FDA 510(k) Premarket Notification

K833397 TRI-MED 511 INFANT RESPIRATION MONITOR

CLEARED

Nov 28, 1983
59 days to decision

K833397 · Product code: **CCK** · Anesthesiology
Source: https://www.510kdatabase.net/k833397/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Analyzer, Gas, Carbon-dioxide, Gaseous-phase (CCK)
Date received	Sep 30, 1983
Decision date	Nov 28, 1983
Days to decision	59 days
Third-party review	No

APPLICANT

Company	**Tri-Med, Inc.**
Location	Mchenry, IL, US
510(k) history	29 submissions · 29 cleared · 1977-2004