FDA 510(k) Premarket Notification

K800482 STEAROTHERMOHILUS SPORE STRIPS

CLEARED

Apr 16, 1980
43 days to decision

K800482 · Product code: **FRC** · General Hospital
Source: https://www.510kdatabase.net/k800482/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Indicator, Biological Sterilization Process (FRC)
Date received	Mar 4, 1980
Decision date	Apr 16, 1980
Days to decision	43 days
Third-party review	No

APPLICANT

Company	**Mdt Corp., Inc.**
Location	Mchenry, IL, US
510(k) history	22 submissions · 22 cleared · 1979-1996