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Handbook of Stability Testing in Pharmaceutical Development: Regulations, Methodologies, and Best Practices is the first volume to cover all aspects of stability testing in pharmaceutical development. It presents a scientific understanding of regulations and balances methodologies and best practices.

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Handbook of Stability Testing in Pharmaceutical Development: Regulations, Methodologies, and Best Practices. In my professional career as a pharmaceutical scientist, I have been involved with several aspects of the drug development process from pre-IND to commercialization and, somehow, I usually found myself coming back to a stability-related issue. The stability area seemed to draw my utmost interest because in my day-to-day work, my opportunities involved more than one product, and no.

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Handbook of Stability Testing in Pharmaceutical

Handbook of Frequency Stability Analysis W.J. Riley Under contract with: Time and Frequency Division Physics Laboratory National Institute of Standards and Technology 325 Broadway Boulder, CO 80305 July 2008 U.S. Department of Commerce Carlos M. Gutierrez, Secretary National Institute of Standards and Technology James M. Turner, Deputy Director

Handbook of Frequency Stability Analysis

Abstract. An important aspect of all stability studies is the stability chambers themselves. This chapter is intended to provide a description of the different sizes and types of chambers that are available, the chamber tolerances required, and to provide some practical information for qualification, calibration, maintenance and monitoring of the chambers.

Qualification, Calibration, and Maintenance of Stability ...

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