FDA 510(k) Premarket Notification

K926293 ANGIOCOR SIDEHOLE INFUSION CATHETER

CLEARED

Mar 7, 1994
448 days to decision

K926293 · Product code: **LJS** · General Hospital
Source: https://www.510kdatabase.net/k926293/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Long-term Greater Than 30 Days (LJS)
Date received	Dec 14, 1992
Decision date	Mar 7, 1994
Days to decision	448 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Angiocor**
Location	Great Neck, NY, US
Contact	ANAND AKERKAR
510(k) history	1 submissions · 1 cleared · 1994-1994