FDA 510(k) Premarket Notification

K833417 ENDOSCOPIC RETROGRADE

CLEARED

Nov 28, 1983
56 days to decision

K833417 · Product code: **ODD** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k833417/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Endoscopic Retrograde Cholangiopancreatography (ercp) Cannula (ODD)
Date received	Oct 3, 1983
Decision date	Nov 28, 1983
Days to decision	56 days
Third-party review	No

APPLICANT

Company	**Microvasive**
Location	Mchenry, IL, US
510(k) history	18 submissions · 18 cleared · 1983-1987