FDA 510(k) Premarket Notification

K896434 AUTO SUTURE SURGIPORT CANNULA DIAMETER REDUCER*

CLEARED

Nov 16, 1989
7 days to decision

K896434 · Product code: **GEA** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k896434/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cannula, Surgical, General & Plastic Surgery (GEA)
Date received	Nov 9, 1989
Decision date	Nov 16, 1989
Days to decision	7 days
Third-party review	No

APPLICANT

Company	**United States Surgical, A Division of Tyco Healthc**
Location	Mchenry, IL, US
Contact	CURTIS RAYMOND
510(k) history	218 submissions · 200 cleared · 1977-2007