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## **DPT Laboratories names Michael Curry as Director of Sterile Operations**

**SAN ANTONIO** – [DPT Laboratories](#), a contract development and manufacturing organization, announced today that Michael Curry has been named Director of Sterile Operations at the company's [Center of Excellence for Sterile & Specialty Products](#) in Lakewood, New Jersey.

As Director of Sterile Operations, Curry will be responsible for managing the manufacturing operations for nasal products, small volume parenterals and other sterile and specialty products. He will manage all aspects of aseptic manufacturing operations and work closely with DPT customers.

“As we continue to grow our business, it is critical that we continue to build a team that is comprised of the best talent possible,” said Gene Ciolfi, General Manager DPT Lakewood. “Based on Michael’s experience, I am confident that he is the right fit for this exciting and challenging role.”

“I am very excited to join the DPT team,” Curry said. “My objectives are to be an immediate contributor to the DPT team and ensure that the highest quality products are manufactured and distributed from the [Lakewood](#) manufacturing site.”

Curry previously was Director of Field Applications for BioVigilant Systems in Tucson, AZ, where he managed the team of Field Applications Scientists for the United States and provided applications support globally. He was responsible for field applications customer support for North America for the implementation of Rapid Microbiological Methods for environmental monitoring with BioVigilant’s Instantaneous Microbial Detection system (IMD-A). He was also responsible for providing environmental monitoring, manufacturing, validation, quality assurance, and CMC regulatory affairs support to customers implementing the IMD-A technology within their environmental monitoring program.

Prior to Biovigilant, Curry worked with The Medicines Company, Dendreon Corporation, Wyeth Pharmaceuticals, Baxter Bioscience, and Schering-Plough in a variety of manufacturing, quality systems and regulatory compliance roles. He has more than 18 years experience within the pharmaceutical and biotechnology industry and is certified as a Lean Six

Sigma Green Belt. He holds a Bachelor of Science degree in Biology from Ramapo College of New Jersey and a Master of Science in Microbiology from Seton Hall University.

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### **About DPT Laboratories**

DPT, a DFB company, is a contract development and manufacturing organization (CDMO) providing companies the best solutions to their sterile and non-sterile drug development and manufacturing needs through innovation, technology, and service. Specializing in semi-solid and liquid dosage forms, DPT has a reputation for quality, unmatched technical expertise, extensive manufacturing capabilities, and an exemplary regulatory compliance record. With five cGMP facilities in San Antonio and Lakewood, N.J., DPT offers full service outsourcing solutions including stand alone development, site transfers, state-of-the-art manufacturing, packaging, and worldwide distribution. For more information, call 210-476-8100 or visit [www.DPTLabs.com](http://www.DPTLabs.com).