FDA 510(k) Premarket Notification

K790372 H-D-EL

CLEARED

May 3, 1979
69 days to decision

K790372 · Product code: **LBR** · Chemistry
Source: https://www.510kdatabase.net/k790372/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Ldl & Vldl Precipitation, Hdl (LBR)
Date received	Feb 23, 1979
Decision date	May 3, 1979
Days to decision	69 days
Third-party review	No

APPLICANT

Company	**General Diagnostics**
Location	Mchenry, IL, US
510(k) history	39 submissions · 39 cleared · 1976-1988

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k790372/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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