

**K033899 LIFESTREAM PLUS CHOLESTEROL MONITOR,
MODIFICATION TO RESOLUTION CHOLESTEROL MONITOR**May 20, 2004
155 days to decisionK033899 · Product code: **NFX** · Chemistry
Source: <https://www.510kdatabase.net/k033899/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Test, Cholesterol, Total, Over The Counter (NFX)
Date received	Dec 17, 2003
Decision date	May 20, 2004
Days to decision	155 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Lifestream Technologies, Inc.
Location	Post Falls, ID, US
Contact	JACKSON CONNOLLY
510(k) history	3 submissions · 3 cleared · 1999-2004

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k033899/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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