FDA 510(k) Premarket Notification

# K912069 AVI MODEL 4225 VENTED PIGGYBACK IV ADMIN SET

**CLEARED**

Oct 14, 1993
888 days to decision

K912069 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k912069/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Set, Administration, Intravascular (FPA) |
| Date received | May 10, 1991 |
| Decision date | Oct 14, 1993 |
| Days to decision | 888 days |
| Third-party review | No |
| Summary / Statement | Summary |

## APPLICANT

| | |
|---|---|
| Company | **3M Avi, Inc.** |
| Location | St. Paul, MN, US |
| Contact | VON BUSCH |
| 510(k) history | 18 submissions · 18 cleared · 1990-1993 |