FDA 510(k) Premarket Notification

K931793 SYMBIOSIS ENT SURGICAL INSTRUMENTS

CLEARED

Jul 22, 1993
101 days to decision

K931793 · Product code: **LRC** · Ear, Nose, Throat
Source: https://www.510kdatabase.net/k931793/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Ent Manual Surgical (LRC)
Date received	Apr 12, 1993
Decision date	Jul 22, 1993
Days to decision	101 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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