FDA 510(k) Premarket Notification

# K040922 VENTURE WIRE CONTROL CATHETER

**CLEARED**

Aug 18, 2004
132 days to decision

K040922 · Product code: **DQY** · Cardiovascular
Source: https://www.510kdatabase.net/k040922/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Catheter, Percutaneous (DQY) |
| Date received | Apr 8, 2004 |
| Decision date | Aug 18, 2004 |
| Days to decision | 132 days |
| Third-party review | No |
| Summary / Statement | Summary |

## APPLICANT

| | |
|---|---|
| Company | **Velocimed, Inc.** |
| Location | Mineapolis, MN, US |
| Contact | Sew-Wah Tay |
| 510(k) history | 5 submissions · 5 cleared · 2004-2006 |