

K810499 PERCUTANEOUS GASTROINTESTINAL TUBEMar 4, 1981
8 days to decisionK810499 · Product code: **KNT** · Gastroenterology & UrologySource: <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

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Device record: <https://www.510kdatabase.net/k810499/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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