FDA 510(k) Premarket Notification

K800373 TENDERLIFT

CLEARED

Mar 12, 1980
21 days to decision

K800373 · Product code: **FMR** · General Hospital
Source: https://www.510kdatabase.net/k800373/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Transfer, Patient, Manual (FMR)
Date received	Feb 20, 1980
Decision date	Mar 12, 1980
Days to decision	21 days
Third-party review	No

APPLICANT

Company	**Competent Design**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1980-1980