FDA 510(k) Premarket Notification

# K760504 PLETHYSMOGRAPH (MDI SYSTEM II)

**CLEARED**

Aug 26, 1976
3 days to decision

K760504 · Product code: **JOM** · Cardiovascular
Source: https://www.510kdatabase.net/k760504/

## SUBMISSION DETAILS

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|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Plethysmograph, Photoelectric, Pneumatic Or Hydraulic (JOM) |
| Date received | Aug 23, 1976 |
| Decision date | Aug 26, 1976 |
| Days to decision | 3 days |
| Third-party review | No |

## APPLICANT

| | |
|---|---|
| Company | **Marketing Development Intl.** |
| Location | Mchenry, IL, US |
| 510(k) history | 1 submissions · 1 cleared · 1976-1976 |