

K813023 FUKUDA DENSHI CARDIMAX FK-11Dec 2, 1981
36 days to decisionK813023 · Product code: **DSP** · Cardiovascular
Source: <https://www.510kdatabase.net/k813023/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Balloon, Intra-aortic And Control (DSP)
Date received	Oct 27, 1981
Decision date	Dec 2, 1981
Days to decision	36 days
Third-party review	No

APPLICANT

Company	Brentwood Instruments, Inc.
Location	Mchenry, IL, US
510(k) history	21 submissions · 20 cleared · 1981-1990

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Device record: <https://www.510kdatabase.net/k813023/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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