

K760057 TEST, RIA, TRIODOTHYRONINESep 13, 1976
89 days to decisionK760057 · Product code: **CDP** · Immunology
Source: <https://www.510kdatabase.net/k760057/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Radioimmunoassay, Total Triiodothyronine (CDP)
Date received	Jun 16, 1976
Decision date	Sep 13, 1976
Days to decision	89 days
Third-party review	No

APPLICANT

Company	Smithkline Diagnostics, Inc.
Location	Mchenry, IL, US
510(k) history	42 submissions · 42 cleared · 1976-1996

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k760057/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 14, 2026