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technology; 3.3 Dry powder inhaler (DPI) formulation
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4.5.2 Controlled evaporation of droplets4.5.3 Evaporation of low-boiling-point solutions; 4.5.4 Spray freeze-drying; 4.6 Supercritical fluid (SCF) technology; 4.7 Conclusion; Acknowledgements; References; 5 Methods for understanding, controlling, predicting, and improving drug product performance; 5.1 Introduction; 5.1.1 The complexities and challenges of aerosol performance; 5.1.2 Understanding powder/particle characteristics: implications for aerosol product performance; 5.1.3 Liquid systems; 5.1.4 Summary; 5.2 Particle sizing; 5.2.1 Sieve analysis; 5.2.2 Image analysis; 5.2.3 Light scatter
5.2.4 Time-of-flight5.2.5 Other methods; 5.3 Powder and particulate characterization systems; 5.3.1 Introduction; 5.3.2 Powder cohesion and adhesion; 5.3.3 Microscopic material characterization; 5.3.4 Methods for studying bulk powders; 5.4 Practical issues in process control; 5.4.1 Common primary and secondary processing methods and issues arising for control; 5.5 Biopharmaceutical powder stability; 5.6 Liquids: solutions and suspensions; 5.6.1 Liquid formulation stability; 5.7 Conclusion; References
6 Aerodynamic assessment for inhalation products: fundamentals and current pharmacopoeial methods

Sommario/riassunto

"Provides students and those in industry with concise clear guide to the essential fundamentals in inhalation drug delivery"--Provided by publisher.