

**K802458 KEVORKIAN-YOUNGE BIOPSY CURETTE #90-6611**Jan 12, 1981  
96 days to decisionK802458 · Product code: **HFB** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k802458/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Forceps, Biopsy, Gynecological (HFB)
Date received	Oct 8, 1980
Decision date	Jan 12, 1981
Days to decision	96 days
Third-party review	No

**APPLICANT**

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Company	<b>J. Sklar Mfg. Co., Inc.</b>
Location	Mchenry, IL, US
510(k) history	18 submissions · 18 cleared · 1977-1984

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k802458/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 30, 2026