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Theories, Models and Methods

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and experts in their respective fields ranging from biomaterials and bioreactors to gene delivery and metabolic engineering. Particular emphasis was placed on including reviews that discuss various aspects of the biochemical processes underlying cell function such as signaling, growth, differentiation and communication. The reviews of research topics cover two main areas: cellular and non-cellular components and assembly, evaluation and optimization of tissue function and integrated reactor or implant system development for research and clinical applications. Many of the reviews illustrate how biochemical engineering methods are used to produce and characterize novel materials, e.g. genetically engineered natural polymers, synthetic scaffolds with specific attachment sites or inductive factors whose unique properties enable increased levels of control over tissue development and architecture.

Cell Culture Engineering Wei-Shu Hu, 2006-08-16. Since the introduction of recombinant human growth hormone and insulin a quarter century ago, protein therapeutics has greatly broadened the horizon of health care. Many patients suffering with life-threatening diseases or chronic dysfunctions which were medically untreatable not long ago can attest to the wonder these drugs have achieved. Although the first generation of protein therapeutics was produced in recombinant *Escherichia coli*, most recent products use mammalian cells as production hosts. Not long after the first production of recombinant proteins in *E. coli*, it was realized that the complex tasks of most post-translational modifications on proteins could only be efficiently carried out in mammalian cells. In the 1990s we witnessed a rapid expansion of mammalian cell-derived protein therapeutics, chiefly antibodies. In fact, it has been nearly a decade since the market value of mammalian cell-derived protein therapeutics surpassed that of those produced from *E. coli*. A common characteristic of recent antibody products is the relatively large dose required for effective therapy, demanding larger quantities for the treatment of a given disease. This coupled with the broadening repertoire of protein drugs has rapidly expanded the quantity needed for clinical applications. The increasing demand for protein therapeutics has not been met exclusively by construction of new manufacturing plants and increasing total volume capacity. More importantly, the productivity of cell culture processes has been driven upward by an order of magnitude in the past decade.

Gene Therapy and Gene Delivery Systems David V. Schaffer, Weichang Zhou, 2005-12-20. 1 D V Schaffer W Zhou Gene Therapy as Future Human Therapeutics 2 J Heidel S Mishra M E Davis Molecular Conjugates 3 M Manthorpe P Hobart G Hermanson M Ferrari A Geall B Goff A Rolland Plasmid Vaccines and Therapeutics From Design to Applications 4 S R Little R Langer Non Viral Delivery of Cancer Genetic Vaccines 5 J C Grieger R J Samulski Adeno Associated Virus as a Gene Therapy Vector Vector Development Production and Clinical Applications 6 J H Yu D V Schaffer Advanced Targeting Strategies for Murine Retroviral and Adeno Associated Viral Vectors 7 N Loewen E M Poeschla Lentiviral Vectors 8 N E Altaras J G Aunins R K Evans A Kamen J O Konz J J Wolf Production and Formulation of Adenovirus Vectors

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