FDA 510(k) Premarket Notification

CLEARED

K800047 LDC UVIII FIXED WAVELENGTH ULTRAVIOLET

Jan 21, 1980
12 days to decision

K800047 · Product code: **KIE** · Toxicology
Source: https://www.510kdatabase.net/k800047/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Apparatus, High Pressure Liquid Chromatography (KIE)
Date received	Jan 9, 1980
Decision date	Jan 21, 1980
Days to decision	12 days
Third-party review	No

APPLICANT

Company	**Laboratory Data Control**
Location	Mchenry, IL, US
510(k) history	4 submissions · 4 cleared · 1980-1980

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k800047/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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