

**K860796 DISPO-URETTE**Apr 8, 1986  
35 days to decisionK860796 · Product code: **PCF** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k860796/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Sampler, Endocervical (PCF)
Date received	Mar 4, 1986
Decision date	Apr 8, 1986
Days to decision	35 days
Third-party review	No

**APPLICANT**

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Company	<b>Roland J. Zwick, Inc.</b>
Location	Walker, MI, US
Contact	ROLAND J ZWICK
510(k) history	2 submissions · 2 cleared · 1984-1986

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k860796/>; 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