FDA 510(k) Premarket Notification

# K873716 ADULT WALKING HEEL PRODUCT NO. 4183-130

**CLEARED**

Oct 2, 1987
17 days to decision

K873716 · Product code: **LDF** · Cardiovascular
Source: https://www.510kdatabase.net/k873716/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Electrode, Pacemaker, Temporary (LDF) |
| Date received | Sep 15, 1987 |
| Decision date | Oct 2, 1987 |
| Days to decision | 17 days |
| Third-party review | No |

## APPLICANT

| | |
|---|---|
| Company | **Martin Medical** |
| Location | Desoto, KS, US |
| Contact | RANDY KILBURN |
| 510(k) history | 14 submissions · 14 cleared · 1986-1987 |