

K813059 LEECO TESTO-QUANT DIAG. KITNov 16, 1981
17 days to decisionK813059 · Product code: **CDZ** · Chemistry
Source: <https://www.510kdatabase.net/k813059/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Radioimmunoassay, Testosterones And Dihydrotestosterone (CDZ)
Date received	Oct 30, 1981
Decision date	Nov 16, 1981
Days to decision	17 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/k813059/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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