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eLearning: Pharmaceuticals: Compliance and Audits

Art. ID	USP-RAPS-02
Unit	Each
Deliverydetails	No Dangerous Good

Description

Many agencies around the world are tasked with regulating the healthcare product industry within their respective countries. These agencies and their suppliers require manufacturers to conduct internal audits of their quality management systems on a regular basis to ensure compliance with the appropriate standards and regulations. Without an effective audit program, a company is at higher risk for nonconformance, regulatory actions, security breaches, poor product quality, loss of certification and registration, increased product liability risk, and a suboptimal process improvement system.