FDA 510(k) Premarket Notification

K944120 FI02 SENSING MODULE

CLEARED

May 10, 1995
260 days to decision

K944120 · Product code: **CCL** · Anesthesiology
Source: https://www.510kdatabase.net/k944120/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Analyzer, Gas, Oxygen, Gaseous-phase (CCL)
Date received	Aug 23, 1994
Decision date	May 10, 1995
Days to decision	260 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Core-M, L.P.**
Location	Allston, MA, US
Contact	NATAN E PARSONS
510(k) history	1 submissions · 1 cleared · 1995-1995