FDA 510(k) Premarket Notification

K810499 PERCUTANEOUS GASTROINTESTINAL TUBE

CLEARED

Mar 4, 1981
8 days to decision

K810499 · Product code: **KNT** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k810499/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tubes, Gastrointestinal (and Accessories) (KNT)
Date received	Feb 24, 1981
Decision date	Mar 4, 1981
Days to decision	8 days
Third-party review	No

APPLICANT

Company	**Health Development Corp.**
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1980-1983