FDA 510(k) Premarket Notification

K801330 ANSPACH LEG SUPPORT

CLEARED

Jun 26, 1980
23 days to decision

K801330 · Product code: **CCX** · Anesthesiology
Source: https://www.510kdatabase.net/k801330/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Support, Patient Position (CCX)
Date received	Jun 3, 1980
Decision date	Jun 26, 1980
Days to decision	23 days
Third-party review	No

APPLICANT

Company	**The Anspach Effort, Inc.**
Location	Mchenry, IL, US
510(k) history	60 submissions · 60 cleared · 1980-2022