

Biotechnology and Its Simultaneous Evolution

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Mini Review

Biotechnology is currently developing across borders rapidly and simultaneously. New equipment is continuously developed and refined to improve production processes and quality control associated with obtaining the most diverse bioproducts such as biosurfactants, natural dyes, antioxidants and antibiotics. There is no industrial biotechnological process that works generically and each requires particular solutions that can sustain a good scale up from scientific and technological points of view.

State-of-the-art centrifuges and decanters “taylor made” with high commercial value have a prominent place in unit operations for the separation of micro-organisms of interest or the culture medium containing bioproducts or only for clarification of the final product as in the industrial procurement of wine.

High-sensitivity flow cytometers with cellular recognition capability utilizing information on biochemical composition, shape and size help in screening and sorting out new strains of microorganisms to be grown for the discovery of promising new molecules (e.g. antivirals to fight HIV).

On the other hand, the huge variety of new newly synthesized reagents – molecular probes – has been assisting in the detection and quantification of target molecules “in situ”, an extremely important fact for rapid evaluation and intervention throughout bioprocesses with short production cycles.

The company Life Technologies Corporation (Figure 1), recently acquired by Thermo Fisher Scientific, has available in its catalog of molecular probes 35 reference standards for flow cytometry at the disposal of the international scientific community, in addition to other catalogs containing hundreds of other equipment, kits and reagents.



Figure 1: Electroporator manufactured by Life Technologies used internally in San Diego/USA (author’s personal archive).

However, the advances made by new equipment and methodologies applied to unit operations can at any time be radically modified by interventions that may change the constitution and expression of the genetic code present in target microorganisms.

The programmed decrease in robustness and mechanical strength of plant cell walls, for example, considerably decreases the energy required - whether thermal or mechanical-for their disruption, facilitating the extraction of target compounds with significantly lower investment and operating costs when designing

a future business venture.

The heat resistance of some microorganisms provided by the biosynthesis of heat shock resistant proteins (chaperones) in turn can radically alter the need to cool closed systems whose crops are typically crashed at temperatures above 35 °C (Figure 2). The absence of coolers for biotechnological processes means a considerable competitive advantage by saving the purchase of this equipment, maintenance and also the energy consumption caused by its use.



Figure 2: Closed system to obtain micro-algal biomass running at Petrobras R&D Center.

Beyond the examples given above there are many other possibilities from the modification and edition of the genetic code with potential to promote major changes in the configuration of future processes or that are already being used to manufacture several categories of bioproducts, including the latest generation biofuels.

A partnership between Exxon Mobil and Synthetic Genomics recently yielded encouraging results published in the Journal

Nature Biotechnology in 2017 on achieving high yield in genetically modified *Nannochloropsis gaditana* crops that successfully achieved high lipid concentration (40% in ash free dry weight).

Other undisclosed advances are certainly underway and could at any time improve or even replace existing commercial bioprocesses and bioproducts (e.g. replacement of vaccines and antidotes produced with the use of animals by those produced using plants).

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