

CIAL | How to read Price List

1. Please visit the [CIAL Website](#) and download the current **Price List**.
2. Tests are grouped by the department performing the testing.
 - a. Microbiology Testing is in **Blue**, pg 1-2
 - b. Chemistry Testing is in **Green**, pg 2-3
 - c. Special Chemistry Testing is in **Cream**, pg 3-4

Test Name	Test Code	USP	Description	Turnaround (bus. days)	Amount of Sample	Price	Notes
Microbiology Testing							
Antimicrobial Effectiveness	AET	<51>	Determine if the antimicrobial agent in the formulation is effective. Required for <795> <797> multi-dose formulations.	40	55 mL	\$1,200	Requires 35 day incubation.
Endotoxin	END	<85>	Determines levels of bacterial endotoxin in finished product. Required per <797> 12.3 when Sterility is performed.	3	1 container	\$135	If raw material (API), quote required.
Chemistry Testing							
Cleaning Validation	CLV	<800>	Tests for residue of API's remaining on compounding surfaces after cleaning procedure.	7	(Quote)	(Quote)	Contact CIAL for cleaning procedure. 1 swab per surface and 4 blank swabs.
Content Uniformity	TMB	<3>	Tests for consistency in typically applied products. Commonly referred to as "Top, Middle, Bottom" testing.	7	1 container	Starting at \$585	3 x respective Potency.
Special Chemistry Testing							
Appearance	AOC	N/A	Documents the physical condition of formula over time.	7	1 container	\$70	Protocol required.
Container Closure Integrity	CCI	<1207.2>	Tests the integrity of the container closure system by dye intrusion technique. Results based on 2 containers.	7	4 containers	\$175	Verified by <381>. Additional +\$20 per container.

3. Locate the Test type that you are looking to request in the *Test Name* section.
 - a. The following boxes are:
 - i. Test Code
 - ii. Any correlating USP Chapter
 - iii. A description of the testing
 - iv. Turnaround time
 - v. The amount of sample needed for the test
 - vi. Price
 - vii. Any special notes about the test or testing parameters
4. When the statement "Quote" is accompanied in any of the boxes, please reach out to lab@compounderslab.com with the appropriate information:
 - a. Phase 1 (SIA) Method Development & Validation
 - i. Formulation ID, if available
 - ii. Active(s) concentration
 - iii. Container size/volume
 - iv. Excipient list (MFR is ideal)
 - b. Phase 2 (BUD) Stability / Potency Over Time study
 - i. Target BUD or final time point
 - ii. Storage condition
 - iii. Test(s) to be performed and time point intervals
 - c. Compendial / Monograph Testing
 - i. Manufacturer CoA
 1. Lot number, if applicable
 - ii. Test(s) to be performed
 1. Acceptance criteria, if no USP monograph is available