FDA 510(k) Premarket Notification

K841083 QUIKPAGE ARRHYTHMIA REPORT GENERATOR

CLEARED

Feb 7, 1985
331 days to decision

K841083 · Product code: **DSI** · Cardiovascular
Source: https://www.510kdatabase.net/k841083/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Mar 13, 1984
Decision date	Feb 7, 1985
Days to decision	331 days
Third-party review	No

APPLICANT

Company	**Agilent Technologies**
Location	Walker, MI, US
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510(k) history	13 submissions · 13 cleared · 1982-2001