

**K780292 TRIA-PEG RIA DIAG. KIT**Mar 2, 1978  
9 days to decisionK780292 · Product code: **CDP** · Toxicology  
Source: <https://www.510kdatabase.net/k780292/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Radioimmunoassay, Total Triiodothyronine (CDP)
Date received	Feb 21, 1978
Decision date	Mar 2, 1978
Days to decision	9 days
Third-party review	No

**APPLICANT**

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Company	<b>Nuclear Diagnostics, Inc.</b>
Location	Mchenry, IL, US
510(k) history	22 submissions · 22 cleared · 1977-1994

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

Device record: <https://www.510kdatabase.net/k780292/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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