FDA 510(k) Premarket Notification

K910831 ENDOPATH SURGICAL INSTRUMENTS

CLEARED

May 31, 1991
93 days to decision

K910831 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k910831/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Feb 27, 1991
Decision date	May 31, 1991
Days to decision	93 days
Third-party review	No

APPLICANT

Company	**Symbiosis Corp.**
Location	Miami, FL, US
Contact	KEVIN W SMITH
510(k) history	34 submissions · 32 cleared · 1989-1996