FDA 510(k) Premarket Notification

K924427 SYNERMED HDL CHOLESTLEROL REAGENT KIT

CLEARED

Nov 9, 1992
69 days to decision

K924427 · Product code: **LBR** · Chemistry
Source: https://www.510kdatabase.net/k924427/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Ldl & Vldl Precipitation, Hdl (LBR)
Date received	Sep 1, 1992
Decision date	Nov 9, 1992
Days to decision	69 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Synermed, Inc.**
Location	Quebec, Canada, CA
Contact	MARCIA J ARENTZ
510(k) history	45 submissions · 45 cleared · 1990-1997