

K800883 CARDIACS I TOXICOLOGY CONTROLApr 29, 1980
12 days to decisionK800883 · Product code: **DIF** · Toxicology
Source: <https://www.510kdatabase.net/k800883/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Drug Mixture Control Materials (DIF)
Date received	Apr 17, 1980
Decision date	Apr 29, 1980
Days to decision	12 days
Third-party review	No

APPLICANT

Company	Utak Laboratories, Inc.
Location	Mchenry, IL, US
510(k) history	23 submissions · 23 cleared · 1980-1991

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Device record: <https://www.510kdatabase.net/k800883/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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