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If not properly planned for upfront, packaging can significantly delay timelines. Here is a breakdown of the design, sterilization, testing requirements, and validation processes medical device companies need to consider when tackling a packaging project. Package design considerations. Most medical devices are packaged either in a pouch or a tray.

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## **Medical packaging 101: Basics medical device companies**

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Thus, medical device manufacturer has to face with medical device packaging requirements to protect high-value products and must also comply with health and safety regulations. At this point, packaging validation requirements for medical devices are so important in order to comply with the approved standards for medical device packaging.

## **Which medical device packaging standards are there? - Safe ...**

Poor packaging can cause a host of problems for medical device companies and their customers. A package that holds a sterile medical device not only has to arrive at the hospital or clinic free of holes, tears and broken seals, it also has to withstand sitting on a shelf, possibly for years, without breaking down.



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MEDICAL DEVICE PACKAGING: INNOVATIONS IN DESIGN AND TESTING ... 2nd Annual Abhishek Gautam Manager, Packaging Engineering CONMED CORPORATION Catherine Olson, MSN, RN Director, Institute for Quality, Safety and Injury Prevention EMERGENCY NURSES ASSOCIATION Philip Desjardins Counsel

## **Medical Device Packaging: Innovations in design and testing**

Medical device packaging design is an integral part of delivering your device to market safely and securely, with the sterile barrier intact. Yet this critical process is often given little attention in the overall product development process. Most single-use sterile medical devices can be opened with a high degree of confidence that the ...

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## **3 steps for designing the ideal medical device packaging**

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The Medical Device Validation Handbook, Second Edition is a useful resource for regulatory and quality personnel, engineers, those involved in sterilization and anybody who works in manufacturing, says Sherman, and he stresses that anyone in these fields, regardless of experience level, can and should learn more about validation.

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