

**K840050 SOPHEIA T3 EIA KIT**Mar 19, 1984  
73 days to decisionK840050 · Product code: **CDP** · Chemistry  
Source: <https://www.510kdatabase.net/k840050/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Radioimmunoassay, Total Triiodothyronine (CDP)
Date received	Jan 6, 1984
Decision date	Mar 19, 1984
Days to decision	73 days
Third-party review	No

**APPLICANT**

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Company	<b>Diagnostic Products Corp.</b>
Location	Mchenry, IL, US
510(k) history	321 submissions · 321 cleared · 1976-2006

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Device record: <https://www.510kdatabase.net/k840050/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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