

K800373 TENDERLIFTMar 12, 1980
21 days to decisionK800373 · 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

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Device record: <https://www.510kdatabase.net/k800373/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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