

K810211 T3 RIA KITFeb 10, 1981
15 days to decisionK810211 · Product code: **CDP** · Toxicology
Source: <https://www.510kdatabase.net/k810211/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Radioimmunoassay, Total Triiodothyronine (CDP)
Date received	Jan 26, 1981
Decision date	Feb 10, 1981
Days to decision	15 days
Third-party review	No

APPLICANT

Company	E. R. Squibb & Sons, Inc.
Location	New York, NY, US
510(k) history	32 submissions · 32 cleared · 1977-1982

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Device record: <https://www.510kdatabase.net/k810211/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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