

K831737 TRICYCLIC ANTIDEPRESSANTS VIJul 19, 1983
49 days to decisionK831737 · Product code: **DIF** · Toxicology
Source: <https://www.510kdatabase.net/k831737/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Drug Mixture Control Materials (DIF)
Date received	May 31, 1983
Decision date	Jul 19, 1983
Days to decision	49 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/k831737/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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