FDA 510(k) Premarket Notification

# K915347 SCIMED VIPER CATHETER

**CLEARED**

Aug 17, 1992
265 days to decision

K915347 · Product code: **LIT** · Cardiovascular
Source: https://www.510kdatabase.net/k915347/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Catheter, Angioplasty, Peripheral, Transluminal (LIT) |
| Date received | Nov 26, 1991 |
| Decision date | Aug 17, 1992 |
| Days to decision | 265 days |
| Third-party review | No |
| Summary / Statement | Summary |

## APPLICANT

| | |
|---|---|
| Company | **Scimed** |
| Location | Maple Grove, MN, US |
| Contact | DARLENE THOMETZ |
| 510(k) history | 8 submissions · 8 cleared · 1990-1999 |