

**K894939 PERCUTANEOUS ENDOSCOPIC GASTROSTOMY TRAY**Dec 8, 1989  
127 days to decisionK894939 · Product code: **FPD** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k894939/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tube, Feeding (FPD)
Date received	Aug 3, 1989
Decision date	Dec 8, 1989
Days to decision	127 days
Third-party review	No

**APPLICANT**

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Company	<b>North American Sterilization &amp; Packaging Co.</b>
Location	Sparta, NJ, US
Contact	HARRY SCHLAKMAN
510(k) history	16 submissions · 16 cleared · 1989-1997

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k894939/>; 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 26, 2026