

K771220 T3 RIA TEST SYSTEMAug 4, 1977
30 days to decisionK771220 · Product code: **CDP** · Chemistry
Source: <https://www.510kdatabase.net/k771220/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Radioimmunoassay, Total Triiodothyronine (CDP)
Date received	Jul 5, 1977
Decision date	Aug 4, 1977
Days to decision	30 days
Third-party review	No

APPLICANT

Company	Monobind
Location	Mchenry, IL, US
510(k) history	21 submissions · 21 cleared · 1977-2003

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k771220/> Data sourced from FDA 510(k) public records (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. - Generated June 29, 2026