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This part of EN 868 provides test methods and values for paper, used in the manufacture of paper bags (specified in EN 868-4) and in the manufacture of pouches and reels (specified in EN 868-5) used as sterile barrier systems and/or packaging systems that are intended to maintain sterility of terminally sterilized medical devices to the point of use.This part of EN 868 only introduces performance requirements and test methods that are specific to the products covered by this part of EN 868 ...

DIN EN 868-3:2017 - Packaging for terminally sterilized ...
DIN EN 868-3 Packaging for terminally sterilized medical devices - Part 3: Paper for use in the manufacture of paper bags (specified in EN 868-4) and in the manufacture of pouches and reels (specified in EN 868-5) - Requirements and test methods. standard by DIN-adopted European Standard, 05/01/2017. View all product details

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BS EN 868-3:2017 - Packaging for terminally sterilized ...
DIN EN 868-6 - 2017-05 Verpackungen für in der Endverpackung zu sterilisierende Medizinprodukte - Teil 6: Papier für Niedertemperatur-Sterilisationsverfahren - Anforderungen und Prüfverfahren; Deutsche Fassung EN 868-6:2017. Jetzt informieren!

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DIN EN 868-2 - 2017-05 Verpackungen für in der Endverpackung zu sterilisierende Medizinprodukte - Teil 2: Sterilisierverpackung - Anforderungen und Prüfverfahren; Deutsche Fassung EN 868-2:2017. Jetzt informieren! Damit wir unsere Webseiten nutzerfreundlicher gestalten und fortlaufend verbessern, verwenden wir Cookies. ...

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BS EN 868-3:2017 Packaging for terminally sterilized medical devices. Paper for use in the manufacture of paper bags (specified in EN 868-4) and in the manufacture of pouches and reels (specified in EN 868-5). Requirements and test methods BS EN 868-7:2017 - TC Tracked Changes. Packaging for terminally sterilized medical devices.

BS EN 868-2:2017 - Packaging for terminally sterilized ...
Anforderungen an Sterilbarrieresysteme werden in der DIN EN 868-2 bis -10 beschrieben und die Handhabung in der DIN 58953-7 bis -9. Sie dienen als Grundlage dieser Leitlinie, die als orientierungshilfe für die Durchfüh-rung der validierung in der Praxis zu verstehen ist.

Leitlinie für die Validierung der Verpackungsprozesse nach
DIN EN 868-5 - 2019-03 Verpackungen für in der Endverpackung zu sterilisierende Medizinprodukte - Teil 5: Siegelfähige Klarsichtbeutel und -schläuche aus porösen Materialien und Kunststoff-Verbundfolie - Anforderungen und Prüfverfahren; Deutsche Fassung EN 868-5:2018. Jetzt informieren!

DIN EN 868-5 - 2019-03 - Beuth.de
BS EN 868-5 provides examples of particular requirements and test methods for heat and self-sealable pouches and reels manufactured from paper complying with EN 868-3 and plastic film complying with clause 4 of this European standard.

BS EN 868-5:2009 - Packaging for terminally sterilized ...
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