

K931793 SYMBIOSIS ENT SURGICAL INSTRUMENTSJul 22, 1993
101 days to decisionK931793 · Product code: **LRC** · Ear, Nose, ThroatSource: <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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k931793/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 22, 2026