

Effect of Bcl-2 overexpression on cell cycle and antibody productivity in chemostat cultures of myeloma NS0 cells

Abstract

Chemostat cultures of NS0 cell lines were carried out at dilution rates ranging from 0.8 d⁻¹ to 0.2 d⁻¹. Compared with the control, the viable cell density of the Bcl-2 cell line was approximately 10% higher at 0.8 d⁻¹ and increased to 55% when the dilution rate was reduced to 0.2 d⁻¹. As the dilution rate was reduced, the viability of the two cultures diverged reaching a difference of 43% at 0.2 d⁻¹. The specific growth rate of the control cells was the same as the dilution rate down to a value of 0.6 d⁻¹. By contrast, the specific growth rate of Bcl-2 cells was parallel to the dilution rate down to a value as low as 0.3 d⁻¹. For both NS0 cell lines, the G1 cell population decreased, while the S and G2/M cell populations increased as the dilution rate was reduced. The antibody titer of the control cells increased from 7 to 21 µg·ml⁻¹ as the dilution rate was reduced from 0.8 to 0.2 d⁻¹. With an initial increase from 2 to 15 µg·ml⁻¹ as the dilution rate was reduced from 0.8 to 0.4 d⁻¹, the antibody titer of the Bcl-2 cells remained constant as the dilution rate was further reduced to 0.2 d⁻¹. A good correlation between specific antibody production rate and the percentage of G2/M cells was observed. © 2005, The Society for Biotechnology.

Keyword: Apoptosis; bcl-2; Bioreactor; Chemostat culture; NS0 myeloma cells