FDA 510(k) Premarket Notification

K810201 TOXICOLOGY CONTROL

CLEARED

Feb 4, 1981
12 days to decision

K810201 · 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