

K955182 NIPRO ARTERIAL VENOUS FISTULA NEEDLEJun 4, 1996
204 days to decisionK955182 · Product code: **FIE** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k955182/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Fistula (FIE)
Date received	Nov 13, 1995
Decision date	Jun 4, 1996
Days to decision	204 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Nissho Corp.
Location	Miami, FL, US
Contact	LUIS CANDELARIO
510(k) history	5 submissions · 5 cleared · 1994-1997

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k955182/> 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 June 30, 2026