FDA 510(k) Premarket Notification

K820820 CARDIO REVUE CENTER

CLEARED

Apr 29, 1982
36 days to decision

K820820 · Product code: **DXG** · Cardiovascular
Source: https://www.510kdatabase.net/k820820/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Computer, Diagnostic, Pre-programmed, Single-function (DXG)
Date received	Mar 24, 1982
Decision date	Apr 29, 1982
Days to decision	36 days
Third-party review	No

APPLICANT

Company	**Diasonics, Inc.**
Location	Mchenry, IL, US
510(k) history	42 submissions · 41 cleared · 1978-1997