

February 17, 2021

Drug Delivery Experts Obtains FDB License for Manufacture of Sterile Injectable Clinical Trial Materials

San Diego, California – Drug Delivery Experts’ (DDE Labs) cGMP manufacturing suite and quality system was inspected and approved by the California Department of Public Health Food and Drug Branch (FDB), permitting DDE Labs to manufacture and ship pharmaceutical products.

The FDB completed a comprehensive cGMP inspection of the facility with no observations and awarded DDE Labs a license for drug product manufacturing.

The fill/finish suite complements DDE Labs array of services in formulation, drug delivery, and drug product development; providing clients a seamless transition to clinical manufacturing for their products.

“This is a tremendous step. We look forward to helping our clients move efficiently from our drug product development support to quick evaluation of their products in the clinic,” said Chris Rhodes, CEO of DDE Labs.



ABOUT DRUG DELIVERY EXPERTS: Drug Delivery Experts (DDE Labs) is a drug product research and development company that specializes in drug delivery systems from preclinical to clinical development and commercialization. DDE Labs has specific expertise with peptides, proteins, and oligonucleotides along with formulation, process, and combination drug product development. With extensive pharmaceutical development experience at a broad range of companies, the leadership team at DDE Labs has worked on products across many routes of administration and dosage forms including various non-invasive and sustained release injection products. DDE Labs is located at 11494 Sorrento Valley Road, Suite J, San Diego, California. For more information, please visit www.ddelabs.com.