bioprocess Engineering, ul. Wólczańska 213, 90-924 Łódź, Poland

Biosynthesis of lovastatin (a polyketide metabolite of *Aspergillus terreus*) in bioreactors of different working volume was studied to indicate how the change of scale of the process influences the formation of this metabolite. The experiments conducted in shake flasks of 150 ml working volume allowed to obtain lovastatin titres at the level of 87.5 mg LOV l⁻¹, when two carbon sources, namely lactose and glycerol were used. The application of the same components in a large stirred-tank bioreactor of 5.3-litre working volume caused a decrease of lovastatin production by 87% compared to the shake flask culture. The deficiency of nitrogen in this bioreactor did not favour the formation of lovastatin, in contrast to the small bioreactor of 1.95-litre working volume, in which lovastatin titres comparable to those in the shake flasks could be achieved, when organic nitrogen concentration was two-fold decreased. When the control of pH and/or pO₂ was used simultaneously, an increase in lovastatin production was observed in the bioreactors. However, these results were still slightly lower than lovastatin titres obtained in the shake flasks.

**Keywords:** lovastatin, *Aspergillus terreus* ATCC 20542, shake flask, stirred tank bioreactor, batch process

1. INTRODUCTION

Lovastatin (C₂₄H₃₆O₅) is a secondary polyketide metabolite produced by such filamentous fungi as *Aspergillus terreus* and *Monascus ruber*. This compound is a natural statin, commonly used as a sovereign antihypercholesterolemia drug. The effect of lovastatin is revealed through the inhibition of (S)-3-hydroxymethylglutaryl-CoA (3-HMG-CoA) reductase, which converts 3-HMG-CoA to mevalonate in the pathway of cholesterol biosynthesis in human cells. The basic strain of *A. terreus* capable of the formation of this metabolite was denoted as ATCC 20542 by American Type Culture Collection.

In many studies this primary producer of lovastatin, i.e. *A. terreus* ATCC 20542, was used. There are several reports concerning it, both in shake flasks and bioreactors, in which different media and process conditions were used. In the experiments performed in shake flasks a significant impact on the course of lovastatin biosynthesis had, first of all, type and concentration of carbon and nitrogen sources (Bizukojć and Ledakowicz, 2007a; Casas Lopez et al., 2003). In bioreactors it was also necessary to use optimal pH (Bizukojć and Ledakowicz, 2008; Lai et al., 2005), oxygen saturation of broth and aeration rate (Casas Lopez et al., 2005; Lai et al., 2005; Rodriguez Porcel et al., 2006). Lovastatin titres obtained with ATCC 20542 were usually moderate, compared to those, which were the effect of the application of other strains, including the mutant ones (Hajjaj et al., 2001; Kumar et al., 2000; Manzoni et al, 1998).
Lai et al. (2002) carried out experiments in shake flasks and in a 5-litre bioreactor using ATCC 20542 strain. They studied the effect of oxygen carrier addition (aliphatic hydrocarbons) at various concentrations (0-5%, w/v) to the medium. They claimed that oxygen carrier enhanced lovastatin biosynthesis only in shake flasks, while in the bioreactor the contrary effect occurred. These findings were some of the important prerequisites on different behaviour of A. terreus in the shake flask culture and bioreactor. The same authors also tested the effect of pH and dissolved oxygen control in lovastatin biosynthesis by A. terreus ATCC 20542 (Lai et al., 2005). The investigations conducted in a 3-litre batch bioreactor showed that the optimal level of dissolved oxygen (DO) was equal to 20% and pH control decreased lovastatin concentration in the broth.

Different results were obtained by Casas Lopez et al. (2004). They proved that by increasing oxygen content in the aeration gas from 20% to 80% (v/v) a four times higher lovastatin concentration of about 80 mg LOV l\(^{-1}\) can be obtained in a shake flask culture. The same researchers later confirmed these results conducting experiments with oxygen-enriched air (80% v/v oxygen) in a 20-litre bubble column bioreactor.

The influence of carbon and nitrogen sources on the metabolism of A. terreus ATCC 20542 in shake flasks was examined by both Casas Lopez et al. (2003) and Bizukojć and Ledakowicz (2007a). They reported that nitrogen deficiency was favourable for lovastatin biosynthesis in a shake flask culture. However, too low nitrogen concentration may result in a decrease of biomass concentration and consequently a decrease of the titre of lovastatin whose formation is to a large extent growth associated (Bizukojć and Ledakowicz, 2007b). At the same time in the bioreactor lovastatin was better produced at extremely low nitrogen levels. (Bizukojć and Ledakowicz, 2008).

Summing up, it must be noticed that many studies on lovastatin production by A. terreus ATCC 20542 were made in shake flasks (Bizukojć and Ledakowicz, 2007a; Casas Lopez et al., 2003; Lai et al., 2002). Some of the presented results came from bioreactor cultures only (Bizukojć and Ledakowicz, 2008; Lai et al., 2005; Rodriguez Porcel et al., 2006). Thus, it is difficult to find a comparative study analysing simultaneously lovastatin biosynthesis both in shake flasks and bioreactors.

Therefore, this work aims at presenting how to increase the scale of lovastatin biosynthesis on the individual and two carbon sources: lactose and/or glycerol using bioreactors of different working volume starting from shake flasks of 150 ml up to two stirred tank bioreactors of 1.95-litre and 5.3-litre working volumes, in order to achieve comparable lovastatin titres in all these bioreactors.

2. MATERIALS AND METHODS

2.1. Strain and media

The strain Aspergillus terreus ATCC 20542 was employed in the experiments of lovastatin biosynthesis. They were conducted in the shake flasks of 150 ml working volume and in the stirred-tank bioreactors of 1.95-litre and 5.3-litre working volume, at 30°C. The speed of the rotary shaker was constant at 110 min\(^{-1}\). In the bioreactor runs with pO\(_2\) and pH control, the dissolved oxygen saturation was controlled at 20%, by changing the air flow rate and rotary speed of the impeller. The initial rotary speed of the impeller was from 150 to 200 min\(^{-1}\) dependent on the bioreactor. The control of pH was performed with either a solution of sodium and potassium carbonates or sodium hydroxide and kept at the levels between 7 and 7.8, dependent on the experiment. The preculture was prepared from spores grown on 10-days malt extract slants. The spores were washed, suspended in the preculture medium to achieve approximately 10\(^9\) spores per litre and precultivated for 24 hours in the shake flasks. The inoculation was performed with the 24-hour preculture. The media contained the following mineral components: potassium dihydrophosphate KH\(_2\)PO\(_4\): 1.51 g l\(^{-1}\), magnesium sulphate MgSO\(_4\)·7H\(_2\)O: 0.52
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g l⁻¹, sodium chloride NaCl: 0.4 g l⁻¹, zinc sulphate ZnSO₄·7H₂O: 1 mg l⁻¹, ferric nitrate Fe(NO)₃·9H₂O: 2 mg l⁻¹, biotin: 0.04 mg l⁻¹ and 1 ml solution of trace elements per 1 l of medium: The solution of trace elements contained sodium tetraborate Na₂B₄O₇·10 H₂O: 100 mg l⁻¹, manganese chloride MnCl₂: 50 mg l⁻¹, sodium molybdate Na₂MoO₄·2H₂O: 50 mg l⁻¹ and copper sulphate CuSO₄·5H₂O: 250 mg l⁻¹. Yeast extract (BD, USA) was used as the nitrogen source at the concentration of 4 g l⁻¹ and 2 g l⁻¹ (8 g l⁻¹ in the preculture) (Bizukojć and Ledakowicz, 2007a). Glycerol and/or lactose monohydrate were carbon substrates. All the process conditions are collected in Table 1.

Table 1. Experimental scheme for all runs

<table>
<thead>
<tr>
<th>Run</th>
<th>c_LAC₀ [g l⁻¹]</th>
<th>c_GLY₀ [g l⁻¹]</th>
<th>c_YE₀ [g l⁻¹]</th>
<th>vvm [l air min⁻¹ l⁻¹]</th>
<th>pH control</th>
<th>pO₂ control</th>
<th>Source of data</th>
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<tbody>
<tr>
<td>SF1</td>
<td>10</td>
<td>10</td>
<td></td>
<td>0.07</td>
<td>-</td>
<td>-</td>
<td>Bizukojć and Pecyna, 2011</td>
</tr>
<tr>
<td>SF2</td>
<td>0</td>
<td>20</td>
<td></td>
<td>0.06</td>
<td>-</td>
<td>-</td>
<td>Bizukojć and Pecyna, 2011</td>
</tr>
<tr>
<td>SF3</td>
<td>20</td>
<td>0</td>
<td></td>
<td>0.10</td>
<td>-</td>
<td>-</td>
<td>Bizukojć and Ledakowicz, 2007a</td>
</tr>
<tr>
<td>SB1</td>
<td>20</td>
<td>0</td>
<td></td>
<td>0.20</td>
<td>0.308</td>
<td>no</td>
<td>Bizukojć and Ledakowicz, 2008</td>
</tr>
<tr>
<td>SB2</td>
<td>20</td>
<td>0</td>
<td>2</td>
<td>0.24</td>
<td>0.308</td>
<td>no</td>
<td>Bizukojć and Ledakowicz, 2008</td>
</tr>
<tr>
<td>LB1</td>
<td>20</td>
<td>0</td>
<td></td>
<td>0.42</td>
<td>0.3</td>
<td>no</td>
<td>present study</td>
</tr>
<tr>
<td>LB2</td>
<td>10</td>
<td>10</td>
<td></td>
<td>0.21</td>
<td>0.283</td>
<td>no</td>
<td>present study</td>
</tr>
<tr>
<td>LB3</td>
<td>10</td>
<td>10</td>
<td>2</td>
<td>0.15</td>
<td>0.282</td>
<td>no</td>
<td>present study</td>
</tr>
<tr>
<td>LB4</td>
<td>10</td>
<td>10</td>
<td></td>
<td>0.29</td>
<td>0.283-0.998</td>
<td>at 7.15 with NaHCO₃ and KHCO₃ at 20%</td>
<td>present study</td>
</tr>
<tr>
<td>LB5</td>
<td>10</td>
<td>10</td>
<td></td>
<td>0.23</td>
<td>0.283-0.779</td>
<td>at 7.15 with NaOH at 20%</td>
<td>present study</td>
</tr>
<tr>
<td>SF/A1</td>
<td>10</td>
<td>10</td>
<td></td>
<td>0.26</td>
<td>-</td>
<td>no</td>
<td>present study</td>
</tr>
<tr>
<td>SF/A2</td>
<td>10</td>
<td>10</td>
<td></td>
<td>0.26</td>
<td>-</td>
<td>with NaHCO₃ and KHCO₃*</td>
<td>-</td>
</tr>
</tbody>
</table>

*pH level could not be set due to lack of pH measurement on-line. Instead, constant volumes of carbonate solution was added at equal intervals of 24 hours*

2.2. Analytical methods

Lovastatin (LOV) and (+)-geodin (GEOD), (a major by-product in lovastatin biosynthesis) were simultaneously analysed with HPLC (Waters, USA). The analytical conditions were as follows: a Waters Symmetry Shield RP18 column (4.6 mm×250 mm×5μm) at the gradient elution CH₃CN – 0.1% H₃PO₄ (40:60, v/v) up to 7 min and CH₃CN – 0.1% H₃PO₄ (60:40, v/v) from 8 to 30 min. The
temperature of the column was 25°C. Lovastatin was detected by a photodiode array detector at $\lambda=238$ nm and (+)-geodin, at $\lambda=280$ nm (Bizukojć and Ledakowicz, 2007b).

Lactose ($LAC$) and glycerol ($GLY$) were determined on an Aminex HPX-87H (BioRad, USA) column. The compounds were eluted with 0.01N $H_2SO_4$ at a flow rate of 0.6 ml min$^{-1}$ and 60°C and detected by a refractive index detector. All the samples before HPLC analysis were filtered through 0.45 $\mu$m syringe filters with membranes made of regenerated cellulose.

Biomass ($X$) was assayed as dry weight. A 40 ml aliquot was filtered through paper filters and washed with a 10-fold amount of distilled water. Biomass was dried at 105°C to a constant weight. Organic nitrogen ($N$) was determined with the use of carbon and nitrogen analyser IL550TOC-TN (HACH, USA). Prior to the analysis the samples were 20-fold diluted to fit in the analytical range of the instrument.

The error of analytical methods used in the study did not exceed 3% for HPLC analyses and for organic nitrogen determination, and 8% for dry weight biomass assay.

3. RESULTS AND DISCUSSION

3.1. Lovastatin biosynthesis on the individual and two carbon sources in the shake flasks

It is generally believed in biochemical engineering that bioreactor cultivations are more efficient and more useful than shake flasks for several reasons. First of all, in many bioreactor systems, including stirred tank bioreactors, it is possible to control the factors, which may have significant influence on the course of the biosynthesis of a metabolite, e.g. $pH$, redox potential, aeration rate and/or oxygen saturation of the medium. Secondly, larger volumes of bioreactors compared with typical shake flasks allow to increase medium volume and thus higher biomass concentration can be applied. It is especially important, as in most of the cases the obtained metabolite concentration and its formation rate is biomass concentration dependent. Quite often it is also proportional to biomass growth rate (growth associated product formation or mixed associated product formation). Thirdly, broths in stirred tank bioreactors are better mixed improving the availability of all substrates. And last but not least, only bioreactors of various constructions can be practically applied in industrial processes.

As mentioned in the Introduction, in processes of lovastatin biosynthesis carried out in shake flasks, the type and concentration of carbon and nitrogen sources had a significant influence on its production. It is clearly evident in the results presented below and these results are going to be treated as reference values in further considerations concerning bioreactor processes.

In Fig. 1 a comparison of lovastatin concentration and lovastatin and (+)-geodin volumetric formation rates was presented for the runs with $c_{LAC,0}=20$ g l$^{-1}$ ($SF3$) (Bizukojć and Ledakowicz, 2007a), $c_{GLY,0}=20$ g l$^{-1}$ ($SF2$), and with a mixture of lactose and glycerol; $c_{LAC,0}=10$ g l$^{-1}$ and $c_{GLY,0}=10$ g l$^{-1}$ ($SF1$) (Bizukojć and Pecyna, 2011). At all these runs organic nitrogen concentration was the same as 4 g of yeast extract per litre was the sole nitrogen source. The selection of carbon and nitrogen substrate concentration was not incidental here, as it was previously shown that 20 g of lactose and 4 g of yeast extract per litre are optimum in the shake flask culture (Bizukojć and Ledakowicz, 2007a). It must be also mentioned here that glycerol has exactly the same amount of carbon as lactose monohydrate, so these are the equivalent substrates with regard to the amount of elemental carbon.

In the process with the initial glycerol concentration equal to 20 g GLY l$^{-1}$, only about 39 mg LOV l$^{-1}$ was obtained. But when glycerol was replaced by lactose at the same concentration (20 g l$^{-1}$), higher lovastatin titre was found (an increase by 26%). The best results were, however, obtained using these
two carbon sources simultaneously, at about 144 hour of the run there was 87.5 mg LOV l\(^{-1}\) in the broth.

![Graph](image)

**Fig. 1.** Comparison of lovastatin concentration and lovastatin and (+)-geodin volumetric formation rates in the runs with \(c_{\text{LAC},0}=20\) g l\(^{-1}\), \(c_{\text{GLY},0}=20\) g l\(^{-1}\), and with the mixture of lactose and glycerol \((c_{\text{LAC},0}=10\) g l\(^{-1}\) and \(c_{\text{GLY},0}=10\) g l\(^{-1}\)). Data for \(c_{\text{LAC},0}=20\) g l\(^{-1}\) come from Bizukojć and Ledakowicz (2007a). Data for \(c_{\text{GLY},0}=20\) g l\(^{-1}\) and for mixture of lactose and glycerol \((c_{\text{LAC},0}=10\) g l\(^{-1}\) and \(c_{\text{GLY},0}=10\) g l\(^{-1}\)) come from Bizukojć and Pecyna (2011).

On the basis of lovastatin volumetric formation rate curves it can be noted that the utilisation of lactose in the medium in the first 48 hours of the run considerably favoured lovastatin biosynthesis. The maximum of lovastatin volumetric formation rate in the run with lactose was exactly at about 48 hour (0.80 mg LOV l\(^{-1}\) h\(^{-1}\)), while in the run with glycerol this maximum was significantly lower (0.34 mg LOV l\(^{-1}\) h\(^{-1}\)) and occurred not earlier than at 96 hour of the experiment. What is more, glycerol as the individual carbon source exerted an undesired effect of elevated (+)-geodin production. In the run with \(c_{\text{GLY},0}=20\) g l\(^{-1}\) (+)-geodin volumetric formation rate reached the highest values compared with the other two processes. In the run with two carbon sources lovastatin volumetric formation rate had two maxima and they were higher both for lactose and glycerol utilisation than in the runs in which these components were used as the individual carbon sources (Bizukojć and Pecyna, 2011).

Similar studies with the use of different carbon sources in shake flasks were performed by Casas Lopez et al. (2003). These authors obtained almost identical results as (Bizukojć and Pecyna, 2011) for lactose as carbon sources \((1.33\) g l\(^{-1}\) of yeast extract), about 90 mg LOV l\(^{-1}\). What is surprising, the same high amount of lovastatin for glycerol was obtained \((90\) mg LOV l\(^{-1}\)), which is in disagreement with the results presented in Fig. 1.

The obtained levels of lovastatin in the shake flasks are satisfactory compared to the results of the other authors for *A. terreus* ATCC 20542 in similar conditions. Thus, at this moment, having obtained such good results as the ones in the shake flask culture, one faces the problem of how to achieve similar lovastatin titres in a bioreactor.
3.2. Lovastatin biosynthesis in stirred tank bioreactors of different working volumes

The experiments on lovastatin biosynthesis conducted in the shake flasks showed that a high lovastatin concentration (from 39 to 90 mg LOV l⁻¹) can be achieved using the optimum carbon (lactose and/or glycerol) and nitrogen (yeast extract) sources with their initial concentration equal to 20 g l⁻¹ and 4 g l⁻¹, respectively. The difference in the quality of glycerol and lactose referred to lovastatin biosynthesis was discussed elsewhere (Bizukojć and Pecyna, 2011).

In order to test lovastatin production in the bioreactors and hopefully obtain higher lovastatin concentration a series of batch processes in stirred-tank bioreactors of 1.95- and 5.3-litre working volumes were performed. Initially, the following conditions: vvm equal to about 0.3 l air min⁻¹ l⁻¹ no dissolved oxygen and pH control were set to somewhat follow the conditions in the shake flasks. In Fig. 2 lovastatin and lactose concentration, and lovastatin formation rates in the runs with \( c_{LAC,0} = 20 \text{ g LAC l}^{-1} \) conducted in the shake flasks (reference run SF3), a small 1.95-litre bioreactor (SB1) and a large 5.3-litre bioreactor (LB1) working volume are presented. Despite the application of lactose at the same concentration in all the three processes, the highest lovastatin concentration (53 mg LOV l⁻¹) was obtained in the shake flasks (SF3), while the lowest (4.2 mg LOV l⁻¹) for the run in the 5.3-litre bioreactor (LB1). Also in the 1.95-litre bioreactor (SB1) the results were not as satisfactory (39 mg LOV l⁻¹) as they were in the shake flasks. It is not easy to explain this phenomenon. Its reason might probably be the level of oxygen dissolved in the medium. No \( \text{pO}_2 \) measurement could be made in the shake flasks due to the technical reasons, so actually it is difficult to compare the conditions in the flasks and bioreactor. Neither can the aeration rate be measured and controlled in the shake flasks. Convective mass transfer coefficient might not be so low in the shake flasks as expected, so the cells might have more oxygen available and thus bioreactor conditions were not properly set. On the other hand, it is clearly seen that volumetric lactose uptake rate for all these runs was approximately constant between 24 and 120 hour of each run and equal to 0.17, 0.12 and 0.15 g LAC l⁻¹ h⁻¹ for SF3, SB1 and LB1, respectively. Only lower consumption of lactose between 24 and 72 hour in SB1 in comparison to the other runs was observed. As volumetric substrate utilisation rate is always proportional to the amount of oxygen available and utilised (oxygen uptake rate), the hypothesis of better or worse aeration in these two runs is not so convincing. What is more, similar conclusions can be drawn from the analysis of nitrogen concentration curves. They were similar too and also in SB1 slower consumption of nitrogen between the same hours was noted (Fig. 2). Also, the changes of substrate and biomass concentrations show no extreme differences between each other. The largest discrepancies are only seen for lovastatin curves. All in all, lovastatin formation is somewhat deteriorated in the bioreactor.

In order to finally confirm this negative effect in the bioreactor process, lovastatin volumetric formation rates were calculated. The highest value of the maximum of lovastatin volumetric rate (0.80 mg LOV l⁻¹ h⁻¹) was observed in the run SF3 (as described earlier in section 3.1. Lovastatin biosynthesis on the individual and two carbon sources in the shake flasks). In run SB1 this maximum occurred about 65 hour and was equal to 0.3 mg LOV l⁻¹ h⁻¹, while in run LB1 it was only 0.18 LOV l⁻¹ h⁻¹ (at about 32 hour).

It was shown previously (Bizukojć and Pecyna, 2011) that in shake flasks the use of two carbon sources led to more efficient lovastatin production, so another bioreactor process was conducted with the use of two carbon sources. The results are presented in Fig. 3 and they were unsatisfactory for the bioreactor run (LB2) again. Firstly, it is well seen that this time in the bioreactor slower consumption of carbon substrates occurred. Conversely, in the run SF1 glycerol was fully utilised within 72 hours, then lactose was almost completely depleted within the next 48 hours. In the bioreactor glycerol consumption was delayed by at least 24 hours and lactose was not fully utilised. The ultimate substrate consumption led to an intensive lovastatin biosynthesis (about 90 mg LOV l⁻¹) in the shake flasks, while in the bioreactor only about 11 mg LOV l⁻¹ was obtained.
Impact of bioreactor scale on lovastatin biosynthesis by Aspergillus terreus ATCC 20542 in a batch culture

Fig. 2. Lovastatin, lactose, nitrogen and biomass concentration and lovastatin volumetric formation rate in the runs SF3, SB1 and LB1; Nitrogen source: $c_{\text{YE}}=4$ g l$^{-1}$ for all runs. Data for $c_{\text{LAC}}=20$ g l$^{-1}$ in the shake flasks come from Bizukojć and Ledakowicz, (2007a). Data for $c_{\text{LAC}}=20$ g l$^{-1}$ in the small stirred-tank bioreactor come from Bizukojć and Ledakowicz, (2008). Processes in the bioreactors without pH and $pO_2$ control; aeration SB1: $v_{vm}=0.308$ l air min$^{-1}$ l$^{-1}$, LB1: $v_{vm}=0.3$ l air min$^{-1}$ l$^{-1}$

It is necessary to answer the question, what the reason of this phenomenon was, i.e. what influenced such a slow utilisation of substrates in the bioreactor and why the substrates in the shake flask were completely consumed. In this case the hypothesis about insufficient bioreactor aeration is more convincing, although a drastic oxygen limitation was still not observed as $pO_2$ was around 10% (Fig. 3). The researchers who conducted lovastatin biosynthesis in bioreactors claimed that there were several factors that may have played an important role in these processes (Bizukojć and Ledakowicz, 2008; Casas Lopez et al, 2004; Lai et al., 2005; Rodriguez Porcel et al., 2006). Among these factors the initial concentration of nitrogen sources, pH control and $pO_2$ control were mentioned. Especially, pH curve is worth further analysis as it changes to a great extent within the cultivation time. One observes the maximum of about 7.6 at about 24 hour and a further decrease down to 6.4 at the end of the run. Thus, all these factors mentioned above are going to be studied further.
Fig. 3. Lovastatin, lactose and glycerol concentration obtained on the mixture of lactose and glycerol in the shake flasks and in the 5.3-litre stirred-tank bioreactor; \(c_{\text{LAC},0}=10 \, \text{g l}^{-1}, c_{\text{GLY},0}=10 \, \text{g l}^{-1}, c_{\text{YE},0}=4 \, \text{g l}^{-1}\). Process in bioreactor without \(pH\) and \(pO_2\) control; Aeration: \(v_{\text{vm}}=0.283 \, \text{l}_{\text{air}} \, \text{min}^{-1} \, \text{l}^{-1}\)

### 3.3. Modification of initial nitrogen concentration

It was proved that an elevated organic nitrogen concentration inhibits lovastatin biosynthesis. For example, Casas Lopez et al. (2003) showed that the deficiency of nitrogen improved lovastatin titre. It was also confirmed by Bizukojć and Ledakowicz (2007a). However, all these results were obtained from shake flasks, not bioreactors.

Heretofore, processes have been presented, in which the initial nitrogen concentration was equal to 4 g l\(^{-1}\). However, it is known from previous reports that this initial nitrogen concentration was optimal only in the runs performed in shake flasks (Bizukojć and Ledakowicz, 2007a), because in a 1.95-litre bioreactor the initial yeast extract concentration of 2 g l\(^{-1}\) was more favourable for lovastatin production (Bizukojć and Ledakowicz, 2008). Thus, the experiment in the 5.3-litre bioreactor with the use of the initial nitrogen concentration equal to 2 g l\(^{-1}\) and two carbon substrates (the best lactose and glycerol composition upon previous experiments) was conducted. In Fig. 4 the evolution curves of lovastatin, nitrogen, lactose, glycerol and biomass for the runs in both sizes of the stirred-tank bioreactors with the use of \(c_{\text{YE},0}=2 \, \text{g l}^{-1}\) are presented.

Despite the fact that in the 5.3-litre bioreactor even two carbon sources were used, which should theoretically improve lovastatin production, the obtained lovastatin titres occurred to be worse than in the 1.95-litre bioreactor (SB2). So the application of a deficient amount of organic nitrogen in the larger stirred-tank bioreactor was not as encouraging as in the smaller one, despite the favourable carbon source sources mixture. The faster nitrogen uptake rate for the first 24 hours in the 5.3-litre bioreactor compared with the run in the 1.95-litre bioreactor was observed and this fact is probably connected with glycerol consumption as this substrate is faster assimilated by \(A. \text{terreus}\) than lactose (Bizukojć
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and Pecyna, 2011). What is more, in the run LB3 lactose consumption was not observed at all and it was not connected with oxygen deficiency (Fig. 4) but rather with the lack of organic nitrogen. It must be also mentioned here again that lactose and glycerol are equivalent substrates with regard to the amount of elemental carbon (Bizukojć and Pecyna, 2011).

To sum up, the decrease of the initial nitrogen concentration in the 5.3-litre bioreactor from 4 to 2 g l⁻¹ did not improve lovastatin production. Additionally, analysing biomass concentration curves, slower biomass growth during first 48 hours in the run LB3 was observed. Nevertheless, the biomass curves did not differ much. Still biomass concentration in the idiophase was at least by 30% lower than in the runs with 4 g of yeast extract per litre (compare Fig. 2 and 4). In the runs SB3 and LB2 the evolution of (+)-geodin was also analysed (data not shown). Enhanced biosynthesis of this metabolite in the large bioreactor (LB3) was observed and the amount of (+)-geodin in the broth exceeded 200 mg l⁻¹. The presence of glycerol in the medium probably had a significant influence on this results, which was mentioned in the previous section.

Fig. 4. Comparison of lovastatin, nitrogen, lactose, glycerol, biomass and dissolved oxygen concentrations for the runs LB3 and SB2. Data for SB2 come from Bizukojć and Ledakowicz (2008); c_{LAC,0}=20 g l⁻¹, vvm=0.308 l air min⁻¹ l⁻¹, LB3: c_{LAC,0}=10 g l⁻¹, c_{GLY,0}=10 g l⁻¹, vvm=0.282 l air min⁻¹ l⁻¹; runs without pH and pO₂ control; Nitrogen source c_{YE,0}=2 g l⁻¹

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3.4. Application of the simultaneous pH and pO₂ control

A study on the influence of pH control onLovastatin biosynthesis by A. terreus ATCC 20542 was previously conducted by Lai et al. (2005). In their experiments, 1 N HCl or 1 N NaOH solution were used to pH adjustment at three different levels i.e. 5.5, 6.5 and 7.5. This way Lai et al. (2005) concluded that pH control caused a decrease ofLovastatin production and biomass concentration. They suggested that unsuitable pH could cause the activation of the enzymes that destroy the structure ofLovastatin. At the same time these authors did not provide data on the impact of pH on (+)-geodin biosynthesis.

Other results of pH control inLovastatin biosynthesis were obtained by Bizukojć and Ledakowicz (2008) in the experiments in a 1.95-litre bioreactor. That pH level was controlled at 7.6 or 7.8 by an addition of sodium and potassium bicarbonate solution. The motivation for this action was the fact that several reactions of primary metabolism of A. terreus are dependent on bicarbonate ions concentration, including malonyl-CoA formation (Bizukojć and Ledakowicz, 2008). Malonyl-CoA is the substrate for polyketide chain elongation and used by both Lovastatin nonaketide synthase and Lovastatin diketide synthase. (Kennedy et al., 1999). The application of pH control also significantly decreased (+)-geodin concentration in the broth (Bizukojć and Ledakowicz, 2008).

Another factor influencing Lovastatin formation was the concentration of dissolved oxygen in the medium. Casas Lopez et al. (2004) conducted experiments in a 5-litre bioreactor, in which aeration rate equal to 1 l air min⁻¹ l⁻¹ was applied. Additionally, aeration gas was oxygen-enriched. This treatment resulted in the increase of Lovastatin production from 20 to 80 mg LOV l⁻¹. Rodriguez Porcel et al. (2006), also proved that the control of dissolved oxygen on the high level (400%) in the broth was necessary for attaining suitable Lovastatin production in the slurry bioreactor. Lai et al. (2005) received the best results, when dissolved oxygen was controlled at the level of 20%. Bizukojć and Ledakowicz (2008), contradicted the result presented earlier, proving that a lower vvm at the level of about 0.3 l air min⁻¹ l⁻¹ was more optimal for Lovastatin biosynthesis. What is more, a high vvm equal to 1 l air min⁻¹ l⁻¹ led to an efficient (+)-geodin formation (Bizukojć and Ledakowicz, 2008).

The actions described in the previous sections did not assure acceptable Lovastatin titles in the 5.3 litre bioreactor and the aforementioned premises are sufficient to verify the simultaneous impact of pH and pO₂ control inLovastatin biosynthesis in this bioreactor using glycerol and lactose as carbon sources.

At first, the experiments with (SF/A2) and without (SF/A1) bicarbonate solution addition in the shake flasks were performed and their results being the reference to the bioreactor runs, are shown in Fig. 5a.

It is clearly seen that a satisfactory Lovastatin concentration (77.3 mg LOV l⁻¹) was only obtained in the run SF/A1 without bicarbonates addition. The appreciable increase in Lovastatin concentration from 27.6 to 67.3 mg LOV l⁻¹ between 96 and 120 hour took place. At the same time the levels of pH, after an initial increase, dropped from 6.67 to 6.35. In this run a decrease of pH in the late idiophase contributed to an increase of (+)-geodin formation compared to the run SF/A2 and an addition of bicarbonates solution that increased pH in run SF/A2 led to a significant decrease of the secretion of this by-product. It was in agreement with the previous observations of Bizukojć and Ledakowicz (2008). Thus, it is easy to notice that in the processes conducted in the shake flasks pH control, actually an addition of bicarbonates solution was not advisable with regard to Lovastatin biosynthesis but it was a suitable method to decrease (+)-geodin concentration in the broth. The consecutive step was to check if the application of pH control would lead to an improvement of Lovastatin production in the 5.3-litre bioreactor, bearing in mind disappointing results from the shake flask culture and previous encouraging results obtained in the 1.95-litre bioreactor (Bizukojć and Ledakowicz, 2008).
Fig. 5. Comparison of lovastatin and (+)-geodin concentration, influence of pH and pO2 control: (a) processes in the shake flasks without SF/A1 and with SF/A2 with pH control; (b) processes in the bioreactor run LB4 pH controlled with NaHCO3 and KHCO3, run LB5 pH controlled with NaOH, run LB2 without pH control; runs SF/A1, SF/A2, LB4, LB5, LB2 (cLAC,0=10 g l−1, cGLY,0=10 g l−1, cYE,0=4 g l−1), runs LB4 and LB5: pO2 control at 20%, run LB2 aeration at vvm=0.283 lair min−1 l−1.

In Fig. 5b the results from the run LB4 (pH controlled at 7.15 with a solution of NaHCO3 and KHCO3) and from LB5 (pH controlled at 7.15 with a solution of NaOH) were compared to the earlier presented run LB2 (pH-uncontrolled). It should be noted that in the runs LB4 and LB5 also pO2 control at 20% saturation was applied for the reasons described further in this section. In the runs conducted in the 5.3-litre bioreactor the application of the simultaneous pH and pO2 control led to a significant increase of lovastatin formation (75.5 mg LOV l−1). This result was now closer to the titre obtained in the shake
flask run (SF/A1) but without pH control (87.4 mg LOV l⁻¹) than in the other bioreactor runs described here. Thus, higher pH seems to be favourable in the bioreactor, not in the shake flasks. But what is the most important thing, it was also observed that the use of bicarbonate ions was most favourable for lovastatin formation by A. terreus ATCC 20542, because in the run LB5, in which the solution of NaOH was used the enhancement of lovastatin production was much weaker compared to the run LB4. Thus, it is not only the effect of pH but of inorganic carbon in the form of bicarbonate that influences lovastatin formation. Additionally, bicarbonate ions to a greater extent than hydroxide ions contributed to the inhibition of (+)-geodin formation. Now, it became understandable why Lai et al. (2005) did not recommend pH control, as they used NaOH and HCl solutions for this purpose.

It was found earlier (Bizukojć and Ledakowicz, 2008) that an increase of aeration in the 1.95-litre bioreactor rate and as a consequence of oxygen saturation in the system led to efficient (+)-geodin formation, whose titre even exceeded 100 mg GEOD l⁻¹, and worse lovastatin formation. So at the first glance it seemed not to be the right action to increase either aeration rate or oxygen saturation in the 5.3-litre bioreactor. In the 1.95-litre bioreactor the optimal vvm was equal to about 0.3 lₘᵦ min⁻¹ l⁻¹. However, it is seen in Fig. 6 that at a low vvm substrate uptake rate deteriorated in the 5.3-litre bioreactor and was even lower than in the shake flasks. So it was decided to hold oxygen saturation at 20% level by increasing the air flow rate and rotary speed of the impeller in this bioreactor. In this way the range of vvm varied from 0.282 to 0.998 lₘᵦ min⁻¹ l⁻¹ dependent on the time of the process (Fig. 6).

![Fig. 6. Influence of pO₂ control on the formation of lovastatin and on the lactose and glycerol uptake rates in runs SF1, LB2, LB4; pO₂ control at 20% and pH control, LB2; vvm=0.283 lₘᵦ min⁻¹ l⁻¹](image)

First of all, this action did not lead to elevated (+)-geodin levels, due to simultaneous pH control. Secondly, as a result of pO₂ control at the level of 20% in the run LB4 up to 72 hour a similar uptake rate of glycerol compared with the run SF/A1 was observed. It was much higher than in the run LB2 without pO₂ and pH control. What is more, the same amount of lovastatin for SF/A1 and LB4 within
this period of time was obtained (Fig. 6). Unfortunately, later despite high concentration of dissolved oxygen in the run LB4 the consumption of lactose was still lower compared with the run SF1 and lovastatin formation slowed down. However, this is the evidence that in the bioreactor systems the application of simultaneous pH and pO2 control is necessary in order to increase lovastatin production and to follow the course of the process as in the efficient shake flask experiment.

To sum up, the discussion above indicates that a simple transfer of the medium for lovastatin production from the efficient shake flask culture to the bioreactor leads to the worsening of the process with regard to lovastatin production. Therefore, several additional measures, namely pO2 control, which is quite obvious, and pH control with bicarbonate solution, which is far more surprising and less obvious, must be taken to achieve at least similar results in the bioreactor compared to those in the shake flasks.

4. CONCLUSIONS

The drawn conclusions are going to give some practical guidance to the problem of how to transfer lovastatin production from the shake flasks to the bioreactor. Thus, on the basis of the presented results it can be said that:

- Using the same type and concentration of carbon and nitrogen sources in the bioreactor, considerably lower lovastatin titres are obtained compared with the similar processes conducted in the shake flasks.

- A low initial nitrogen concentration (2 g l⁻¹) may increase lovastatin formation but not always as it was observed only in the 1.95-litre stirred tank bioreactor. On the other hand it may also lead to a decrease of substrate uptake rate and worse biomass formation aggravating lovastatin production too, which was observed in the 1.95-litre bioreactor.

- The application of pH control using bicarbonates solution in the processes conducted in the shake flasks does not lead to the enhancement of lovastatin production, although this control slightly decreases the formation of the by-product, (+)-geodin. When pH and pO2 control are used simultaneously in the 5.3-litre bioreactor, a satisfactory lovastatin titre is obtained; even seven times higher lovastatin concentration compared with the bioreactor runs without pH and pO2 control. It is definitely important to control pH at a mild alkaline level slightly above 7 with bicarbonates solution, as the use of sodium hydroxide does not give such good effects. Thus, not only pH but the effect of bicarbonate ions is favourable for lovastatin production.

- A controlled level of dissolved oxygen in the medium for the run in the 5.3-litre bioreactor is required for the efficient utilisation of the substrates, which somewhat enhances lovastatin production.

The authors wish to acknowledge National Science Centre (Republic of Poland) for the partial financial support of this work, project no. NN209 765240.

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Received 23 September 2011
Received in revised form 03 January 2012
Accepted 03 January 2012