FDA 510(k) Premarket Notification

K923119 WALLSTENT ESOPHAGEAL PROTHESIS

CLEARED

Sep 1, 1992
67 days to decision

K923119 · Product code: **ESW** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k923119/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Esophageal (ESW)
Date received	Jun 26, 1992
Decision date	Sep 1, 1992
Days to decision	67 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Schneider U.S. Stent Div.**
Location	Plymouth, MN, US
Contact	RUSSELL W OLSON
510(k) history	3 submissions · 3 cleared · 1991-1994