FDA 510(k) Premarket Notification

K780454 ACTIVE-LIFE

CLEARED

Apr 21, 1978
32 days to decision

K780454 · Product code: **EXB** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k780454/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Collector, Ostomy (EXB)
Date received	Mar 20, 1978
Decision date	Apr 21, 1978
Days to decision	32 days
Third-party review	No

APPLICANT

Company	**E. R. Squibb & Sons, Inc.**
Location	New York, NY, US
510(k) history	32 submissions · 32 cleared · 1977-1982