FDA 510(k) Premarket Notification

K812765 DATASCRIBE #DS-40

CLEARED

Nov 16, 1981
42 days to decision

K812765 · Product code: **BXW** · Anesthesiology
Source: https://www.510kdatabase.net/k812765/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Calibrator, Volume, Gas (BXW)
Date received	Oct 5, 1981
Decision date	Nov 16, 1981
Days to decision	42 days
Third-party review	No

APPLICANT

Company	**Timeter Instrument Corp.**
Location	Mchenry, IL, US
510(k) history	29 submissions · 29 cleared · 1976-1988