

K924568 TUBE, LEAD, AND CORD HOLDER NO. 102Nov 23, 1992
75 days to decisionK924568 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k924568/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated July 7, 2026