

K820251 T4 DIAGNOSTIC KITMay 5, 1982
96 days to decisionK820251 · Product code: **CDP** · Chemistry
Source: <https://www.510kdatabase.net/k820251/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Radioimmunoassay, Total Triiodothyronine (CDP)
Date received	Jan 29, 1982
Decision date	May 5, 1982
Days to decision	96 days
Third-party review	No

APPLICANT

Company	Diagnostic Reagents, Inc.
Location	Walker, MI, US
510(k) history	66 submissions · 66 cleared · 1982-1998

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Device record: <https://www.510kdatabase.net/k820251/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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