FDA 510(k) Premarket Notification

# K921755 8 FR SINGLE LUMEN HIGH PERF. GROSHONG CENT. VENOUS

**CLEARED**

Oct 23, 1992
193 days to decision

K921755 · Product code: **LJS** · General Hospital
Source: https://www.510kdatabase.net/k921755/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Catheter, Intravascular, Therapeutic, Long-term Greater Than 30 Days (LJS) |
| Date received | Apr 13, 1992 |
| Decision date | Oct 23, 1992 |
| Days to decision | 193 days |
| Third-party review | No |
| Summary / Statement | Summary |

## APPLICANT

| | |
|---|---|
| Company | **Davol, Inc.** |
| Location | Mchenry, IL, US |
| Contact | JACK SPEER |
| 510(k) history | 50 submissions · 47 cleared · 1977-2025 |