

**K923119 WALLSTENT ESOPHAGEAL PROTHESIS**Sep 1, 1992  
67 days to decisionK923119 · Product code: **ESW** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k923119/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Esophageal (ESW)
Date received	Jun 26, 1992
Decision date	Sep 1, 1992
Days to decision	67 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Schneider U.S. Stent Div.</b>
Location	Plymouth, MN, US
Contact	RUSSELL W OLSON
510(k) history	3 submissions · 3 cleared · 1991-1994

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k923119/> 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 July 2, 2026