FDA 510(k) Premarket Notification

K924568 TUBE, LEAD, AND CORD HOLDER NO. 102

CLEARED

Nov 23, 1992
75 days to decision

K924568 · Product code: **KMK** · General Hospital
Source: https://www.510kdatabase.net/k924568/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Intravascular Catheter Securement (KMK)
Date received	Sep 9, 1992
Decision date	Nov 23, 1992
Days to decision	75 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Ansley Medical Products, Inc.**
Location	Dallas, TX, US
Contact	SHARON D CHEATWOOD
510(k) history	5 submissions · 5 cleared · 1992-1993