FDA 510(k) Premarket Notification

K071959 ANGIODYNAMICS, INC.NEVERTOUCH 600UM FIBER AND VENACURE PROCEDURE KIT

CLEARED

Aug 1, 2007
16 days to decision

K071959 · Product code: **GEX** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k071959/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Jul 16, 2007
Decision date	Aug 1, 2007
Days to decision	16 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**AngioDynamics, Inc.**
Location	Glens Falls, NY, US
Contact	TERI JUCKETT
Website	http://www.angiodynamics.com/
510(k) history	87 submissions · 82 cleared · 1995-2025

AngioDynamics, Inc. is a global leader in vascular and oncology medical technologies, with a manufacturing facility in Glens Falls, US. The company develops advanced devices addressing blood flow restoration, cancer therapies, vascular access, and varicose vein treatment. AngioDynamics has received FDA 510(k) clearances from total submissions since its first clearance in 1995. The company specializes in cardiovascular devices, with recent cleared products including mechanical aspiration systems, infusion systems, and angiographic catheters. The latest FDA 510(k) clearance...