

**K071822 URGENT PC NEUROMODULATION SYSTEM, MODELS
UPC200 AND UPC250-12**Aug 20, 2007
48 days to decisionK071822 · Product code: **NAM** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k071822/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Peripheral Nerve, Non-implanted, For Urinary Incontinence (NAM)
Date received	Jul 3, 2007
Decision date	Aug 20, 2007
Days to decision	48 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Uroplasty, Inc.
Location	Minneapolis, MN, US
Contact	LISA GALLATIN
510(k) history	8 submissions · 8 cleared · 2005-2012

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k071822/> Data sourced from FDA 510(k) public records (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. - Generated June 14, 2026