

K760368 MONITORING & STRESS TESTING EQUIPMENTFeb 28, 1977
208 days to decisionK760368 · Product code: **DSI** · Cardiovascular
Source: <https://www.510kdatabase.net/k760368/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Aug 4, 1976
Decision date	Feb 28, 1977
Days to decision	208 days
Third-party review	No

APPLICANT

Company	Advanced Medical, Inc.
Location	Mchenry, IL, US
510(k) history	3 submissions · 3 cleared · 1977-1989

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Device record: <https://www.510kdatabase.net/k760368/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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