

**K882606 EURO-MED ENDO-CURETTE**Jul 25, 1988  
31 days to decisionK882606 · Product code: **PCF** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k882606/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Sampler, Endocervical (PCF)
Date received	Jun 24, 1988
Decision date	Jul 25, 1988
Days to decision	31 days
Third-party review	No

**APPLICANT**

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Company	<b>Buckman Co., Inc.</b>
Location	Mchenry, IL, US
Contact	DAVID W SCHLERF
510(k) history	111 submissions · 104 cleared · 1983-1998

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k882606/>; 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 May 21, 2026