FDA 510(k) Premarket Notification

K930196 CEPTI-SEAL I.V. PREP KIT

CLEARED

Jul 2, 1993
169 days to decision

K930196 · Product code: **LRS** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k930196/

SUBMISSION DETAILS

Decision	Substantially Equivalent - KD
Submission type	Traditional
Device classification	I.v. Start Kit (LRS)
Date received	Jan 14, 1993
Decision date	Jul 2, 1993
Days to decision	169 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Medi-Flex Hospital Products, Inc.**
Location	Overland Park, KS, US
Contact	ORLANDO CORDOVA
510(k) history	13 submissions · 9 cleared · 1991-1997