



PEDI Receives California FDB Device Manufacturing License

Carlsbad, CA – October 27, 2003 – PEDI (Plastics Engineering & Development, Inc.), a contract manufacturer and custom injection molding company for the medical and biomedical device industry, announced certification by the State of California Food & Drug Branch (FDB) to manufacture medical devices. The FDB, which partners with the FDA, has authorized PEDI to operate under the rigorous quality guidelines of California's Device Manufacturing License.

Prior to issuing the license the FDB inspects the manufacturing facility and PEDI demonstrated compliance with all applicable state laws including the federal Good Manufacturing Practice (GMP) Quality System Regulation (QSR) regulation adopted as a California regulation. Manufacturers must renew their license annually and the FDB conducts periodic renewal inspections.

For more information on PEDI's services call 1-760-931-1844 or visit www.pedioplastics.com.

About PEDI

Since 1985, a full service contract manufacturer and custom plastic injection molding company. PEDI can manufacture a single component or handle every aspect of a product's development from design, tooling and molding to ultrasonic welding, assembly, and packaging. PEDI specializes in the medical and biomedical device markets with additional expertise in manufacturing implantables using PEEK™ material. Other industries served include electronics, consumer, industrial and recreation products. PEDI holds a California FDB Device Manufacturing License and operates a 50,000 sq. ft. facility with a Class 100,000 cleanroom in San Diego County. www.pedioplastics.com