FDA 510(k) Premarket Notification

K791888 CATHETER CARE KIT REORDER 7940&7941

CLEARED

Oct 17, 1979
36 days to decision

K791888 · Product code: **EFQ** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k791888/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Gauze/sponge, Internal (EFQ)
Date received	Sep 11, 1979
Decision date	Oct 17, 1979
Days to decision	36 days
Third-party review	No

APPLICANT

Company	**Ascep, Inc.**
Location	Mchenry, IL, US
510(k) history	9 submissions · 9 cleared · 1979-1983