FDA 510(k) Premarket Notification

# K032624 MICRUS MICROCATHETER

**CLEARED**

Sep 10, 2003
15 days to decision

K032624 · Product code: **DQO** · Cardiovascular
Source: https://www.510kdatabase.net/k032624/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Catheter, Intravascular, Diagnostic (DQO) |
| Date received | Aug 26, 2003 |
| Decision date | Sep 10, 2003 |
| Days to decision | 15 days |
| Third-party review | Yes |
| Summary / Statement | Summary |

## APPLICANT

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|---|---|
| Company | **Micrus Corp.** |
| Location | Mountain View, CA, US |
| Contact | MARGARET WEBBER |
| 510(k) history | 6 submissions · 6 cleared · 2001-2004 |