FDA 510(k) Premarket Notification

K881214 IMMPULSE T, ASSAY REAGENTS

CLEARED

May 27, 1988
67 days to decision

K881214 · Product code: **CDP** · Toxicology
Source: https://www.510kdatabase.net/k881214/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Radioimmunoassay, Total Triiodothyronine (CDP)
Date received	Mar 21, 1988
Decision date	May 27, 1988
Days to decision	67 days
Third-party review	No

APPLICANT

Company	**Sclavo, Inc.**
Location	Mchenry, IL, US
Contact	MICHAEL BRINKLEY
510(k) history	82 submissions · 82 cleared · 1977-1992