imprimis Rx®

WHO WOULD YOU TRUST IN PROVIDING MEDICATIONS FOR YOUR PATIENTS?

Your patients are important to us and we want them to receive the highest quality medications. Don't trust your patient's safety with just anyone. You and your patients deserve high quality medications made to standards that are required to meet applicable USP Chapter <797> and cGMP standards.

ImprimisRx can provide you with the **correct** answers to all of the questions below.

10 IMPORTANT QUESTIONS YOU SHOULD ASK WHEN CHOOSING A PHARMACY OR SUPPLIER

1. Where are your formulations made? Does your company own and manage these facilities?
2. How long has your pharmacy been making these formulations? Have they served over **1 million** patient eyes?
3. Are all of the formulations tested in an FDA-registered laboratory?
4. Is the laboratory testing every batch of your formulations for **potency**, **sterility**, and **endotoxins**? Can you provide me a report for all of these with every shipment I receive?
5. Is beyond use date (BUD) determined using validated stability indicating methods and data from an FDA-registered laboratory that follows cGMP guidelines?
6. Does your laboratory and company test the active pharmaceutical ingredient (API) raw material for potency before making new batches?
7. Is your supplier using validated warehouses to store finished product?
8. Can I order and store inventory of your product on my shelf to provide to patients when they need them?
9. Does your supplier follow **cGMP requirements** in making and testing all of your formulations?
10. Does your pharmacy participate in MedWatch: The FDA Safety Information and Adverse Event Reporting Program?

GO DROP LESS®

LESS DROPS®

Imprimis delivers high-quality formulations following FDA's current good manufacturing practice regulations and guidance. Our 503B outsourcing facility is FDA-registered and required to meet cGMP guidelines to safeguard the quality and safety of every compounded medication dispensed from our facilities.

IMPRIMIS PROVIDES STERILE COMPOUNDED FORMULATIONS YOU CAN TRUST

- Our cGMP facility is inspected by the FDA
- We use bulk active and inactive substances obtained from FDA-registered manufacturers, and accompanied by a certificate of analysis
- Release and stability testing performed at FDA-registered labs
- Sterility, endotoxin and potency testing performed on each batch
- Continuous environmental monitoring to maintain product quality standards

THE IMPRIMIS COMMITMENT

Quality compounded formulations brought to you only by Imprimis

	503A Pharmacy	503B Outsourcing Facility
API TESTING / RAW MATERIALS	No requirement	Identity test for each incoming lot
POTENCY TESTING	No requirement	All finished product lots
STERILITY TESTING	Sterile lots per USP <71>	Provide results for sterile lots per USP <71>
ENDOTOXIN TESTING	Sterile Injectable lots per USP <85>	Sterile Injectable lots per USP <85>
PRE-SHIPMENT QUARANTINE	Not required, but recommended	14 days for sterility result
ENVIRONMENTAL TESTING	Every 6 months	Continuous (24 hours/day; 7 days/week)
TEST RESULTS INCLUDED WITH ORDER	No requirement	Sterility Results Endotoxin Results
BEYOND USE DATING (EXPIRATION)	Literature and pharmacist experience based, but not required. Stability study over time recommended.	Stability study data from three lots using validated stability indicating methods

DYNALABS Continuous Quality Improvement (CQI) Center of Excellence

To contact an Imprimis Pharmacist, call
844-4IMMYRX (844-446-6979)
or email **info@imprimisrx.com** today!

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