FDA 510(k) Premarket Notification

K083704 NUTRAC PELVATOR, MODEL PEL 200

CLEARED

Jul 14, 2009
211 days to decision

K083704 · 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