

How terminal sterilization puts quality and safety first



With a proprietary process that combines aseptic processing with terminal sterilization—a proven pharmaceutical industry standard—we are able to achieve unprecedented sterility assurance and extended Beyond-Use Date (BUD).

COMBINING TWO STERILIZATION TECHNIQUES

Aseptic processing: A sterilization technique used by 503A compounding pharmacies to produce sterile drug products.



Used to sterilize the patient-specific prescriptions dispensed by AIS Healthcare Targeted Drug Delivery



Multiple 0.22-micron sterilizing-grade filtration steps to ensure medication sterility, going beyond USP guidelines for preparing sterile solutions



Goal is to remove any potential contaminants

Aseptic processing provides a sterility assurance level of 1:1,000.

Terminal sterilization: A sterilization technique used by AIS Healthcare Targeted Drug Delivery to ensure the highest level of sterility assurance.



Used in addition to aseptic processing to further sterilize patient-specific prescriptions dispensed by AIS Healthcare Targeted Drug Delivery



A proven sterilization technique that is the pharmaceutical industry standard and the preferred method for the commercial preparation of injectable medications



Our validated, proprietary heat and steam terminal sterilization cycles are used to eradicate all theoretical bioburdens without compromising medication potency or purity

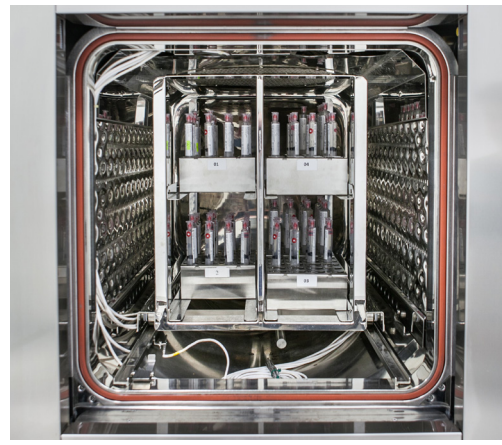


Because the terminal sterilization process happens after the medication is in the syringe, there's no risk of further contamination from human handling post-sterilization



Goal is the destruction of all theoretical bioburdens

Terminal sterilization provides a sterility assurance level of 1:1,000,000



Take a tour

Contact us to tour our industry-leading compounding pharmacies in Dallas, Texas (headquarters), or Ridgeland, Mississippi, and see how our proprietary process can benefit you and your patients.

Can't visit in person?

Take our virtual tour at tour.aishealthcarepharmacy.com.



aiscaregroup.com | 877.443.4006

¹ All patient-specific formulary dispenses within maximum concentrations (based on AIS Healthcare in-house data for 120,000 aseptically processed and terminally sterilized patient-specific prescriptions in 2019).

² USP-NF: https://online.uspnf.com/uspnf/document/1_GUID-08D442A0-208E-4DCE-98BE-21C84C2D48AA_1_en-US?source= Accessed March 7, 2020.

³ U.S. Food and Drug Administration. Summary of Recent Findings Related to Glass Delamination. November 2018. <http://www.fda.gov/drugs/pharmaceutical-quality-resources/summary-recent-findings-related-glass-delamination>. Accessed March 7, 2020.

⁴ Iacocca RG, Totti N, Allgeier M, et al. Factors Affecting the Chemical Durability of Glass Used in the Pharmaceutical Industry. *AAPS PharmSciTech*. 2010;11(3):1340-1349. doi:10.1208/s12249-010-9506-9. Accessed March 7, 2020.

⁵ U.S. Food and Drug Administration. FDA Form 483 Frequently Asked Questions. January 2020. [https://www.fda.gov/inspections-compliance-enforcement-and-criminal-investigations/inspection-references/fda-form-483-frequently-asked-questions#:~:text=A%3A%20An%20FDA%20Form%20483,FD%26C\)%20Act%20and%20related%20Acts](https://www.fda.gov/inspections-compliance-enforcement-and-criminal-investigations/inspection-references/fda-form-483-frequently-asked-questions#:~:text=A%3A%20An%20FDA%20Form%20483,FD%26C)%20Act%20and%20related%20Acts). Accessed July 6, 2020.

How AIS Healthcare Targeted Drug Delivery achieves unprecedented sterility assurance and extended BUD

Advancing quality through terminal sterilization



Advancing quality. Improving lives.



21-45
DAY BUD AT ROOM
TEMPERATURE¹


By following aseptic processing with terminal sterilization, AIS Healthcare Targeted Drug Delivery achieves sterility assurance levels of 1:1,000,000 and BUD of 21–45 days (average of 34 days).¹

Dedicated to a better terminal sterilization process

AIS Healthcare goes above and beyond standard terminal sterilization. Our industry-leading process and equipment reflect our dedication to do more to advance quality and patient safety in every prescription we deliver.


The AIS Healthcare difference	Why it matters	Did you know?
Significant investment	AIS Healthcare has made a multimillion-dollar, multi-year investment in our terminal sterilization equipment and processes.	USP <797> standards do not require terminal sterilization. We not only use terminal sterilization but have also made a significant investment in our equipment and processes, reflecting our long-term commitment to raising the bar for quality and sterility and doing what's right, not just what's expected.
Specialty designed	Our custom, 1.5-ton autoclaves were designed in collaboration with industry experts and built by the world leader in the production of solutions for sterile processes and equipment in the pharmaceutical sector.	Our custom autoclaves allow our pharmacists control over critical parameter settings, including temperature, pressure, counter pressure and time, to eradicate all theoretical bioburdens in the syringes.
Proprietary process	We use a deeply researched and proprietary terminal sterilization process, developed in conjunction with industry experts, for patient-specific prescriptions.	This proprietary process applies the right temperature, pressure and time parameters to the syringes, effectively leveraging steam heat sterilization to achieve the highest levels of sterility assurance.
Extensively studied	We have conducted and continue to conduct rigorous and independent testing and validation studies, including extractable and leachable studies, of our terminal sterilization program.	These ongoing studies prove that our terminal sterilization process effectively eradicates contaminants that may have been introduced into our medications.
Lower heat	Our terminal sterilization process uses steam heat cycles with temperatures that peak well below the conventional temperatures associated with autoclaving.	Lower temperatures and our proprietary terminal sterilization processes have been proven to effectively sterilize while reducing any potential damage to the medication during terminal sterilization.
Enhanced convenience and safety	Prescriptions are terminally sterilized in plastic syringes, which are then packaged and shipped directly to providers.	Delivering the medication in a syringe, which is the final container for each patient-specific prescription, instead of glass vials allows for fewer manipulations by the clinic and clinic staff, decreasing the possibility of introducing potential contaminants. This also eliminates the need for additional training on safely transferring the medication from a vial to a syringe.
Pharmacists only	All patient-specific prescriptions are only compounded by licensed pharmacists to ensure every prescription meets our industry-leading quality standards.	Licensed pharmacists have additional expertise and experience and are the most qualified to compound medications of the highest quality.
Inspected and accredited	We are regularly inspected by the various state boards of pharmacy, the DEA and the FDA (with no 483 observations on last inspection). We are also dually accredited by URAC and the Accreditation Commission for Health Care (ACHC). Our Ridgeland, Mississippi, facility received the first URAC accreditation for Home Infusion Therapy Supplier (HITS).	The absence of 483 observations and our dual accreditations show our commitment to quality and dedication to producing products under the highest possible standards.






“I call AIS Healthcare my poster child of how to do it right.”

Eric Kastango
President of Clinical IQ and CriticalPoint



ACCREDITED
Specialty Pharmacy
11/01/2022



Our compounding process

Receipt and scheduling	Customer service coordinates order intake and schedules orders for shipping.
Entry	AIS Healthcare Targeted Drug Delivery uses double-blind order entry and verification. All orders are entered and reviewed by two pharmacy technicians.
Verification and reverification	Two pharmacists then do a blind review of the order before the medication is compounded. This means that multiple licensed professionals have eyes on every order to reduce the chances of any human error.
API procurement and stock solution preparation	All USP-grade active pharmaceutical ingredients are sourced from FDA-registered manufacturers, and we obtain certificates of analysis from both manufacturers and distributors. All of our stock solutions are prepared by a licensed pharmacist and tested by an outside, independent lab for sterility, potency, particulate matter, pH and endotoxins before use in compounding.
Aseptic processing	Prescriptions are compounded from sterile stock solutions in our state-of-the-art cleanroom. The pharmacist uses the compounding record to select the correct inventory items, and a pharmacy technician confirms the selection twice. The prescriptions are then put into previously sterilized syringes.
Terminal sterilization prep	After aseptic processing, a pharmacist visually inspects each syringe and then passes it to a pharmacy technician for labeling. A terminal sterilization label is put on each syringe and a bubble test is performed by a technician as a second visual inspection. Then, the syringes are loaded into the autoclave for the terminal sterilization cycle.
Terminal sterilization	In the autoclave, each syringe gets enough heat and steam to kill all theoretical bioburdens that may be present in that environment. Once terminal sterilization is complete, parametric release is achieved and the syringes are unloaded from the steam autoclave. Another check is then performed. No further stability or sterility testing is required.
Staging	An AIS Healthcare technician stages every labeled syringe, and a pharmacist inspects each syringe for any issues, matches it with the appropriate shipping packaging and double-checks the volume.
Packaging	AIS Healthcare pharmacists package each prescription into an amber bag for light protection and then seal it in a shipping tube that can be surrounded by bubble wrap. Another check is then performed to ensure the packing label reflects the packing slip.
Shipping	The shipping tube is then placed in an International Safe Transit Association–certified box with tamper-evident tape and other proprietary safety precautions. We ship via first-class overnight air at no additional cost, which means patients across the country get their medication the next morning. We also monitor potential weather-related delays and proactively adjust when and how we ship medication, including shipping from our redundancy operations or using another shipping route, to make sure patients' prescriptions arrive on time.