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Renee Nemitz

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shelf life and sustainable packaging along with cost of environmental benefits to leverage savings on energy and carbon footprint In addition insight into safe processing of particulates using two and three dimensional thermal processing followed by prompt cooling is provided In the third edition the editors have compiled contemporary topics with information synthesized from internationally recognized authorities in their fields In addition to updated information 12 new chapters have been added in this latest release with content on Design of the aseptic processing system and thermal processing Thermal process equipment and technology for heating and cooling Flow and residence time distribution RTD for homogeneous and heterogeneous fluids Thermal process and optimization of aseptic processing containing solid particulates Aseptic filling and packaging equipment for retail products and food service Design of facility infrastructure and utilities Cleaning and sanitization for aseptic processing and packaging operations Microbiology of aseptically processed and packaged products Risk based analyses and methodologies Establishment of validated state for aseptic processing and packaging systems Quality and food safety management systems for aseptic and extended shelf life ESL manufacturing Computational and numerical models and simulations for aseptic processing Also there are seven new appendices on original patents examples of typical thermal process calculations and particulate studies single particle and multiple type particles and Food and Drug Administration FDA filing The three editors and 22 contributors to this volume have more than 250 years of combined experience encompassing manufacturing innovation in processing and packaging R D quality assurance and compliance Their insight provides a comprehensive update on this rapidly developing leading edge technology for the food processing industry. The future of aseptic processing and packaging of foods and beverages will be driven by customer facing convenience and taste use of current and new premium clean label natural ingredients use of multifactorial preservation or hurdle technology for maximizing product quality and sustainable packaging with claims and messaging 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 *A Manual for Hospital Central Services* Marie M. Lech, United States. Bureau of Health Planning and Resources Development, 1977

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