

K830808 LEECO T3-RIA DIAGNOSTIC KITApr 8, 1983
25 days to decisionK830808 · Product code: **CDP** · Chemistry
Source: <https://www.510kdatabase.net/k830808/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Radioimmunoassay, Total Triiodothyronine (CDP)
Date received	Mar 14, 1983
Decision date	Apr 8, 1983
Days to decision	25 days
Third-party review	No

APPLICANT

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

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Device record: <https://www.510kdatabase.net/k830808/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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