

Cantrell Drug Relocates to State-of-the-Art Facility

Cantrell Drug Company, originally opened in 1955, has recently built and relocated to a new, state-of-the-art facility, boasting over 18,000 sq. ft.

The new facility offers expanded office, production, quality control and warehouse space. It is a current Good Manufacturing Practice (c-GMP) compliant facility, employs advanced “smart technology” monitoring of critical parameters,” and boasts a 5,000 sq. ft. Iso7 clean room.

“This modern facility provides us with the needed production capacity, quality controls and testing space to keep pace with the demands for compounding sterile preparations and most any dose form,” stated the company president, Dell McCarley. Cantrell Drug provides the highest quality medications within the industry’s strict guidelines. A FDA registration is pending for this new facility. Cantrell Drug Company is also accredited by the Pharmacy Compounding Accreditation Board. Cantrell Drug is well positioned to become one of the nation’s most advanced compounding companies.

Cantrell Drug Company, a national leader in interventional pain management and high-risk sterile products, provides customized pain medications, admixture services, research preparations and other custom compounding services to patients, physicians, clinics and healthcare institutions.

For more information, contact Dell McCarley at 877-666-5222, or send e-mail inquiries to llanier@cantrelldrug.com

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