FDA 510(k) Premarket Notification

K881006 STOCKERT-SHILEY BUBBLE MONITOR 23-23-01

CLEARED

May 24, 1988
76 days to decision

K881006 · Product code: **DRY** · Cardiovascular
Source: https://www.510kdatabase.net/k881006/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Blood-gas, On-line, Cardiopulmonary Bypass (DRY)
Date received	Mar 9, 1988
Decision date	May 24, 1988
Days to decision	76 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