FDA 510(k) Premarket Notification

K893420 MASTERLAB PFT AND MASTERLAB BODY BOX

CLEARED

Jun 23, 1989
57 days to decision

K893420 · Product code: **CCM** · Anesthesiology
Source: https://www.510kdatabase.net/k893420/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Plethysmograph, Pressure (CCM)
Date received	Apr 27, 1989
Decision date	Jun 23, 1989
Days to decision	57 days
Third-party review	No

APPLICANT

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