FDA 510(k) Premarket Notification

K830808 LEECO T3-RIA DIAGNOSTIC KIT

CLEARED

Apr 8, 1983
25 days to decision

K830808 · 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