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Nota di contenuto	Management of Chemical and Biological Samples for Screening Applications; Contents; Preface; List of Contributors; 1 Introduction to Sample Management; References; 2 Generating a High-Quality Compound Collection; 2.1 Defining Current Screening Collections; 2.2 Design Criteria for Enriching a Compound Collection with Drug-Like Compounds; 2.2.1 Physicochemical Tailoring of a Compound Collection; 2.2.2 Lipophilicity Design Considerations; 2.2.3 Other Physicochemical Roadblocks; 2.2.4 Assessing Risk - from Rule of 5 to Rule of 3/75; 2.2.5 Tools Enabling Desk Top In Silico Design 2.3 Concluding RemarksReferences; 3 Assessing Compound Quality; 3.1 Introduction; 3.2 Process Quality and Analytical Quality in Compound Management; 3.2.1 Process Quality (QA); 3.2.2 Analytical Quality (Sample QC); 3.3 Identity; 3.4 Purity/Stability; 3.4.1 Measuring Purity; 3.4.2 Determining the Most Appropriate Purity Cut-Off for Solutions; 3.4.3 Stability of Solutions; 3.5 Concentration/Solubility; 3.6 Conclusions; Acknowledgments; References; Further Reading; 4 Delivering and Maintaining Quality within Compound Management; 4.1 Introduction 4.2 What is Quality from a Compound Management Perspective?4.3

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Sommario/riassunto

Filling an obvious gap in the scientific literature, this practice-oriented reference is the first to tie together the working knowledge of largescreening centers in the pharmaceutical and biotechnological field. It spans the entire fi eld of this emerging discipline, from compoundacquisition to collection optimization for specific purposes, to technology and quality control. In so doing, it applies two decades of expertise gathered by several large pharmaceutical companies to current and future challenges in high-throughput screening. With its treatment of libraries of small molecules