Why Choose a 503B Outsourcing Facility?
• Products do not require a patient specific prescription.
• Required by the FDA to follow strict testing standards for sterility, potency, and endotoxins.
• Products are compounded in ISO clean rooms that are continuously monitored for particulate matter and environmental conditions.
• Ability to provide drugs on the FDA Short List.
• Must comply with current good manufacturing practice (cGMP) requirements and U.S. Pharmacopeia Guidelines (USP) and follow state regulations and voluntarily register with the FDA.

Why Choose RXQ Compounding, LLC?
• Highest quality sterile and non-sterile manufacturing.
• ISO 5 clean room sterile production.
• Safety, availability, and compliance.
• Integrated delivery network and ability to deliver custom compounding.
• 50,000 sq. ft. future production facility.