

K941850 IMMOBILE NON-STERILE AND IMMOBILE A/C NON-STERILEJul 14, 1994
91 days to decisionK941850 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k941850/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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