FDA 510(k) Premarket Notification

K030315 MODIFICATION TO ULTROID HEMORRHOID MANAGEMENT SYSTEM

CLEARED

Feb 27, 2003
28 days to decision

K030315 · Product code: **KNS** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k030315/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Unit, Electrosurgical, Endoscopic (with Or Without Accessories) (KNS)
Date received	Jan 30, 2003
Decision date	Feb 27, 2003
Days to decision	28 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Ultroid, LLC**
Location	Carson City, NV, US
Contact	RONALD R NEWTON
510(k) history	2 submissions · 2 cleared · 2003-2003