FDA 510(k) Premarket Notification

# K072316 PRECIOUS PLUS GUIDE CATHETER

**CLEARED**

Sep 17, 2007
31 days to decision

K072316 · Product code: **DQY** · Cardiovascular
Source: https://www.510kdatabase.net/k072316/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Special |
| Device classification | Catheter, Percutaneous (DQY) |
| Date received | Aug 17, 2007 |
| Decision date | Sep 17, 2007 |
| Days to decision | 31 days |
| Third-party review | No |
| Summary / Statement | Summary |

## APPLICANT

| | |
|---|---|
| Company | **Asahi Intecc Co., Ltd.** |
| Location | Seto-Shi, JP |
| Contact | YOSHI TERAI |
| Website | https://www.asahi-intecc.com |
| 510(k) history | 84 submissions · 84 cleared · 2003-2026 |