

# Setting new quality standards for targeted drug delivery

#### AIS Healthcare is a leader in sterility assurance and extended BUD

At our two 503A compounding pharmacies, AIS Healthcare goes above and beyond to advance medication quality. With a proven, proprietary process that combines aseptic processing with terminal sterilization—a pharmaceutical industry standard that only AIS Healthcare uses—we are able to achieve unprecedented sterility assurance and extended BUDs.



of all stock solutions1



21-45

DAY BUD AT ROOM TEMPERATURE

for all formulary medications<sup>2</sup>



for unprecedented quality and patient safety (versus 1:1,000 sterility assurance levels with aseptic processing alone)

#### Maximum concentrations for extended-BUD stock solutions

MEDICATION	MAXIMUM CONCENTRATIONS	AIS STOCK SOLUTION 3RD-PARTY TESTED <sup>1</sup>	ASEPTICALLY PROCESSED	TERMINALLY STERILIZED <sup>3</sup>	BUD
Baclofen <sup>4</sup>	4,000 mcg/mL	✓	<b>~</b>	<b>~</b>	21-45 DAYS AT RT AVG.=34 DAYS
Bupivacaine <sup>5</sup>	40 mg/mL	✓	<b>~</b>	<b>~</b>	
Clonidine	5,000 mcg/mL	✓	<b>~</b>	<b>~</b>	
Fentanyl citrate	10,000 mcg/mL	✓	<b>~</b>	<b>~</b>	
Hydromorphone	100 mg/mL	✓	<b>~</b>	<b>~</b>	
Morphine sulfate	62.5 mg/mL	✓	<b>~</b>	<b>~</b>	
Sufentanil	1,000 mcg/mL	✓	<b>~</b>	<b>~</b>	

These medications account for 97% of all AIS Healthcare syringe dispenses, including trials. For more information about trial medications, ask your representative.

BUD that is never less than 21 days<sup>2</sup>—just one of the ways AIS Healthcare does more to support patients and providers.



### More medications available from AIS Healthcare

Ask your AIS Healthcare representative about other medications.

MEDICATION	ASEPTICALLY PROCESSED	BUD
Gabapentin	<b>~</b>	
Ketamine	<b>✓</b>	
Lidocaine	<b>✓</b>	24
Magnesium sulfate	<b>✓</b>	HOURS AT RT
Meperidine	<b>✓</b>	711111
Ropivacaine	<b>✓</b>	
Prialt	<b>~</b>	Combo: 3 days refrigerated Mono: 9 days refrigerated

AIS Healthcare is dually accredited by URAC and the Accreditation Commission for Health Care (ACHC).





Specialty Pharmacy 11/01/2022

These medications account for only 3% of all AIS Healthcare syringe dispenses.

## Exceeding industry standards

AIS Healthcare goes above and beyond USP requirements, doing more to put patient safety first.

AIS HEALTHCARE QUALITY STANDARD	USP <797> GUIDELINES	
All stock solutions are prepared by a licensed pharmacist and tested by an outside, FDA-registered lab for sterility, potency, particulate matter, pH and endotoxins before use in compounding	Not required	
Compounding done only by licensed pharmacists	Compounder not required to be licensed pharmacist	
Enhanced contamination control procedures, including full-body sterile garb for cleanroom personnel	Only sterile gloves required for cleanroom personnel	
Multiple 0.22-micron sterilizing-grade filtration steps, including filter integrity testing, to ensure medication sterility	Only one 0.22-micron sterilizing-grade filtration step required	
Four independent cleanroom certifications per year	Only two required per year	
Non-viable particle count done 12 times per year, combined viable air and surface sampling done no fewer than 50 times per year	Non-viable particle count and viable air sampling required twice per year; surface sampling required periodically	

Discover unmatched quality and patient safety. Get started with AIS Healthcare today. aiscaregroup.com | 877.443.4006

<sup>&</sup>lt;sup>1</sup> Third-party testing for pH, sterility, potency, endotoxins and particulate matter.

<sup>2</sup> All patient-specific dispenses within maximum concentrations (based on AIS Healthcare in-house data for 120,000 aseptically processed and terminally sterilized patient-specific prescriptions in 2019).

<sup>3</sup> Only patient-specific dispenses are terminally sterilized.

<sup>4</sup> FDA regulations require baclofen to be compounded at a concentration above or below 10% of the commercially available concentration of Gablofen® and Lioresal®.