FDA 510(k) Premarket Notification

K820431 E.R.C.P. CANNULAS

CLEARED

Apr 9, 1982
51 days to decision

K820431 · Product code: **ODD** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k820431/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Endoscopic Retrograde Cholangiopancreatography (ercp) Cannula (ODD)
Date received	Feb 17, 1982
Decision date	Apr 9, 1982
Days to decision	51 days
Third-party review	No

APPLICANT

Company	**American Endoscopy, Inc.**
Location	Mchenry, IL, US
510(k) history	19 submissions · 19 cleared · 1982-1985