FDA 510(k) Premarket Notification

K900129 MODIFIED AUTO SUTURE ENDOSCOPIC GIA SURG. STAPLER*

CLEARED

Feb 27, 1990
55 days to decision

K900129 · Product code: **GDW** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k900129/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Staple, Implantable (GDW)
Date received	Jan 3, 1990
Decision date	Feb 27, 1990
Days to decision	55 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