

**K910691 NORTEC URETERAL STENT**Aug 15, 1991  
177 days to decisionK910691 · Product code: **FAD** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k910691/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stent, Ureteral (FAD)
Date received	Feb 19, 1991
Decision date	Aug 15, 1991
Days to decision	177 days
Third-party review	No

**APPLICANT**

---

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

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k910691/>, 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