FDA 510(k) Premarket Notification

K840108 NEUROMUSCULAR TRANSMISSION MONITOR

CLEARED

Feb 17, 1984
38 days to decision

K840108 · Product code: **KOI** · Anesthesiology
Source: https://www.510kdatabase.net/k840108/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Peripheral, Electric (KOI)
Date received	Jan 10, 1984
Decision date	Feb 17, 1984
Days to decision	38 days
Third-party review	No

APPLICANT

Company	**Datex Division Instrumentarium Corp.**
Location	Walker, MI, US
510(k) history	55 submissions · 55 cleared · 1982-1997