FDA 510(k) Premarket Notification

K834348 LEECO T3-QUANT DIAG. KIT

CLEARED

Feb 21, 1984
70 days to decision

K834348 · Product code: **CDP** · Chemistry
Source: https://www.510kdatabase.net/k834348/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Radioimmunoassay, Total Triiodothyronine (CDP)
Date received	Dec 13, 1983
Decision date	Feb 21, 1984
Days to decision	70 days
Third-party review	No

APPLICANT

Company	**Leeco Diagnostics, Inc.**
Location	Mchenry, IL, US
510(k) history	49 submissions · 49 cleared · 1979-1989

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k834348/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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