FDA 510(k) Premarket Notification

K801632 THE BELLCO FISTULA NEEDLES

CLEARED

Aug 4, 1980
20 days to decision

K801632 · Product code: **FIE** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k801632/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Fistula (FIE)
Date received	Jul 15, 1980
Decision date	Aug 4, 1980
Days to decision	20 days
Third-party review	No

APPLICANT

Company	**Bellco Artifidial Organ Spec., Inc.**
Location	Mchenry, IL, US
510(k) history	3 submissions · 3 cleared · 1980-1980