

**K881214 IMPULSE T, ASSAY REAGENTS**May 27, 1988  
67 days to decisionK881214 · Product code: **CDP** · Toxicology  
Source: <https://www.510kdatabase.net/k881214/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Radioimmunoassay, Total Triiodothyronine (CDP)
Date received	Mar 21, 1988
Decision date	May 27, 1988
Days to decision	67 days
Third-party review	No

**APPLICANT**

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Company	<b>Sclavo, Inc.</b>
Location	Mchenry, IL, US
Contact	MICHAEL BRINKLEY
510(k) history	82 submissions · 82 cleared · 1977-1992

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k881214/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated June 29, 2026