FDA 510(k) Premarket Notification

K924380 MDT/HARVEY MODEL 7000 & 8000 CHEMICLAVE

CLEARED

Feb 16, 1994
534 days to decision

K924380 · 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