

**K032290 FUKUDA DENSHI DYNASCOPE MODEL DS-7100
SERIES PORTABLE PATIENT MONITOR**Oct 10, 2003
78 days to decisionK032290 · Product code: **MHX** · Cardiovascular
Source: <https://www.510kdatabase.net/k032290/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Monitor, Physiological, Patient(with Arrhythmia Detection Or Alarms) (MHX)
Date received	Jul 24, 2003
Decision date	Oct 10, 2003
Days to decision	78 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k032290/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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