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FDA Registered 503B Outsourcing Facility by Cantrell Drug Company

Our experience with cGMP focused policies and procedures; along with carefully designed facilities, equipment, training, and testing; provide confidence to our customers in the quality of our services.

CEO:	Dell McCarley, Pharm.D.
President:	Mike Pierce
Founded:	1952
Employees:	150+
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Company Background

Established in 1952, Cantrell Drug Company has a long and successful history. Amidst many changes in the field of pharmacy, Cantrell Drug Company has remained committed to the highest standard of quality. Innovation and investment in cutting-edge technology at Cantrell Drug Company is evident by being the first in the industry to utilize a rapid microbiological sterility testing program. Our commitment to quality is readily apparent from a completely autonomous and independent Quality Assurance Department and one of the first sterile compounders to register as a 503B with the FDA.

Product Overview

Cantrell Drug Company is an FDA registered 503B Human Drug Compounding Outsourcing Facility that specializes in providing customized IVs, OR syringes, epidural preparations, PCA syringes, and drug shortage solutions with the highest level of quality. Product lines are compounded under a comprehensive quality system model that supports and sustains robust quality systems consistent with cGMP-focused regulations. Cantrell provides sterile and non-sterile compounded preparations that meet the needs of patients, physicians, clinics, and health care institutions. Cantrell retains a DEA manufacturing license, an FDA registration, and is appropriately licensed in all 50 states. When you need Quality, Service, and Experience — Trust Cantrell Drug Company.

Drug Shortage Solutions

Drug shortages lead to increased cost and procurement challenges for pharmacies to meet patient and physician needs, and much of the financial impact is felt in acute care settings. Cantrell Drug Company offers compounded solutions to many of these shortages. We give our clients a way to “fill the gap” during their time of need for hard to find preparations.

Cantrell Drug Compounds From Active Pharmaceutical Ingredients (API) Whenever Possible for a Number of Reasons

■ Drug Shortage Insulation

Utilizing API helps ensure we will be able to supply our customers even when commercial products cannot be sourced.

■ Commercial Price Insulation

The commercial drug market is in a constant state of flux. By utilizing API we are able to keep our costs consistent so our clients can better plan their pharmacy spending.

■ Reduced Bioburden in the Hood

API allows us to prepare a single bag and limit the number of non-sterile commercial containers in our clean room area thus reducing the bioburden. Cantrell Drug prepares a single bag vs. hundreds of vials in the class five environment.

■ Fewer Controlled Movements in the Hood

API allows a more efficient process, removing the need of pooling commercial vials. Fewer manipulations saves time and operator fatigue.

