

K800482 STEAROTHERMOHILUS SPORE STRIPSApr 16, 1980
43 days to decisionK800482 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k800482/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 22, 2026