

K810818 KEY-MED FIBRESCOPE DISINFECTION TROLLEYApr 8, 1981
15 days to decisionK810818 · Product code: **FEB** · General HospitalSource: <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

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Device record: <https://www.510kdatabase.net/k810818/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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