



Using Helium Leak Detection CCI Testing to Inform Container Closure System Design

A Prefilled Syringe Case Study

Container Closure Integrity: Regulations, Test Methods, Application Vienna, Austria; 9-10 November 2017

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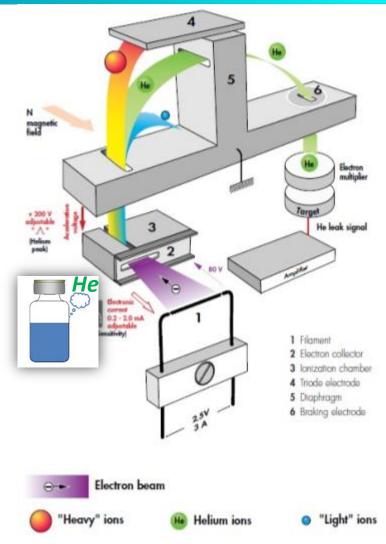
Overview of Helium Leak Detection (HeLD)

- Method development for syringes
 - Fixture design
 - Helium charging
- □ Applications of Helium Leak Detection
 - Determination of inherent package integrity
 - Evaluation of syringe system and sub-system design
 - Assessment of container closure robustness

Introduction – Helium Leak Detection



- Sensitivity & Quantitative
 - Mass spectrometer as detector
- Selectivity
 - Low atmospheric interference: Helium in the atmosphere (~5 ppm)
 - Do need to minimize lab ambient helium and permeation
- Flows through cracks ~2.7x faster than air



Leak Detection Associates, Blackwood, NJ





Calibration

- Internal calibration standards
- Quantitation Range: 1x10⁻¹¹ to 1x10⁻³ atm-cc/sec
- System suitability
 - NIST-treaceable standard leaks
 - Verified range: 4x10⁻¹⁰ to 6x10⁻⁴ atm-cc/sec
- Results reported as helium leak rate
 - Converted to a nominal leak orifice size using USP <1207> method, where appropriate





- Quantitatively determine inherent package integrity
 - The leakage rate (or the equivalent leak size) of a well-assembled package using nodefect components. Inherent package integrity is a measure of the leak tightness of a container–closure system, given anticipated variables of material composition, dimension, processing, assembly; package storage, distribution and use. (USP<1207>)
- Demonstrate conformance to Maximum Allowed Leakage Rate (MALL)
 - To preserve sterility and product formulation content

MALL <= $6x10^{-6}$ mbar·L/s (atm-cc/sec) (USP<1207>)

- Inform packaging system design and process development
 - Assess critical seal elements, sub-systems, and system design
 - Evaluate sealing robustness (e.g. impact of potential defects on package integrity)



Prefilled Syringe System

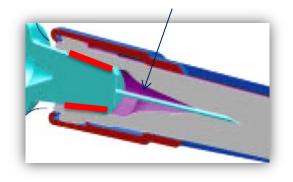


Drug Product Compartment

- Plunger-barrel seal
- Needle shield seal
 - Needle tip seal
 - Glued needle stem
 - Needle shield/syringe head

Needle Stem Compartment

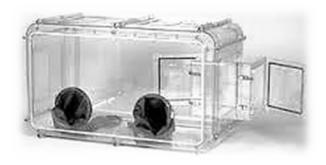
• Needle shield/syringe head



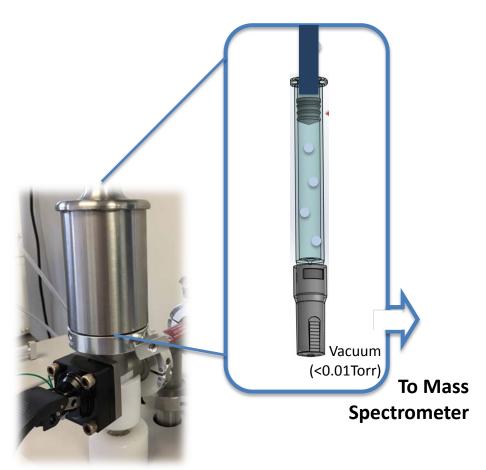




1. Filling: syringes charged with helium and plungered in a glove box



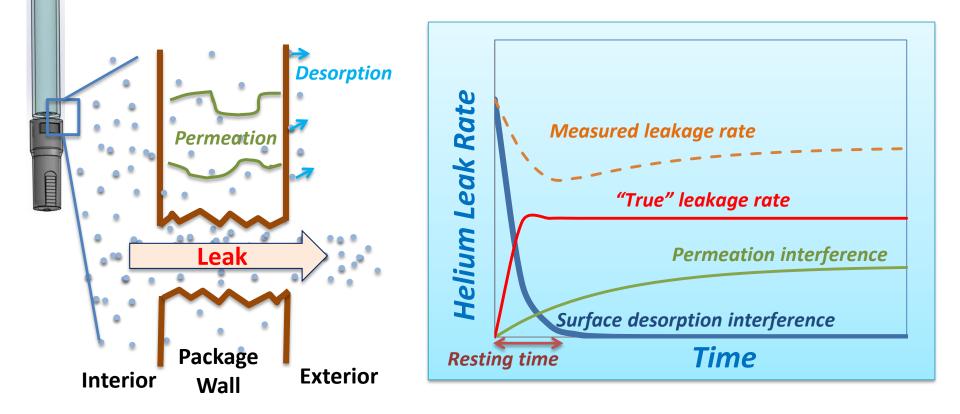
- 2. **Testing**: filled syringe is placed in a sample chamber for testing
 - A plunger rod is used to retain plunger during testing





Potential Interferences

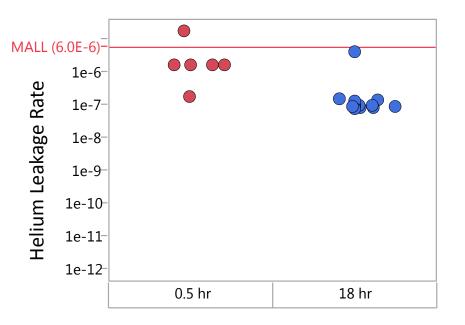
- **Desorption** of exterior surface-adsorbed helium
- Helium permeation





✓ Estimated overall leakage rate

- Results can be artificially high due to surface desorption and permeation
- ✓ Demonstrated conformance to MALL



Improvement Needs

- □ Reduce helium background noise to measure "true" leakage rate
 - Exterior surface desorption
 - Permeation

□ Need to evaluate sealing capability of critical seal elements and sub-systems

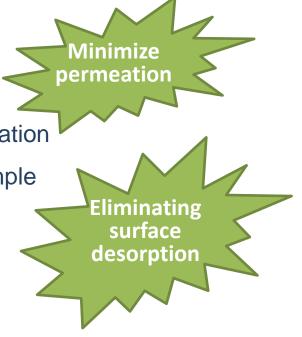
PDA Method Improvement Strategy

"Fast and Clean" helium charging

- Fast: allow testing to start prior to significant permeation
- Clean: eliminate helium contact with interfering sample surfaces

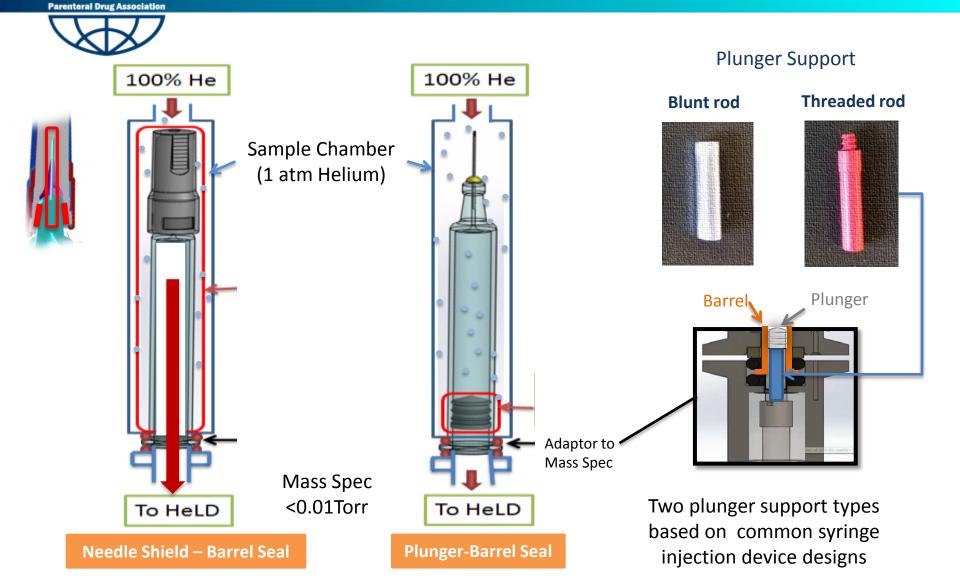
Divide and Conquer" critical seal elements

Isolate critical seal elements for independent
 assessment

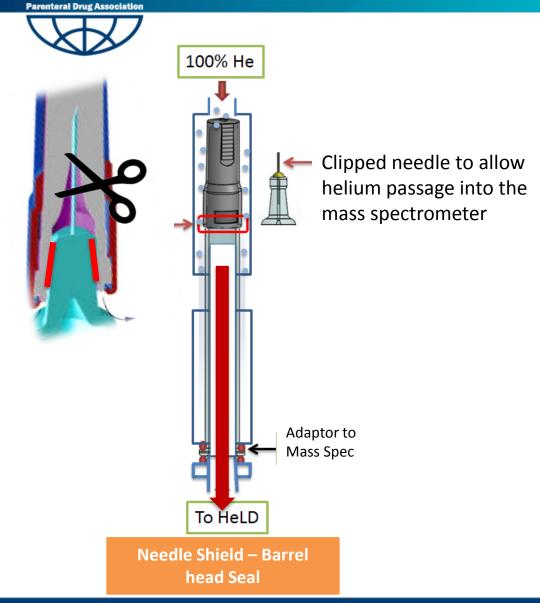


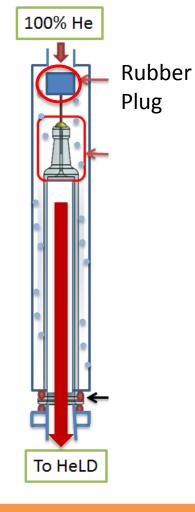
Key Enabler: Sample Fixture Design

PDA Testing Fixture Design



PDA^{*} Testing Fixture Design

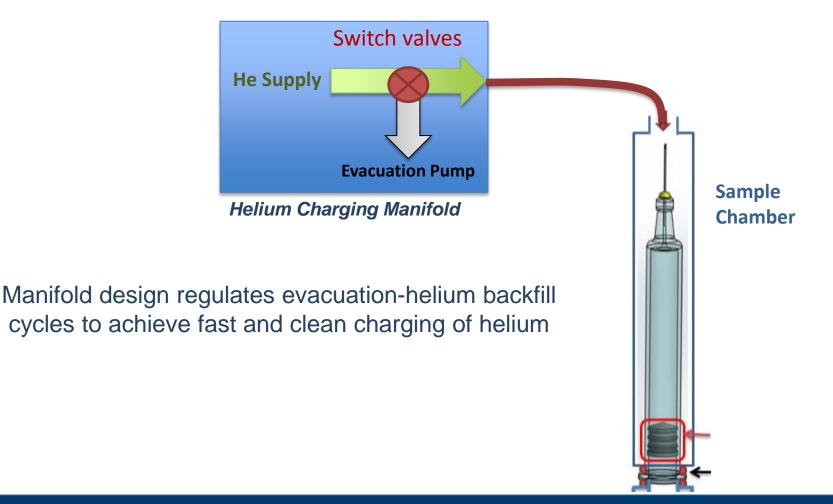




Needle Stem Seals

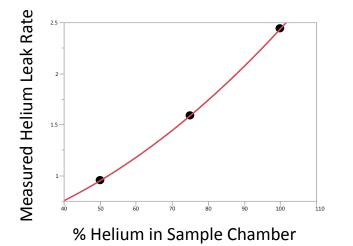


chamber to allow accurate quantitative leakage rate testing



PDA Pronteral Drug Association Optimize Helium Charging Parameters

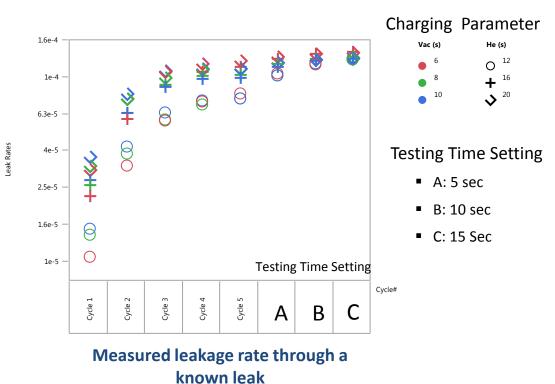
Leakage Rates dependence on He% in Sample Chamber



Key Parameters

- Charging cycle
 - Evacuation time (sec)
 - He back-fill time (sec)
- Number of Cycles
- Post-charging testing time (sec)

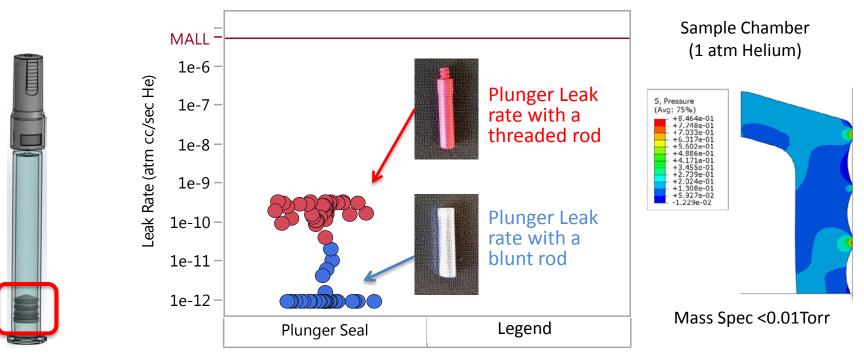
Leakage Rates vs. Charging Parameters Study



DA Evaluation of Plunger-Barrel Seal

Physically mated (compression) seal

- Critical to product sterility and formulation content protection



- Leakage rate <= 4 x 10⁻¹⁰ atm-cc/sec
 - Confirm to MALL for preserving sterility and product formulation content
- Plunger rod selection affects plunger seal during testing
 - In agreement with FEA modeling results

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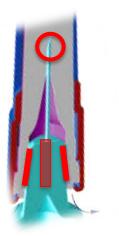
Parenteral Drug Associati

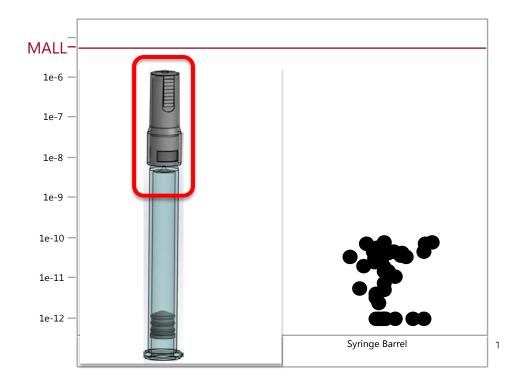
Evaluation of Needle Shield – Barrel Seal



A combination of 3 seal elements

- Needle shield Syringe head
- Needle tip Needle shield
- Glued staked needle
- Rate (atm cc/sec He) Critical to product sterility and formulation content protection

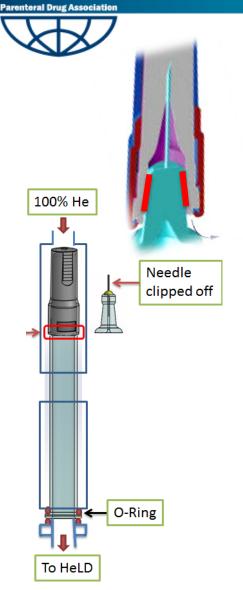




Leakage Rate <= 8 x 10⁻¹¹ atm-cc/sec

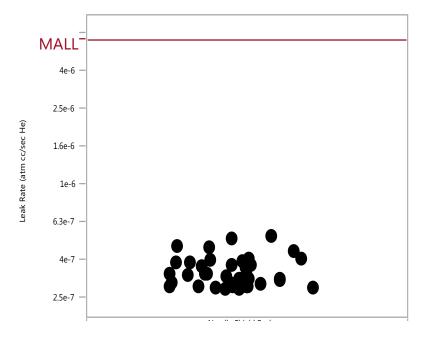
Confirm to MALL for preserving sterility and product formulation content

PDA Evaluation of Needle Shield – Barrel Head Seal



Physically mated (compression) seal

- Critical to needle stem sterility protection
- Contributes to DP compartment seal



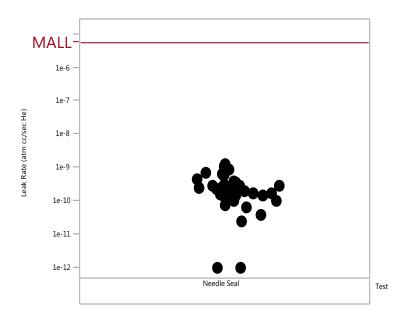
Leakage Rate <= 6x10⁻⁷ atm-cc/sec

Confirm to MALL for preserving sterility



Evaluation of Needle Stem Seals

- Glued (physicochemically bonded) at the base
- Physically mated (compression) seal at the tip
 - Not definitive sterility barriers
 - Poor seals may result to product loss or injection issues



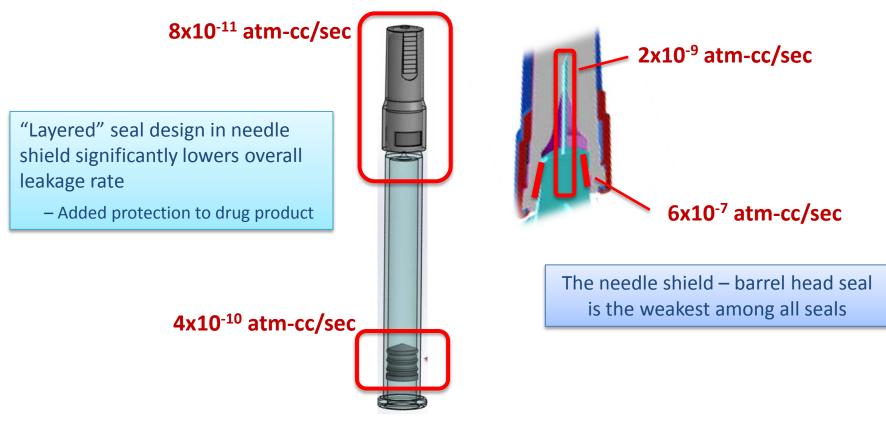
Leakage Rate <= 2x10⁻⁹ atm-cc/sec

 Confirm to MALL for preserving sterility and product formulation content



Inherent Package Integrity Summary

- DP compartment (Plunger & Needle Shield) <= 5 x 10⁻¹⁰ atm-cc/sec
- Needle stem compartment <= 6 x 10⁻⁷ atm-cc/sec
- All individual critical seals conform to MALL



Package Integrity was verified for preserving sterility and formulation content



Leak Rate (atm cc/sec He)

Design Robustness: Plunger Ribs

Each plunger rib assessed individually for sealing capability

- Assessed by compromising 2 of the 3 ribs, leaving 1 intact rib
- Evaluate impact of potential plunger molding defects

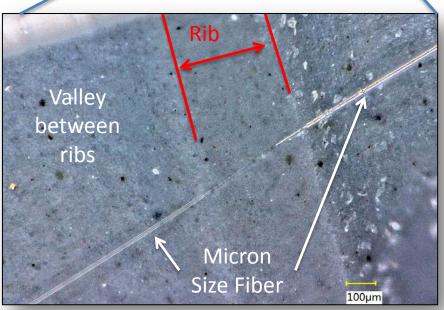


PDA Design Robustness: Fiber Interfernce

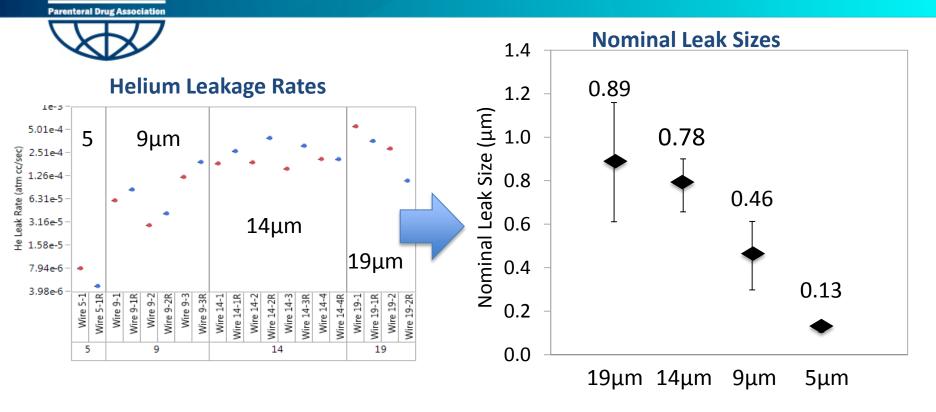




- Industry needs a practical means to fabricate and characterize sub-micron size defects
- Assess impact of interfering fibers of various sizes



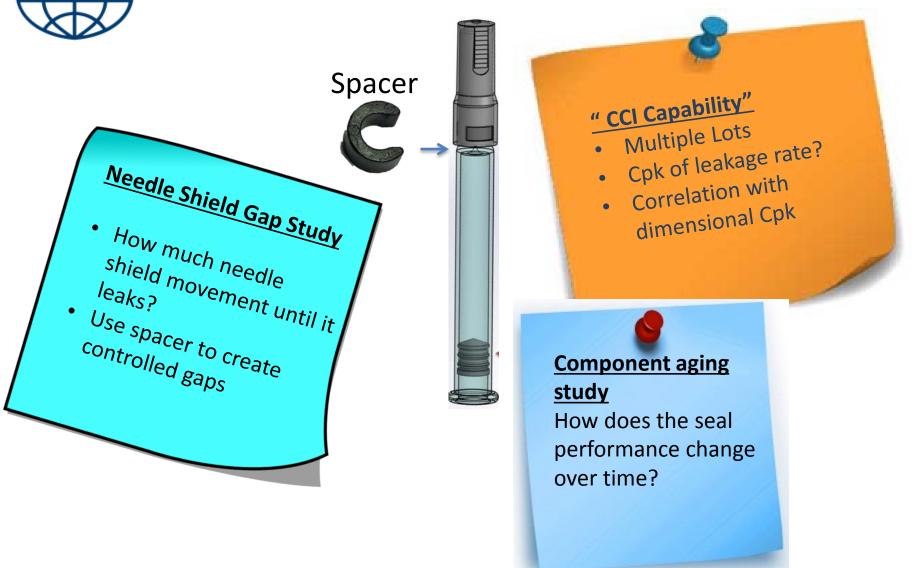
Design Robustness: Fiber Interfernce



- Presence of interfering fibers could compromise plunger seal integrity
- Interfering fibers provide a practical means for fabricating sub-micron defects
 - Enable development of other CCI testing technologies

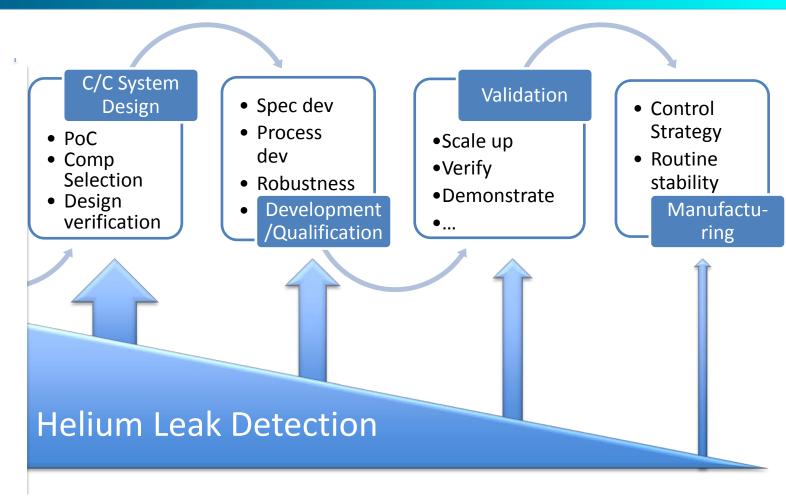


Potential Opportunities

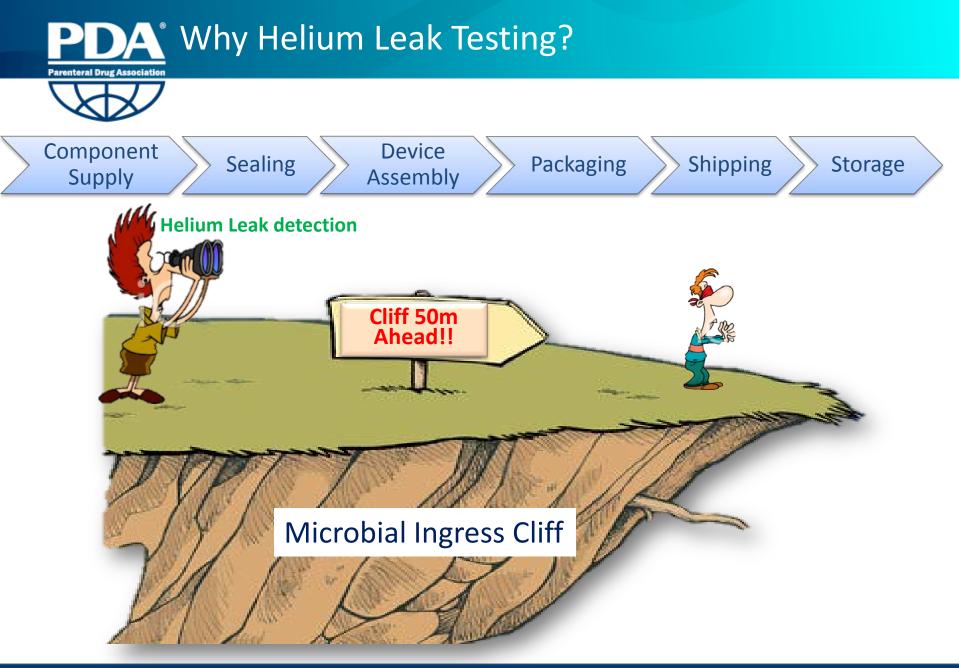


PDA Application of Helium Leak Testing





- Sensitive and quantitative testing to inform design & development
- Use in routine manufacturing DP filled container may be limited



PDA Parenteral Drug Association Conclusion: Helium Leak Detection

□ HeLD a capable CCI technology: sensitive, precise, and quantitative

- Enable data-driven decision making: rich information; high throughput
- Fixture design and helium charging critical for method development

□ Inform container closure design and process development

- Inform packaging component selection and system design
- Assess container closure robustness against design, process variability
- Demonstrate conformance to MALL
- Enable foundational CCI research
 - Characterize micron and sub-micron leaks



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