

Protein Purification: Design and Scale Up of Downstream Processing

Wheelwright, Scott M.

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Protein Purification Design And Scale Up Of Downstream Processing

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Protein Purification Design And Scale Up Of Downstream Processing:

Protein Purification Scott M. Wheelwright, 1991 A comprehensive introduction to the art and science used by the biochemist and process engineer in the design and optimization of large scale protein purification processes Covers the basics of proteins properties the unit operations involved in protein purification as well as important related issues of process design integration with upstream operations cost and timing to market Includes an abundance of figures and tables

Scale-Up and Optimization in Preparative Chromatography Anurag Rathore, Ajoy Velayudhan, 2002-09-26 This text offers fundamental analysis of the underlying physicochemical principles to select optimal modes of interaction design efficient separations and increase productivity in the manufacture of novel therapeutics and pharmaceuticals

Handbook of Downstream Processing E. Goldberg, 2012-12-06 The last two decades have seen a phenomenal growth of the field of genetic or biochemical engineering and have witnessed the development and ultimately marketing of a variety of products typically through the manipulation and growth of different types of microorganisms followed by the recovery and purification of the associated products The engineers and biotechnologists who are involved in the full scale process design of such facilities must be familiar with the variety of unit operations and equipment and the applicable regulatory requirements This book describes current commercial practice and will be useful to those engineers working in this field in the design construction and operation of pharmaceutical and biotechnology plants It will be of help to the chemical or pharmaceutical engineer who is developing a plant design and who faces issues such as Should the process be batch or continuous or a combination of batch and continuous How should the optimum process design be developed Should one employ a new revolutionary separation which could be potentially difficult to validate or use accepted technology which involves less risk Should the process be run with ingredients formulated from water for injection deionized water or even filtered tap water Should any of the separations be run in cold rooms or in glycol jacketed lines to minimize microbial growth where sterilization is not possible Should the process equipment and lines be designed to be sterilized in place cleaned in place or should every piece be broken down cleaned and autoclaved after every turn

Downstream Processing of Proteins Mohamed A.

Desai, 2008-02-05 Considerable effort and time is allocated to introducing cell culture and fermentation technology to undergraduate students in academia generally through a range of courses in industrial biotechnology and related disciplines Similarly a large number of textbooks are available to describe the applications of these technologies in industry However there has been a general lack of appreciation of the significant developments in downstream processing and isolation technology the need for which is largely driven by the stringent regulatory requirements for purity and quality of injectable biopharmaceuticals This is particularly reflected by the general absence of coverage of this subject in many biotechnology and related courses in educational institutions For a considerable while I have felt that there is increasing need for an introductory text to various aspects of downstream processing particularly with respect to the needs of the

biopharmaceutical and biotechnology industry. Although there are numerous texts that cover various aspects of protein purification techniques in isolation, there is a need for a work that covers the broad range of isolation technology in an industrial setting. It is anticipated that *Downstream Processing of Proteins: Methods and Protocols* will play a small part in filling this gap and thus prove a useful contribution to the field. It is also designed to encourage educational strategists to broaden the coverage of these topics in industrial biotechnology courses by including accounts of this important and rapidly developing element of the industrial process.

Process Scale Bioseparations for the Biopharmaceutical Industry

Abhinav A. Shukla, Mark R. Etzel, Shishir Gadgil, 2006-07-07. The biopharmaceutical industry has become an increasingly important player in the global economy and the success of these products depends on the development and implementation of cost-effective, robust and scalable production processes. Bioseparations, also called downstream processing, can be a key source of competitive advantage to biopharmaceuticals.

Biosimilars and Interchangeable Biologics

Sarfaraz K. Niazi, 2016-01-05. What's the Deal with Biosimilars? Biosimilars are gaining momentum as new protein therapeutic candidates that can help fill a vital need in the healthcare industry. The biological drugs are produced by recombinant DNA technology that allows for large-scale production and an overall reduction in costs and development time. Part of a two-volume set that covers varying aspects of biosimilars, *Biosimilars and Interchangeable Biologics: Tactical Elements* explores the development and manufacturing of biosimilars and targets challenges surrounding the creation of these products. This includes manufacturing, production costs and intellectual property barriers, particularly in regulated markets. Regulatory agencies are still in the process of developing guidelines. It addresses the complexity of biological drugs and discusses specific structural elements vital to the functionality, immunogenicity and safety of biosimilar products. Of specific interest to practitioners, researchers and scientists in the biopharmaceutical industry, this volume provides an overall understanding of the hurdles, difficulties and practicalities of developing a strong plan. It introduces a step-by-step approach for creating a strategy that helps develop and manufacture a biosimilar product while reducing overall production costs and meeting the requirements of biosimilarity based on analytical and functional pharmacokinetic/pharmacodynamic where applicable and nonclinical toxicology or toxicokinetic similarity where appropriate while remaining competitive in the market.

Handbook of Methods and Instrumentation in Separation Science, 2009-11-11. *Handbook of Methods and Instrumentation in Separation Science: Volume 1* provides concise overviews and summaries of the main methods used for separation. It is based on the *Encyclopedia of Separation Science*. The handbook focuses on the principles of methods and instrumentation. It provides general concepts concerning the subject matter; it does not present specific procedures. This volume discusses the separation processes including affinity methods, analytical ultracentrifugation, centrifugation, chromatography and use of decanter centrifuge and dialysis. Each methodology is defined and compared with other separation processes. It also provides specific techniques, principles and theories concerning each process. Furthermore, the handbook presents the applications

benefits and validation of the processes described in this book This handbook is an excellent reference for biomedical researchers environmental and production chemists flavor and fragrance technologists food and beverage technologists academic and industrial librarians and nuclear researchers Students and novices will also find this handbook useful for practice and learning One stop source for information on separation methods General overviews for quick orientation Ease of use for finding results fast Expert coverage of major separation methods Coverage of techniques for all sizes of samples pico level to kilo level

Bioprocess Engineering Principles Pauline M. Doran, 2012-04-23 This welcome new edition discusses bioprocess engineering from the perspective of biology students It includes a great deal of new material and has been extensively revised and expanded These updates strengthen the book and maintain its position as the book of choice for senior undergraduates and graduates seeking to move from biochemistry microbiology molecular biology to bioprocess engineering All chapters thoroughly revised for current developments with over 200 pgs of new material including significant new content in Metabolic Engineering Sustainable Bioprocessing Membrane Filtration Turbulence and Impeller Design Downstream Processing Oxygen Transfer Systems Over 150 new problems and worked examples More than 100 new illustrations

Protein Chromatography Sinéad T Loughran, John Joseph Milne, 2023-08-30 This third edition expands on the previous editions with updated and new chapters on protein chromatography Chapters detail protein stability and storage avoiding proteolysis protein quantitation methods generation and purification of recombinant proteins recombinant antibody production and the tagging of proteins Written in the format of the highly successful *Methods in Molecular Biology* series each chapter includes an introduction to the topic lists necessary materials and reagents includes tips on troubleshooting and known pitfalls and step by step readily reproducible protocols Authoritative and cutting edge *Protein Chromatography Methods and Protocols* Third Edition aims to provide commonly used methods and new approaches to help both new researchers and experts expand their knowledge

Biotechnology and Biopharmaceutical Manufacturing, Processing, and Preservation Kenneth E. Avis, Vincent L. Wu, 2020-08-13 In this unique book experts describe practices applicable to the large scale processing of biotechnological products Beginning with processing and bulk storage preservation techniques the book provides strategies for improving efficiency of process campaigns of multiple products and manufacturing facilities for such processing techniques Large scale chromatography for the purification of biomolecules in manufacturing and lyophilization of protein pharmaceuticals are discussed Includes a case study on blow fill seal processing technology and a chapter on economic and cost factors for bioprocess engineering

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