

**K070378 ANGIODYNAMICS, INC. 600UM FIBER AND
VENACURE PROCEDURE KIT**Mar 9, 2007
29 days to decisionK070378 · Product code: **GEX** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k070378/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Feb 8, 2007
Decision date	Mar 9, 2007
Days to decision	29 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...

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Device record: <https://www.510kdatabase.net/k070378/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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