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En 868 5 And Astm

- ASTM F2054 – Standard test method for burst testing of flexible package seals using internal air pressurization within restraining plates
- EN 868-5:1999, Annex D – Packaging materials and systems for medical devices which are to be sterilized –Part 5 Heat and self sealable pouches and reels of paper and plastic film construction –

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EN 868-5 is used to determine the peel strength of the heat seal joints used for heat and self-sealable pouches, as well as for reels used in packaging of medical devices that are sterilized. The EN 868-5 test method is suitable for use with a range of materials manufactured from paper and plastic film used in packaging.

EN 868-5 Packaging Materials Heat Self-Sealable Pouches

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BS EN 868-5:2018

EN868-5 – Annexe D. This test involves cutting a 15mm wide strip on the package and using a motorized tensile testing machine to separate the two sealed materials. The strip must be cut perpendicular to the seal. A sample of each packaging seal must be collected and tested at its center (see diagram below):

Peel tests on terminally sterilized medical packaging ...

The tensile test is carried out according to EN 868-5 and ASTM F88/F88M. During the tensile test, one or more strips of defined width (e.g. 15 mm) are cut out of the packaging at right angles to the sealing seam. A calibrated tensile testing machine is used to measure the force required to open the sealing.

Testing laboratory packaging tests/sterile barrier systems

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BS EN 868-5, 2019 Edition, January 31, 2019 - Packaging for terminally sterilized medical devices Part 5: Sealable pouches and reels of porous materials and plastic film construction - Requirements and test methods There is no abstract currently available for this document

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En 868 5 And Astm F88 - FPF Tech

ment and, as per en 868-5, must be great-er than or equal to 1.5 n/15 mm width14. if the maximum tensile strength of one of the three tests is less than 1.5 n/15 mm width, PQ is deemed to have failed. in addition the quality properties listed in iso 11607-2, § 5.3.2 b must be assured: – intact seal for a specified seal width

Guideline for Validation of Packaging Processes according

...
 EN 868-1:1997, which provided general requirements for packaging materials for sterile medical devices, was withdrawn and replaced by EN ISO 11607-1 in 2006. The remaining EN 868 standards, however, remain and revised editions have been published between 2017 and 2019. This series comprises Parts 2 to 10.

EN 868 series of European standards revised

Part 1 of the BS EN 868 series of European standards specifies general requirements and test methods for all packaging

materials and systems intended for use as packaging for medical devices which are to be terminally sterilized in their packaging. 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.

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11607-1:2006, BS EN

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This part of EN 868 provides test methods and values for sealable pouches and reels manufactured from porous materials complying with either EN 868 part 2, 3, 6, 7, 9 or 10 and plastic film complying with Clause 4 used as sterile barrier systems and/or packaging systems that are intended to maintain sterility of terminally sterilized medical ...

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