FDA 510(k) Premarket Notification

# K880761 TAPER-22 STEERABLE GUIDEWIRE

**CLEARED**

Aug 19, 1988
176 days to decision

K880761 · Product code: **DQX** · Cardiovascular
Source: https://www.510kdatabase.net/k880761/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Wire, Guide, Catheter (DQX) |
| Date received | Feb 25, 1988 |
| Decision date | Aug 19, 1988 |
| Days to decision | 176 days |
| Third-party review | No |

## APPLICANT

| | |
|---|---|
| Company | **Target Therapeutics** |
| Location | Los Angeles, CA, US |
| Contact | JONI M SNYDER |
| 510(k) history | 70 submissions · 70 cleared · 1985-1998 |