

**K823517 THE KLINE APPLICATOR**Jan 18, 1983  
50 days to decisionK823517 · Product code: **HGD** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k823517/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k823517/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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