

K802472 BIOMETRIC ERG SYSTEMNov 26, 1980
48 days to decisionK802472 · Product code: **DSI** · Cardiovascular
Source: <https://www.510kdatabase.net/k802472/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Oct 9, 1980
Decision date	Nov 26, 1980
Days to decision	48 days
Third-party review	No

APPLICANT

Company	Biometric Analysis Group, Inc.
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1980-1980

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Device record: <https://www.510kdatabase.net/k802472/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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