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CARBON DIOXIDE FROM ALCOHOLIC FERMENTATION AS A CARBON SOURCE FOR FED-BATCH CULTIVATION OF *Arthrospira platensis*

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Abstract

World's production of ethanol has increased dramatically in recent years. Brazil is the world's largest exporter of bioethanol and second-largest producer after the United States. Considering the increasing demand for this fuel and the fact that alcoholic fermentation is responsible for a CO₂ release, on weight basis, almost coincident with ethanol production, it would be interesting to develop a process for CO₂ fixation able to turn it into a useful product. Photosynthetic microorganisms can fix CO₂ efficiently producing biomass that contains high-value bioactive products and may provide a very promising alternative for the current CO₂ mitigation strategies. Nowadays, there are numerous commercial applications of *Arthrospira platensis* biomass such as the enhancement of the nutritional value of foods and animal feed, bioremediation, and use in cosmetics. The objective of this work was to evaluate the *Arthrospira platensis* cultivation using CO₂ from alcoholic fermentation and either urea or nitrate as nitrogen source at different light intensities ($60 \leq I \leq 240 \mu\text{mol photons m}^{-2} \text{s}^{-1}$). The CO₂ source (pure CO₂ or from alcoholic fermentation) did not influence the maximum cell concentration (X_m), cell productivity (P_X) and nitrogen-to-cell conversion factor ($Y_{X/N}$). On the other hand, the use of urea instead of nitrate led to higher $Y_{X/N}$ values. X_m and P_X increased when I was increased from 60 to 120-240 $\mu\text{mol photons m}^{-2} \text{s}^{-1}$. Using CO₂ from alcoholic fermentation, the best performance ($X_m=2952 \pm 35 \text{ mg L}^{-1}$, $P_X=425 \pm 5.9 \text{ mg L}^{-1} \text{ d}^{-1}$ and $Y_{X/N}=15 \pm 0.20 \text{ mg mg}^{-1}$) was obtained at $I = 120 \mu\text{mol photons m}^{-2} \text{s}^{-1}$ with urea. The results obtained in this work demonstrate that urea and CO₂ from alcoholic fermentation could be simultaneously used in large-scale cultivations to reduce the environmental impact associated to the release of this greenhouse gas as well as to decrease the production cost of this cyanobacterium.
