FDA 510(k) Premarket Notification

# K881290 MODIFIED SHILEY DUAL LUMEN SUBCLAVIAN CANNULA

**CLEARED**

Apr 27, 1988
30 days to decision

K881290 · Product code: **LFJ** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k881290/

## SUBMISSION DETAILS

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|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Catheter, Subclavian (LFJ) |
| Date received | Mar 28, 1988 |
| Decision date | Apr 27, 1988 |
| Days to decision | 30 days |
| Third-party review | No |

## APPLICANT

| | |
|---|---|
| Company | **Shiley, Inc.** |
| Location | Mchenry, IL, US |
| Contact | GANZ POBUDA |
| 510(k) history | 174 submissions · 174 cleared · 1976-1993 |