FDA 510(k) Premarket Notification

**CLEARED**

# K024126 CLEAR OR BRAIDED CONTRAST MEDIA INJECTION LINE OR WITHOUT ROTATOR, CATH LAB KIT

Jan 14, 2003
29 days to decision

K024126 · Product code: **DQO** · Cardiovascular
Source: https://www.510kdatabase.net/k024126/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Special |
| Device classification | Catheter, Intravascular, Diagnostic (DQO) |
| Date received | Dec 16, 2002 |
| Decision date | Jan 14, 2003 |
| Days to decision | 29 days |
| Third-party review | No |
| Summary / Statement | Statement |

## APPLICANT

| | |
|---|---|
| Company | **Abbott Laboratories** |
| Location | Abbott Park, IL, US |
| Contact | NICOHL R WILDING |
| 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...