FDA 510(k) Premarket Notification

K800419 ADT PERCUTANEOUS TRANSHEPATIC DISP. CHOL

CLEARED

Mar 25, 1980
28 days to decision

K800419 · Product code: **FCG** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k800419/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Biopsy Needle (FCG)
Date received	Feb 26, 1980
Decision date	Mar 25, 1980
Days to decision	28 days
Third-party review	No

APPLICANT

Company	**A.D.T. Lab Industries**
Location	Mchenry, IL, US
510(k) history	21 submissions · 21 cleared · 1980-1981