FDA 510(k) Premarket Notification

K832999 RESPIRATION MONITOR

CLEARED

Feb 13, 1984
179 days to decision

K832999 · Product code: **FLS** · Anesthesiology
Source: https://www.510kdatabase.net/k832999/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Apnea, Facility Use (FLS)
Date received	Aug 18, 1983
Decision date	Feb 13, 1984
Days to decision	179 days
Third-party review	No

APPLICANT

Company	**Litton Medical Electronics**
Location	Walker, MI, US
510(k) history	38 submissions · 38 cleared · 1982-1985