FDA 510(k) Premarket Notification

CLEARED

K952325 ET TUBE PLACEMENT DETECTOR (WITH COLIBRI INDICATOR)

Jul 31, 1995
74 days to decision

K952325 · Product code: **CCK** · Anesthesiology
Source: https://www.510kdatabase.net/k952325/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Analyzer, Gas, Carbon-dioxide, Gaseous-phase (CCK)
Date received	May 18, 1995
Decision date	Jul 31, 1995
Days to decision	74 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Icor AB**
Location	Bromma, SE
Contact	ANDRAS GEDEON
510(k) history	16 submissions · 16 cleared · 1986-1996