FDA 510(k) Premarket Notification

K002427 SANDHILL UNITIP CATHETER PRESSURE SENSOR, MODEL K 5251-L3-0026

CLEARED

Jun 22, 2001
318 days to decision

K002427 · Product code: **FFX** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k002427/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	System, Gastrointestinal Motility (electrical) (FFX)
Date received	Aug 8, 2000
Decision date	Jun 22, 2001
Days to decision	318 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Sandhill Scientific, Inc.**
Location	Mchenry, IL, US
Contact	LIUNDA L DIEDERICH
510(k) history	16 submissions · 16 cleared · 1984-2011