

K990935 JACKSON ESOPHAGEAL DILATORMay 17, 1999
59 days to decisionK990935 · Product code: **KNQ** · Gastroenterology & UrologySource: <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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k990935/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 26, 2026