FDA 510(k) Premarket Notification

K823517 THE KLINE APPLICATOR

CLEARED

Jan 18, 1983
50 days to decision

K823517 · Product code: **HGD** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k823517/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Applicator, Vaginal (HGD)
Date received	Nov 29, 1982
Decision date	Jan 18, 1983
Days to decision	50 days
Third-party review	No

APPLICANT

Company	**Kline Technologies, Ltd.**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1983-1983