FDA 510(k) Premarket Notification

K914475 IV START KIT

CLEARED

Mar 18, 1992
162 days to decision

K914475 · Product code: **LRS** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k914475/

SUBMISSION DETAILS

Decision	Substantially Equivalent - KD
Submission type	Traditional
Device classification	I.v. Start Kit (LRS)
Date received	Oct 8, 1991
Decision date	Mar 18, 1992
Days to decision	162 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Cypress Medical Products, Ltd.**
Location	Mc Henry, IL, US
Contact	VARUN SONI
510(k) history	26 submissions · 19 cleared · 1989-2002