

**K802468 T3 UPTAKE TEST SYSTEM**Oct 31, 1980  
22 days to decisionK802468 · Product code: **KHQ** · Chemistry  
Source: <https://www.510kdatabase.net/k802468/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Radioassay, Triiodothyronine Uptake (KHQ)
Date received	Oct 9, 1980
Decision date	Oct 31, 1980
Days to decision	22 days
Third-party review	No

**APPLICANT**

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Company	<b>American Diagnostic Corp.</b>
Location	Mchenry, IL, US
510(k) history	39 submissions · 39 cleared · 1980-2017

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k802468/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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