FDA 510(k) Premarket Notification

K834080 MATCHED COMPONENT SET FOR DIGITOXIN

CLEARED

Feb 4, 1984
68 days to decision

K834080 · Product code: **LCW** · Toxicology
Source: https://www.510kdatabase.net/k834080/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Radioimmunoassay, Digitoxin (125-i) (LCW)
Date received	Nov 28, 1983
Decision date	Feb 4, 1984
Days to decision	68 days
Third-party review	No

APPLICANT

Company	**Cambridge Medical Technology**
Location	Mchenry, IL, US
510(k) history	14 submissions · 14 cleared · 1982-1989

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k834080/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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