FDA 510(k) Premarket Notification

K091305 CLEARPATH

CLEARED

Oct 23, 2009
172 days to decision

K091305 · Product code: **FDF** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k091305/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Colonoscope And Accessories, Flexible/rigid (FDF)
Date received	May 4, 2009
Decision date	Oct 23, 2009
Days to decision	172 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Easyglide , Ltd.**
Location	Bonita Springs, FL, US
Contact	PAUL DRYDEN
510(k) history	5 submissions · 5 cleared · 2009-2012