

**K821722 T3 UPTAKE DIAGNOSTIC KIT**Jun 24, 1982  
14 days to decisionK821722 · Product code: **KHQ** · Chemistry  
Source: <https://www.510kdatabase.net/k821722/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Radioassay, Triiodothyronine Uptake (KHQ)
Date received	Jun 10, 1982
Decision date	Jun 24, 1982
Days to decision	14 days
Third-party review	No

**APPLICANT**

---

Company	<b>Diagnostic Reagents, Inc.</b>
Location	Walker, MI, US
510(k) history	66 submissions · 66 cleared · 1982-1998

---

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

Device record: <https://www.510kdatabase.net/k821722/> 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 28, 2026