FDA 510(k) Premarket Notification

K930286 DRI-VENDT/PROVENT/ACCUVENT ARTERIAL BLOOD GAS KITS

CLEARED

Jun 22, 1993
152 days to decision

K930286 · Product code: **CBT** · Anesthesiology
Source: https://www.510kdatabase.net/k930286/

SUBMISSION DETAILS

Decision	Substantially Equivalent - KD
Submission type	Traditional
Device classification	Arterial Blood Sampling Kit (CBT)
Date received	Jan 21, 1993
Decision date	Jun 22, 1993
Days to decision	152 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Concord/Portex**
Location	Keene, NH, US
Contact	ROBERT WHEELER
510(k) history	23 submissions · 20 cleared · 1989-1993