

K924427 SYNERMED HDL CHOLESTLEROL REAGENT KITNov 9, 1992
69 days to decisionK924427 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k924427/> 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 1, 2026