

K924380 MDT/HARVEY MODEL 7000 & 8000 CHEMICLAVEFeb 16, 1994
534 days to decisionK924380 · Product code: **MLR** · General Hospital
Source: <https://www.510kdatabase.net/k924380/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Sterilizer, Chemical (MLR)
Date received	Aug 31, 1992
Decision date	Feb 16, 1994
Days to decision	534 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k924380/> 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