

K830494 CONTOX DRUG SCREEN CONTROLMar 11, 1983
23 days to decisionK830494 · Product code: **DIF** · Toxicology
Source: <https://www.510kdatabase.net/k830494/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Drug Mixture Control Materials (DIF)
Date received	Feb 16, 1983
Decision date	Mar 11, 1983
Days to decision	23 days
Third-party review	No

APPLICANT

Company	Kaulson Laboratories, Inc.
Location	Mchenry, IL, US
510(k) history	4 submissions · 4 cleared · 1983-1983

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Device record: <https://www.510kdatabase.net/k830494/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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