FDA 510(k) Premarket Notification

CLEARED

K033576 LIFESHIELD LATEX-FREE PRIMARY IV PUMP SET DISTAL MICROBORE PATIENT LINE, CONVERTIBLE PIN, 72 INCH, WITH 2 PRESSURE

Dec 4, 2003
21 days to decision

K033576 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k033576/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Set, Administration, Intravascular (FPA)
Date received	Nov 13, 2003
Decision date	Dec 4, 2003
Days to decision	21 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Abbott Laboratories**
Location	Abbott Park, IL, US
Contact	PATRICIA MELERSKI
Website	http://www.abbott.com
510(k) history	883 submissions · 868 cleared · 1976-2026

Abbott Laboratories is an American multinational medical devices and health care company headquartered in Abbott Park, Illinois. The company operates in over 160 countries and produces pharmaceuticals, diagnostics, nutritional products, and medical devices. Abbott maintains a substantial FDA 510(k) regulatory record with FDA 510(k) cleared devices from total submissions since 1976. The company's cleared devices span chemistry, microbiology, hematology, immunology, and toxicology categories. The latest clearance in 2025 reflects continued regulatory activity and product de...