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Regulatory 101: Bulk Holding Time Requirements

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Regulatory 101 provides a forum for pharma professions to share information about how they manage their respective quality and compliance responsibilities with regulatory considerations. Topics such as regulatory inspections, FDA-483s and Warning Letters, GMP manufacturing compliance topics, product technical management, and similar topics are planned for discussion -- all of which are supportive to the work of the quality and compliance function. Topics previously discussed in Regulatory 101 include the following: Environmental Excursions and Drug Product Stability. JGXP V23,...