FDA 510(k) Premarket Notification

K760182 SHEATH, RESECTOSCOPE

CLEARED

Dec 2, 1976
149 days to decision

K760182 · Product code: **FDC** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k760182/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Resectoscope, Working Element (FDC)
Date received	Jul 6, 1976
Decision date	Dec 2, 1976
Days to decision	149 days
Third-party review	No

APPLICANT

Company	**V. Mueller O.V. Baxter Healthcare Corp.**
Location	Mchenry, IL, US
510(k) history	41 submissions · 41 cleared · 1976-1981