FDA 510(k) Premarket Notification

K990935 JACKSON ESOPHAGEAL DILATOR

CLEARED

May 17, 1999
59 days to decision

K990935 · Product code: **KNQ** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k990935/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dilator, Esophageal (KNQ)
Date received	Mar 19, 1999
Decision date	May 17, 1999
Days to decision	59 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Pilling Weck Surgical**
Location	Fort Washington, PA, US
Contact	ELIZABETH LAZARO
510(k) history	5 submissions · 5 cleared · 1998-2000