

K810201 TOXICOLOGY CONTROLFeb 4, 1981
12 days to decisionK810201 · Product code: **LAS** · Toxicology
Source: <https://www.510kdatabase.net/k810201/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Drug Specific Control Materials (LAS)
Date received	Jan 23, 1981
Decision date	Feb 4, 1981
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/k810201/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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