

K083704 NUTRAC PELVATOR, MODEL PEL 200Jul 14, 2009
211 days to decisionK083704 · Product code: **KPI** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k083704/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Electrical, Non-implantable, For Incontinence (KPI)
Date received	Dec 15, 2008
Decision date	Jul 14, 2009
Days to decision	211 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Verity Scientific , Ltd.
Location	Oxford, OH, US
Contact	BRENT REIDER
510(k) history	1 submissions · 1 cleared · 2009-2009

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k083704/> 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 22, 2026