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ISO 11607-1:2006 Preview Packaging for terminally sterilized medical devices -- Part 1: Requirements for materials, sterile barrier systems and packaging systems This standard was last reviewed and confirmed in 2015.

ISO 11607-1:2006 - Packaging for terminally sterilized
ISO 11607 is the principal guidance document. Packaging for terminally sterilised medical devices - Part 1: Requirements for Materials, Sterile Barrier Systems and Packaging

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EN ISO 11607-1:2009 - Packaging for terminally sterilized
6 7 HALYARD* STERILISATION WRAP COMPLIANCE TO EN ISO 11607-1:2006 INTRODUCTION Dear Customer, In July 2014, the technical committee ISO/TC 198 (Sterilisation of health care products)

COMPLIANCE TO EN ISO 11607-1:2006/ AMD 1:2014
Katie Tran Presenter Â©2014, Westpak, Inc. 1 Greg Schwinghammer Moderator Review and Updates on Standardized Test Methods of ISO 11607 Herb Schueneman

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Packaging for terminally sterilized medical devices
11 Frequently Asked Questions about ISO 11607-1 ISO 11607-1 is the principal guidance document for validating terminally sterilized medical device packaging systems. Packaging must comply with ISO 11607-1 in order to satisfy European regulations and obtain a CE Mark. ISO 11607-1 is also a FDA Recognized Consensus Standard which is used in satisfying portions of device premarket review ...

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