FDA 510(k) Premarket Notification

K820438 QUINTON Q2000 ETT MONITORING SYSTEM

CLEARED

Apr 29, 1982
71 days to decision

K820438 · Product code: **DSI** · Cardiovascular
Source: https://www.510kdatabase.net/k820438/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Feb 17, 1982
Decision date	Apr 29, 1982
Days to decision	71 days
Third-party review	No

APPLICANT

Company	**Quinton, Inc.**
Location	Mchenry, IL, US
510(k) history	164 submissions · 160 cleared · 1976-2003