FDA 510(k) Premarket Notification

K771220 T3 RIA TEST SYSTEM

CLEARED

Aug 4, 1977
30 days to decision

K771220 · 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