FDA 510(k) Premarket Notification

K954772 QUICKELS QS 200 SYSTEM

CLEARED

Jul 11, 1996
272 days to decision

K954772 · Product code: **DRX** · Cardiovascular
Source: https://www.510kdatabase.net/k954772/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Oct 13, 1995
Decision date	Jul 11, 1996
Days to decision	272 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Ferguson Medical**
Location	Chico, CA, US
Contact	FRANK FERGUSON
510(k) history	57 submissions · 55 cleared · 1985-2004