

K042763 MODIFICATION TO MEDLINE STRIDER MIDI 3Oct 7, 2004
2 days to decisionK042763 · Product code: **INI** · Physical MedicineSource: <https://www.510kdatabase.net/k042763/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Vehicle, Motorized 3-wheeled (INI)
Date received	Oct 5, 2004
Decision date	Oct 7, 2004
Days to decision	2 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Statement

APPLICANT

Company	Medline Industries, Inc.
Location	Mchenry, IL, US
Contact	ANDREA HAFERKAMP
510(k) history	238 submissions · 234 cleared · 1977-2024

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k042763/> 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 June 28, 2026