FDA 510(k) Premarket Notification

K782033 CARDIAC CARE Q/C TEST KIT

CLEARED

Dec 15, 1978
9 days to decision

K782033 · Product code: **DRL** · Cardiovascular
Source: https://www.510kdatabase.net/k782033/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tester, Defibrillator (DRL)
Date received	Dec 6, 1978
Decision date	Dec 15, 1978
Days to decision	9 days
Third-party review	No

APPLICANT

Company	**Instrumentation Laboratory CO**
Location	Mchenry, IL, US
510(k) history	321 submissions · 320 cleared · 1976-2023