

**K900129 MODIFIED AUTO SUTURE ENDOSCOPIC GIA SURG.
STAPLER***Feb 27, 1990
55 days to decisionK900129 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k900129/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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