FDA 510(k) Premarket Notification

K781518 PREGELLED DISPOSABLE ELECTRODE

CLEARED

Oct 6, 1978
36 days to decision

K781518 · Product code: **DRX** · Cardiovascular
Source: https://www.510kdatabase.net/k781518/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Aug 31, 1978
Decision date	Oct 6, 1978
Days to decision	36 days
Third-party review	No

APPLICANT

Company	**Vente Medical Products, Inc.**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1978-1978