FDA 510(k) Premarket Notification

K941850 IMMOBILE NON-STERILE AND IMMOBILE A/C NON-STERILE

CLEARED

Jul 14, 1994
91 days to decision

K941850 · Product code: **KMK** · General Hospital
Source: https://www.510kdatabase.net/k941850/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Intravascular Catheter Securement (KMK)
Date received	Apr 14, 1994
Decision date	Jul 14, 1994
Days to decision	91 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Tnt Moborg Intl. , Ltd.**
Location	Buffalo, NY, US
Contact	DENNIS R TOLLINI
510(k) history	3 submissions · 3 cleared · 1992-1994