FDA 510(k) Premarket Notification

K821518 MYO/CLIP

CLEARED

Jul 8, 1982
48 days to decision

K821518 · Product code: **LDF** · Cardiovascular
Source: https://www.510kdatabase.net/k821518/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Pacemaker, Temporary (LDF)
Date received	May 21, 1982
Decision date	Jul 8, 1982
Days to decision	48 days
Third-party review	No

APPLICANT

Company	**Alto Development Corp.**
Location	Mchenry, IL, US
510(k) history	10 submissions · 10 cleared · 1982-1996