FDA 510(k) Premarket Notification

K760174 ADAPTER, RESECTOSCOPE

CLEARED

Aug 30, 1976
55 days to decision

K760174 · Product code: **FDC** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k760174/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Resectoscope, Working Element (FDC)
Date received	Jul 6, 1976
Decision date	Aug 30, 1976
Days to decision	55 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