FDA 510(k) Premarket Notification

K030556 STEELEX ELECTRODE SET

CLEARED

Apr 22, 2003
60 days to decision

K030556 · Product code: **LDF** · Cardiovascular
Source: https://www.510kdatabase.net/k030556/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Pacemaker, Temporary (LDF)
Date received	Feb 21, 2003
Decision date	Apr 22, 2003
Days to decision	60 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Aesculap, Inc.**
Location	Burlingame, CA, US
Contact	GEORG KELLER
510(k) history	207 submissions · 201 cleared · 1991-2025