FDA 510(k) Premarket Notification

# K042763 MODIFICATION TO MEDLINE STRIDER MIDI 3

**CLEARED**

Oct 7, 2004
2 days to decision

K042763 · Product code: **INI** · Physical Medicine
Source: 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 |