

K853352 FUKUDA DENSHI DS-1060 DYNASCOPE PATIENT MONITORNov 14, 1985
94 days to decisionK853352 · Product code: **DSI** · Cardiovascular
Source: <https://www.510kdatabase.net/k853352/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Aug 12, 1985
Decision date	Nov 14, 1985
Days to decision	94 days
Third-party review	No

APPLICANT

Company	Fukuda Denshi USA, Inc.
Location	Mchenry, IL, US
Contact	DON ANDRESEN
510(k) history	68 submissions · 68 cleared · 1984-2018

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k853352/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 18, 2026