

K843731 KENZ 80Z MARK IIJul 11, 1985
290 days to decisionK843731 · Product code: **DSI** · CardiovascularSource: <https://www.510kdatabase.net/k843731/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Sep 24, 1984
Decision date	Jul 11, 1985
Days to decision	290 days
Third-party review	No

APPLICANT

Company	Suzuken Co., Ltd.
Location	Walker, MI, US
510(k) history	13 submissions · 13 cleared · 1981-2018

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Device record: <https://www.510kdatabase.net/k843731/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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