FDA 510(k) Premarket Notification

K780062 CATHETER/HEMOSTATIC/GERIATRIC

CLEARED

Feb 14, 1978
33 days to decision

K780062 · Product code: **EZL** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k780062/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Retention Type, Balloon (EZL)
Date received	Jan 12, 1978
Decision date	Feb 14, 1978
Days to decision	33 days
Third-party review	No

APPLICANT

Company	**Kendall Research Center**
Location	Mchenry, IL, US
510(k) history	14 submissions · 14 cleared · 1976-1980