

K910855 ENDOPATH VERESS NEEDLEApr 18, 1991
49 days to decisionK910855 · Product code: **FHP** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k910855/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Pneumoperitoneum, Simple (FHP)
Date received	Feb 28, 1991
Decision date	Apr 18, 1991
Days to decision	49 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k910855/>; 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