FDA 510(k) Premarket Notification

K092850 ENDOFLIP

CLEARED

Dec 15, 2009
90 days to decision

K092850 · Product code: **FFX** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k092850/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Gastrointestinal Motility (electrical) (FFX)
Date received	Sep 16, 2009
Decision date	Dec 15, 2009
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Crospon, Ltd.**
Location	Bonita Springs, FL, US
Contact	PAUL DRYDEN
510(k) history	12 submissions · 12 cleared · 2009-2019