

**K820340 T3 UPTAKE-SQUIBB DIAGNOSTIC KIT**Mar 4, 1982  
24 days to decisionK820340 · Product code: **KHQ** · Toxicology  
Source: <https://www.510kdatabase.net/k820340/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Radioassay, Triiodothyronine Uptake (KHQ)
Date received	Feb 8, 1982
Decision date	Mar 4, 1982
Days to decision	24 days
Third-party review	No

**APPLICANT**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k820340/> 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