FDA 510(k) Premarket Notification

K955182 NIPRO ARTERIAL VENOUS FISTULA NEEDLE

CLEARED

Jun 4, 1996
204 days to decision

K955182 · 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