FDA 510(k) Premarket Notification

K810818 KEY-MED FIBRESCOPE DISINFECTION TROLLEY

CLEARED

Apr 8, 1981
15 days to decision

K810818 · Product code: **FEB** · General Hospital
Source: https://www.510kdatabase.net/k810818/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accessories, Cleaning, For Endoscope (FEB)
Date received	Mar 24, 1981
Decision date	Apr 8, 1981
Days to decision	15 days
Third-party review	No

APPLICANT

Company	**Keymed, Inc.**
Location	Mchenry, IL, US
510(k) history	29 submissions · 29 cleared · 1981-2010