

K101339 ARCHER SUPER STIFF GUIDEWIRESAug 31, 2010
111 days to decisionK101339 · Product code: **DQX** · Cardiovascular
Source: <https://www.510kdatabase.net/k101339/>**SUBMISSION DETAILS**

| | |
|-----------------------|------------------------------------|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Wire, Guide, Catheter (DQX) |
| Date received | May 12, 2010 |
| Decision date | Aug 31, 2010 |
| Days to decision | 111 days |
| Third-party review | No |
| Summary / Statement | Summary |

APPLICANT

| | |
|----------------|---|
| Company | Medtronic, Inc. |
| Location | Mounds View, MN, US |
| Contact | COLLEEN MULLINS |
| Website | https://www.medtronic.com |
| 510(k) history | 209 submissions · 208 cleared · 1981-2026 |

Medtronic, Inc. is a global medical device manufacturer headquartered in Mounds View, United States. The company develops and markets a broad range of medical devices across multiple therapeutic areas. Medtronic maintains a strong FDA 510(k) regulatory track record with FDA 510(k) cleared devices from total submissions since 1981. The company specializes primarily in Cardiovascular devices, which represent 82% of its submission portfolio. Recent clearances include coronary perfusion cannulae, intracoronary shunts, venous cannulae, guidewires, deflectable catheter systems,...
