FDA 510(k) Premarket Notification

K821090 CARDI-O-LIFE MODEL 8823

CLEARED

Jul 8, 1982
80 days to decision

K821090 · Product code: **DPS** · Cardiovascular
Source: https://www.510kdatabase.net/k821090/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrocardiograph (DPS)
Date received	Apr 19, 1982
Decision date	Jul 8, 1982
Days to decision	80 days
Third-party review	No

APPLICANT

Company	**Amertek**
Location	Walker, MI, US
510(k) history	2 submissions · 2 cleared · 1981-1982