FDA 510(k) Premarket Notification

K844075 URIC ACID REAGENT SET

CLEARED

Oct 30, 1984
11 days to decision

K844075 · Product code: **LFQ** · Chemistry
Source: https://www.510kdatabase.net/k844075/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Acid, Uric, Acid Reduction Of Ferric Ion (LFQ)
Date received	Oct 19, 1984
Decision date	Oct 30, 1984
Days to decision	11 days
Third-party review	No

APPLICANT

Company	**Livonia Diagnostics, Inc.**
Location	Mchenry, IL, US
Contact	WALTER H LAMBERT
510(k) history	16 submissions · 16 cleared · 1984-1985

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k844075/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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