FDA 510(k) Premarket Notification

K032777 MODIFICATION TO: NIPRO SAFE TOUCH SAFETY FISTULA NEEDLE

CLEARED

Nov 14, 2003
67 days to decision

K032777 · Product code: **FOZ** · General Hospital
Source: https://www.510kdatabase.net/k032777/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Catheter, Intravascular, Therapeutic, Short-term Less Than 30 Days (FOZ)
Date received	Sep 8, 2003
Decision date	Nov 14, 2003
Days to decision	67 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Nipro Medical Corp.**
Location	Miami, FL, US
Contact	RICHARD D BLISS
510(k) history	27 submissions · 27 cleared · 1997-2010